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## Review



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# Engineering *Saccharomyces cerevisiae* cells for production of fatty acid-derived biofuels and chemicals

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The yeast *Saccharomyces cerevisiae* is a widely used cell factory for the production of fuels and chemicals, in particular ethanol, a biofuel produced in large quantities. With a need for high-energy-density fuels for jets and heavy trucks, there is, however, much interest in the biobased production of hydrocarbons that can be derived from fatty acids. Fatty acids also serve as precursors to a number of oleochemicals and hence provide interesting platform chemicals. Here, we review the recent strategies applied to metabolic engineering of *S. cerevisiae* for the production of fatty acid-derived biofuels and for improvement of the titre, rate and yield (TRY). This includes, for instance, redirection of the flux towards fatty acids through engineering of the central carbon metabolism, balancing the redox power and varying the chain length of fatty acids by enzyme engineering. We also discuss the challenges that currently hinder further TRY improvements and the potential solutions in order to meet the requirements for commercial application.

## 1. Introduction

The growing demand for liquid transport fuels alongside concerns about climate change caused by greenhouse gas emissions from the use of fossil fuels has become one of the greatest challenges for modern society. The generation of biofuels from biomass is a sustainable solution that could substantially decrease the usage of fossil fuels. In past decades, various policies have been established to stimulate global biofuel production [1]. Ethanol is the predominant biofuel and has been used in Europe and the USA since the 1900s [2], and global ethanol production is expected to expand from 120 billion litres in 2017 to 131 billion litres by 2027 [3]. Even though the use of ethanol allows a reduction in CO<sub>2</sub> emissions of up to 80% compared with using petrol, its low energy density and hygroscopicity have become an obstacle for its wider application [4,5]. Fatty acid-derived biofuels, such as fatty alcohols and hydrocarbons, have been proposed as an option for use within transport sectors where there is a need for high-density fuels, e.g. aviation and heavy trucks [6]. Fatty acids are naturally produced by cells for both chemical and energy storage functions. Therefore, producing fatty acid-derived biofuels by microorganisms as an alternative biofuel production method has drawn more and more attention, and significant progress has been achieved owing to the dramatic advances in biotechnology.

The diversity of microorganisms, such as fungi, bacteria and algae, allows for the usage of a wider range of substrates, which enables expansion from the use of solely starch-based agricultural products to lignocellulosic biomass waste, CO<sub>2</sub>, methane, etc. [7]. Bioethanol is currently mainly derived from corn starch or cane sugar, which are fermented by microorganisms, such as

yeast [7–9]. *Yarrowia lipolytica*, a model oleaginous yeast, and other oleaginous yeasts are being established as promising platforms for biofuel generation [10]. However, owing to its essential role in bioethanol production, the yeast *Saccharomyces cerevisiae* has become one of the most intensively industrially applied cell factories, offering the possibility of alternative advanced biofuel production based on existing infrastructures and assets without any extra facility costs. Furthermore, the ease of genetic manipulation and its robustness and tolerance towards harsh conditions in industrial production also contribute to the popularity of yeast as a platform to generate various chemicals [11,12]. We will therefore focus on *S. cerevisiae* in this review.

As the major component of cell membranes, fatty acids and their metabolism have been comprehensively studied in yeast [13]. Metabolic engineering strategies, such as blocking competing pathways, increasing precursor supply and balancing cofactor regeneration in the cell, have been applied to establish optimal native and heterologous pathways for sustainable production of fatty acids [14,15]. In addition, several advanced biofuels including alkanes, fatty alcohols and fatty acid ethyl esters (FAEEs), for which fatty acids serve as precursors, were successfully generated in yeast [16–19]. Progress in synthetic and systems biology has also enabled the construction of yeast strains that produce fatty acids and fatty alcohols with a chain length that cells lack the capacity to generate naturally. Here, we will highlight the major contributions to the production of fatty acid-derived biofuels and chemicals through different metabolic engineering strategies in yeast, and point to the major challenges and directions for future laboratory-scale studies and industrial applications.

## 2. Engineering of central carbon metabolism

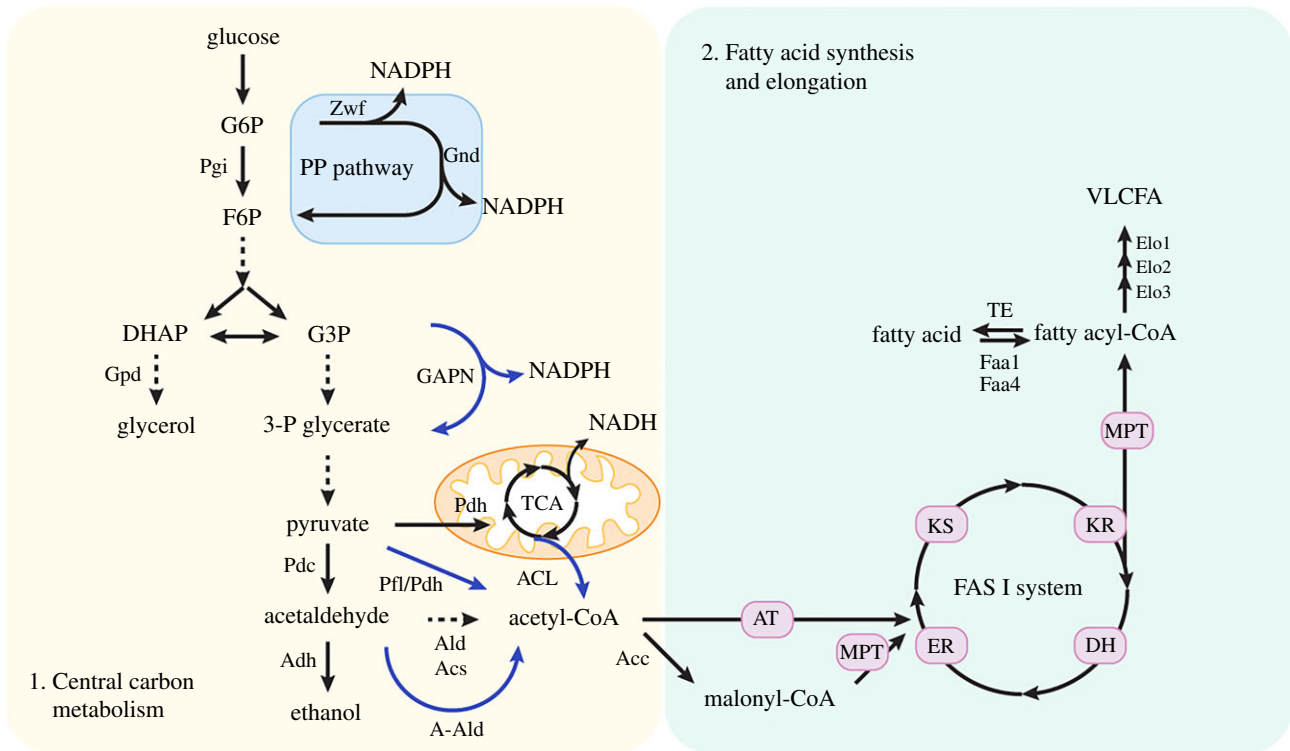
### 2.1. Enhancing the precursor supply for fatty acid synthesis

Fatty acids with a long aliphatic chain are naturally produced by yeast in either their saturated or monounsaturated form. Acetyl-coenzyme A (acetyl-CoA) as the main C2 metabolite is the essential building block for fatty acid synthesis (FAS). Although acetyl-CoA is involved in the metabolic network of *S. cerevisiae* in the cytosol, nucleus, peroxisome and mitochondrion, it is not transported freely across membranes in the absence of the carrier carnitine. However, different shuttle mechanisms exist. The major substrate for de novo FAS in yeast is cytosolic acetyl-CoA, which is generated from pyruvate via three reactions catalysed by pyruvate decarboxylase (Pdc), acetaldehyde dehydrogenase (ALD) and acetyl-CoA synthetase (ACS) (figure 1). Acetaldehyde, as the intermediate generated from pyruvate, is also the precursor for ethanol production, which is an undesired by-product when aiming for high-yield fatty acid production. Thus, many efforts have been directed towards improving the cytosolic acetyl-CoA pool. Alcohol dehydrogenase (ADH) genes were deleted to prevent the conversion from acetaldehyde to ethanol and/or ALD and endogenous or heterologous ACS were overexpressed to enhance the carbon flux to

acetyl-CoA [20]. However, ethanol production is hard to eliminate in yeast by simply deleting ADH genes, as there is a large number of promiscuous ADHs that could catalyse the reaction to generate ethanol and many of these are also involved in other important reactions within the cell [21,22]. In order to overcome this problem, ethanol formation was inhibited by the elimination of all three Pdc enzymes (Pdc1, Pdc5 and Pdc6) [23]. Nevertheless, such a Pdc-negative strain is unable to grow in excess glucose possibly because of repression of the respiratory metabolism and a deficiency in cytosolic C2 supply. Pronk and co-workers [23] solved this problem by evolving the Pdc-deficient strain and succeeded in obtaining a C2-independent Pdc-negative strain which could also grow in excess glucose. Later studies revealed that the adaptation mechanism was associated with an internal deletion in a transcriptional regulator, Mth1, which is involved in glucose sensing in yeast. Such a mutated version of Mth1 reduced the glucose influx and thus resulted in decreased repression of respiration in the evolved strain [24,25].

Although ethanol synthesis is blocked in a Pdc-negative strain, this also decreases the amount of cytosolic acetyl units that serve as precursors for various downstream products including fatty acids. A route relying on mitochondrial Ach1, the CoA transferase hydrolysing mitochondrial acetyl-CoA to acetate that enters the cytosol to provide the C2 unit for cytoplasmic acetyl-CoA synthesis, was shown to compensate for the lack of cytosolic acetyl-CoA synthesis [26]. However, this strategy is restricted by the limited mitochondrial acetyl-CoA supply owing to the stringent regulation of the pyruvate dehydrogenase (PDH) complex and this route cannot function under glucose-repressed conditions (figure 1). Alternatively, a heterologous PDH complex from *Enterococcus faecalis* was expressed in an ACS-deficient yeast strain and shown to fully complement the cytosolic acetyl-CoA supply [27]. Other attempts implemented in yeast to benefit cytosolic acetyl-CoA supply as well include introducing heterologous acetylating ALD (A-Ald), pyruvate-formate lyase (PFL), a phosphoketolase (PHK) pathway or pyruvate oxidase (figure 1) [27–30]. Recently, our group reprogrammed the yeast central metabolism to demonstrate a feasible strategy for industrial production of fatty acids with high titre and yield [31]. A heterologous ATP citrate lyase (ACL) was overexpressed to provide cytosolic acetyl-CoA in an engineered fatty acid overproducing strain (figure 1). Based on that, the three Pdc genes were deleted to abolish ethanol production. After adaptive laboratory evolution (ALE) in glucose, the evolved strain exhibited a pure lipogenesis metabolism, resulting in a great improvement in fatty acid production. Using a similar strategy, expression of ACL from *Y. lipolytica*, downregulation of malate synthase (Mls1) and deletion of glycerol-3-phosphate dehydrogenase (Gpd1) were carried out in *S. cerevisiae*—the latter two being key enzymes involved in competing with FAS for carbon flux—leading to a 70% improvement in free fatty acid production [32].

The conversion of acetyl-CoA to malonyl-CoA via acetyl-coenzyme A carboxylase (ACCase) encoded by *ACC1* is the first committed and rate-limiting step in de novo FAS in yeast. Increasing malonyl-CoA supply is a promising strategy that benefits fatty acid production. In order to break through the limitation of low efficiency of this reaction, the Snf1-dependent phosphorylation of ACCase was—at least



**Figure 1.** Fatty acid synthesis (FAS) pathway in *S. cerevisiae* and engineering strategies of the central carbon metabolism for increasing fatty acid production. The solid arrows and dashed arrows represent single catalytic steps and multiple catalytic steps, respectively. The arrows in blue represent the heterologous pathway that was introduced into yeast. G6P, glucose-6-phosphate; F6P, fructose-6-phosphate; DHAP, dihydroxyacetone phosphate; G3P, glyceraldehyde-3-phosphate; VLCFA, very-long-chain fatty acid; Pgi, phosphoglucose isomerase; Zwf, D-glucose-6-phosphate dehydrogenase; Gnd, phosphogluconate dehydrogenase; Gpd, glycerol-3-phosphate dehydrogenase; GAPN, glyceraldehyde-3-phosphate dehydrogenase; Pdc, pyruvate decarboxylase; Pdh, pyruvate dehydrogenase; A-Ald, acetylating acetaldehyde dehydrogenase; Pfl, pyruvate-formate lyase; Adh, alcohol dehydrogenase; Ald, acetaldehyde dehydrogenase; Acs, acetyl-CoA synthetase; ACL, ATP citrate lyase; Acc, acetyl-coenzyme A carboxylase; TE, thioesterase; Elo1/2/3, fatty acid elongase; Faa1/4, fatty acyl-CoA synthetases; AT, acetyl transferase; MPT, malonyl/palmitoyl transferase; KS, ketoacyl synthase; KR, ketoacyl reductase; CH, dehydratase; ER, enoyl reductase.

partially—abolished by introducing two mutations corresponding to Ser1157 and Ser659 of *ACC1* (*ACC1*<sup>S1157A,S659A</sup>) [33]. An even more efficient catalytic activity was observed when S686A was introduced into the double mutant *ACC1* (*ACC1*<sup>S1157A,S659A,S686A</sup>) [34]. The resulting higher ratio of malonyl-CoA/acetyl-CoA shifts production towards C18 fatty acids, and the overexpression of either wild-type *ACC1* or mutant *ACC1* can lead to an improvement in fatty acid production [16,35].

## 2.2. Balancing cofactor supply and reducing power

Reducing power is an essential element involved in many metabolite conversions. NADH and NADPH, the two major reducing equivalents in yeast, play distinct functions in the cell, i.e. NADH predominantly participates in catabolic reactions and NADPH is mainly required for anabolic reactions [36]. The ratios of the two pyridine nucleotide cofactor systems NADH/NAD<sup>+</sup> and NADPH/NADP<sup>+</sup> are vital for the determination of the cellular redox status and the formation of various metabolites.

Cellular NADPH, the essential reducing equivalent for fatty acid formation and other metabolic conversions in yeast, is predominantly generated from the pentose phosphate (PP) pathway (figure 1). D-Glucose-6-phosphate (G6P) is oxidized through D-glucose-6-phosphate dehydrogenase (G6PDH) encoded by *ZWF1*, which is the first rate-limiting step in the PP pathway. Then, ribulose-5-phosphate is generated through 6-phosphogluconate dehydrogenase

(6PGDH), encoded by *GND1* and *GND2*, thereby yielding two molecules of NADPH. For de novo FAS in yeast, two molecules of NADPH are required as cofactors for each cycle of elongation. Since yeast cells naturally produce excess NADH as the electron carrier, the NADPH supply is often limiting for anabolic reactions. Thus, attempts at metabolic engineering have been made to increase the NADPH supply in the cytosol. To facilitate NADPH regeneration and reduce loss of the carbon source, a non-phosphorylating NADP<sup>+</sup>-dependent glyceraldehyde-3-phosphate dehydrogenase (GAPN) from *Bacillus cereus* was expressed in a yeast strain carrying a deletion in the *GPD1* gene encoding NAD<sup>+</sup>-dependent glycerol-3-phosphate dehydrogenase [37] (figure 1). Extra NADPH can also be produced by overexpression of the otherwise mitochondrial malic enzyme (ME), which converts malate into pyruvate in the cytosol [19,38]. The PHK pathway, which uses xylulose-5-phosphate as a precursor, was introduced into *S. cerevisiae* to increase the NADPH supply in the cytosol. The combination of the PP pathway and PHK pathway as an interesting alternative for fatty acid derivative production resulted in improved production of FAEEs [39]. Fine tuning the flux distribution between the PP pathway and glycolysis by overexpression of phosphogluconate dehydrogenase (encoded by *GND1*), transketolase (encoded by *TKL1*) and transaldolase (encoded by *TAL1*) together with downregulation of phosphoglucose isomerase (encoded by *PGI1*) in yeast to provide additional NADPH led to a 28% increase in free fatty acid production [29].

### 3. De novo fatty acid synthesis in yeast

The biosynthesis of fatty acids in yeast can take place in the cytosol and the mitochondria, where it is carried out by a type I FAS and a type II FAS, respectively. Experimental results suggested that the mitochondrial FAS II pathway is the sole source of the octanoic acid required for lipoic acid production. Lipoic acid serves as an essential cofactor for PDH,  $\alpha$ -ketoglutarate dehydrogenase and the glycine cleavage system [40,41]. However, the range of fatty acids produced by the mitochondrial FAS II pathway and other potential roles in cellular metabolism are still uncertain [41].

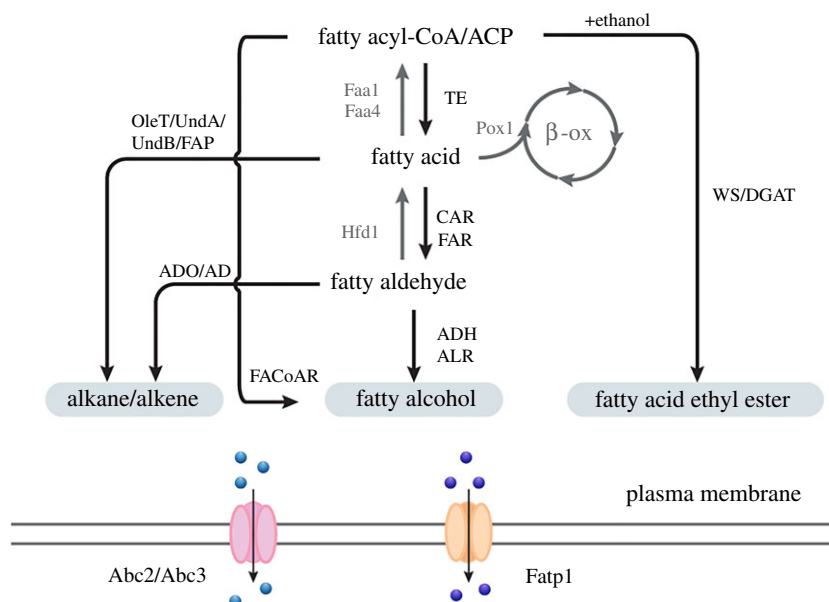
As a type I FAS is responsible for the cytosolic de novo FAS, we will mainly focus on it in this review. The type I FAS in yeast comprises two subunits,  $\alpha$ -subunit Fas2 and  $\beta$ -subunit Fas1. Six copies of eight independent functional domains are assembled into an  $\alpha_6\beta_6$  molecular complex of 2.6 MDa [42] (figure 1). The yeast FAS is activated by its phosphopantetheinyl transferase domain located at the C-terminus of the  $\alpha$ -subunit, and all the reactions occur in the limited space of the  $\alpha_6\beta_6$  complex [42]. The yeast FAS initiates the reaction by transferring acetyl primer and malonyl elongation substrate from acetyl-CoA/malonyl-CoA to the acyl carrier protein (ACP) pantetheine arm by the acetyl transferase (AT) and malonyl/palmitoyl transferase (MPT), respectively. The ketoacyl synthase (KS) condenses them to acetoacetyl-ACP in a malonyl decarboxylation reaction, which is considered the first step of the elongation cycle. Subsequently, the  $\beta$ -ketoacyl-ACP is reduced by the ketoacyl reductase (KR) in the  $\alpha$ -subunit, followed by a dehydration reaction catalysed by the dehydratase (DH) and the second reduction reaction catalysed by the enoyl reductase (ER) in the  $\beta$ -subunit yielding acyl-ACP [43] (figure 1). The ACP domain, which plays the central role in shuttling intermediates between the active sites in the complex, brings the processed acyl chain back to the KS domain for the next elongation cycle [44]. This repetitive process occurs using malonyl-CoA as the provider of 2C units until the carbon chain length of the fatty acid reaches 16 or 18. The end product will be shuttled by ACP from ER to MPT, where it is transferred to CoA and then released. Previous results showed that the overexpression of native *FAS1* and *FAS2* in *S. cerevisiae* could contribute to the fatty acid production as well as introducing heterologous type I or type II FASs [16,19,45,46]. Additionally, in order to overproduce fatty acids in yeast, a common strategy is to overexpress heterologous acyl-ACP or acyl-CoA thioesterases, which can relieve feedback inhibition and increase the fatty acid release [16]. Moreover, the engineering of type I FAS towards short/medium-chain fatty acid (S/MCFA) production has recently attracted attention. A thioesterase from *Acinetobacter baylyi* (*AcTesA*) that has a substrate preference for short/medium-chain acyl-ACP/CoA was embedded into the type I FAS, which benefitted S/MCFA production significantly and led to a 5- to 13-fold increase in S/MCFA production compared with wild-type FAS [47]. Rational modification of (i) the KS domain to restrict chain length elongation, (ii) the MPT domain to reduce the affinity to its substrate malonyl-CoA, and (iii) the AT domain to increase its affinity to acetyl-CoA has succeeded in altering the chain length of fatty acid products resulting in production of extracellular S/MCFAs, mainly hexanoic acid and octanoic acid, of 464 mg l<sup>-1</sup> in total [48]. Very-long-chain fatty acids

(VLCFAs) are the precursors for various valuable chemicals and the essential components for yeast cell membrane structures. VLCFA synthesis occurs at the endoplasmic reticulum membrane with distinct enzymes similar to the domains in the FAS system [49]. Of the three fatty acid elongases (equivalent to the KS domain in FAS), Elo1 is responsible for elongation of C12–16 fatty acids to C16–18 fatty acids, while Elo2 and Elo3 are more specifically responsible for the synthesis of up to C22 fatty acid and C26 fatty acid, respectively [50,51] (figure 1). Yeast was successfully engineered for the production of VLCFAs and derived products by the selective modification of the endogenous yeast fatty acid elongation system together with the expression of a heterologous FAS I system from *Mycobacterium vaccae* [52].

## 4. Fatty acid-derived biofuels and chemicals

### 4.1. Fatty alcohols

Fatty alcohols are important oleochemicals with wide industrial applications ranging from cosmetics to substitutes for petroleum-derived compounds such as biofuels [53]. Fatty alcohols can be generated from fatty acyl-CoAs, fatty acyl-ACPs and fatty acids with fatty aldehydes as the intermediates via the corresponding enzymes fatty acyl-CoA reductase, fatty acyl-ACP reductase and carboxylic acid reductase, respectively [54–56]. These enzymes usually catalyse the first step of the two consecutive reduction steps, i.e. fatty aldehyde formation, followed by the second reaction towards fatty alcohol production via aldehyde reductases/alcohol dehydrogenases (ALRs/ADHs) (figure 2). However, some fatty acyl-CoA/ACP reductases can catalyse the entire four-electron reduction step to generate fatty alcohols directly, for example the well-known FACoAR enzymes from jojoba plant and *Arabidopsis thaliana* [57]. The heterologous pathways including these enzymes have been successfully introduced into yeast. The production of fatty alcohols in *S. cerevisiae* was drastically improved by rewiring central metabolic pathways. The production of free fatty acids which served as the precursors for fatty alcohol formation in this strain was improved significantly by deleting fatty acyl-CoA oxidase (encoded by *POX1*) and fatty acyl-CoA synthetases (encoded by *FAA1* and *FAA4*) (figure 2). Reversal of aldehyde formation was abolished by deleting aldehyde dehydrogenase (encoded by *HFD1*) [19] (figure 2). Combined with (over)expression of the genes involved in fatty alcohol synthesis, which were in this case fatty acyl-CoA reductase FaCoAR from *Marinobacter aquaeolei* VT8, carboxylic acid reductase (CAR) from *Mycobacterium marinum* and native alcohol dehydrogenase Adh5 (figure 2), this led to production of up to 120 mg l<sup>-1</sup> fatty alcohols in shake flasks [19]. In another study, after expression of a high-activity heterologous fatty acid reductase (FAR) (figure 2), blocking competing pathways by deletion of *DGH1*, *HFD1* and *ADH6* together with limiting NADPH and carbon usage by deleting glutamate dehydrogenase encoded by *GDH1*, a strain producing 1.2 g l<sup>-1</sup> fatty alcohols in shake flasks was obtained [58]. However, impaired growth occurred due to intracellular fatty alcohol accumulation. The expression of the multi-functional transporter FATP1 from human in a fatty alcohol producing yeast strain was shown to facilitate



**Figure 2.** Heterologous pathways for fatty acid-derived biofuel synthesis and secretion. The arrows in grey represent the steps involved in fatty acid degradation in yeast. Faa1/4, fatty acyl-CoA synthetases; Pox1, fatty acyl-CoA oxidase; TE, thioesterase; Hfd1, aldehyde dehydrogenase; CAR, carboxylic acid reductase; FAR, fatty acid reductase; ADH, alcohol dehydrogenase; ALR, aldehyde reductase; ADO, aldehyde deformylating oxygenase; AD, aldehyde decarbonylase; WS/DGAT, wax ester synthase/acyl-CoA: diacylglycerol acyltransferase; FAP, fatty acid photodecarboxylase; Abc2/3, alkane transporter from *Yarrowia lipolytica*; Fatp1, mammalian fatty alcohol transporter;  $\beta$ -ox,  $\beta$ -oxidation.

fatty alcohol export (figure 2), which benefited production levels as well as the cell fitness, and resulted in 4.5-fold more extracellular fatty alcohols than the control strain [59]. Based on a yeast strain producing *S*/MCFAs, 1-octanol was successfully generated with the two-step pathway via *M. marinum* CAR and aldehyde reductase Ahr from *Escherichia coli*, demonstrating that the chain length specificity of FAS is the decisive factor for producing fatty alcohols of a specific chain length [60].

## 4.2. Alka(e)nes

Alkenes and alkanes serve as the major constituents of gasoline, diesel and jet fuel. Several alka(e)ne biosynthesis pathways have been successfully demonstrated in microbes in recent years [18,61–63]. Fatty acids, fatty acyl-CoA/ACP and fatty aldehydes are the major precursors that can be used to generate alka(e)nes via corresponding enzymes. Both aldehyde deformylating oxygenase (ADO) and aldehyde decarbonylase (AD) use fatty aldehydes as substrate to facilitate the formation of alkanes (figure 2). ADO is a non-haem di-iron oxygenase requiring molecular oxygen and an external reducing system to provide four electrons, yielding hydrogen peroxide ( $H_2O_2$ ) and formate as by-products [64] (figure 2). The well-known ADs, *Drosophila melanogaster* CYP4G1 and *Arabidopsis* CER1, are naturally involved in long-chain alkane biosynthesis and have been successfully expressed in *S. cerevisiae* to generate  $C_{n-1}$  alkanes from  $C_n$  fatty aldehydes [62]. Fatty acid decarboxylases can catalyse the one-step decarboxylation from  $C_n$  fatty acids to  $C_{n-1}$  1-alkenes in a process that avoids the formation of fatty aldehydes as intermediates. OleT was reported as a cytochrome P450 enzyme that is responsible for the conversion of C12–C20 fatty acids to corresponding 1-alkenes using  $H_2O_2$  as the sole electron and oxygen donor [65] (figure 2). In addition, UndA and UndB were identified as fatty acid decarboxylases for medium-chain 1-alkene synthesis that specifically convert C10–C14

fatty acids and C10–C16 fatty acids, respectively (figure 2). Fatty acyl-CoA/ACPs as substrates can be used to synthesize long-chain alkenes by olefin synthase, a multi-domain polyketide synthase (PKS) from cyanobacteria, through an elongation/sulfonation/decarboxylation mechanism [66]. Furthermore, an algal fatty acid photodecarboxylase (FAP) driven by light was recently found which can convert fatty acids to corresponding alka(e)nes, and it was successfully expressed in *E. coli* to generate hydrocarbons in the presence of visible light [67] (figure 2).

The implementation of alka(e)ne biosynthesis in yeast has made significant progress during recent years. However, the low efficiency of pathway enzymes and the strong competition of fatty alcohol accumulation for metabolic precursors and intermediates are considered the major obstacles of further alka(e)ne production improvement in *S. cerevisiae* [14,68]. Therefore, compartmentalization in yeast organelles turned out to be a promising strategy that provides a suitable environment for alka(e)ne production via isolating the synthesis pathway from the competing pathways in the cytosol. Peroxisomes represent a suitable location for alka(e)ne synthesis not only because of the absence of ALRs/ADHs, but also because of the potential NADPH supply from the peroxisomal NADP-dependent isocitrate dehydrogenase isoenzyme Idp3 [69]. Recently, the alkane synthesis pathway consisting of *Synechococcus elongatus* ADO (*Se*ADO) together with *Mycobacterium marinum* CAR (*Mm*CAR) was targeted to the peroxisomes in yeast, yielding around  $0.12 \text{ mg l}^{-1}$  alkanes, which was a 90% higher alkane titre than that yielded by the cytosolic pathway [14]. After further increasing the precursor supply in the peroxisomes and deleting the cytosolic ALR/ADH genes *ADH5* and *SFA1*, alkane production reached  $1.2 \text{ mg l}^{-1}$  with significantly decreased fatty alcohol accumulation. Moreover, an additional study focusing on medium-chain alkane synthesis in yeast indicates that the compartmentalization in peroxisomes could work as an efficient strategy in this context as well [70].

The toxicity caused by the accumulation of alka(e)nes negatively affects the cell growth and limits the production yield. Consequently, some studies towards improving solvent tolerance were conducted in yeast. The native plasma membrane efflux pumps Snq2 and Pdr5 were identified in *S. cerevisiae* as contributing to alkane export and tolerance by reducing intracellular levels, specifically for C10 and C11 alkanes [71]. The heterologous transporters Abc2 and Abc3 from *Yarrowia lipolytica* significantly increased tolerance against decane and undecane in *S. cerevisiae* through maintaining lower intracellular alkane levels [72] (figure 2). Furthermore, mammalian FATP1 previously identified as a fatty alcohol exporter was expressed in yeast to benefit 1-alkene secretion. The implementation of dynamic regulation, through expressing PfUndB under the control of the GAL7 promoter with deletion of GAL80 to separate the cell growth and production process and replacing the electron transfer system by the NADH-based putidaredoxin reductase system, finally enabled a yeast cell factory to produce 35.3 mg l<sup>-1</sup> 1-alkenes with more than 80% being secreted, which is a 10-fold improvement compared with earlier reported hydrocarbon production by *S. cerevisiae* [63,73].

### 4.3. Fatty acid ethyl esters

The biosynthesis of FAEEs, considered as potential diesel fuel replacement, was demonstrated in yeast. Ethanol and acyl-CoAs are the essential precursors involved in FAEE synthesis that can be catalysed by a wax ester synthase/acyl-CoA: diacylglycerol acyltransferase (WS/DGAT) (figure 2). Generally, most WSs naturally accept acyl groups with a chain length of C16 or C18 and linear alcohols with a chain length ranging from C12 to C20, and various WSs have different substrate chain length preferences [17]. Five heterologous WSs from bacteria and mammals were expressed and evaluated in *S. cerevisiae* to investigate their substrate preferences [17]. The results showed that the WS from *Marinobacter hydrocarbonoclasticus* had the best performance using ethanol as the substrate *in vitro* compared with the other enzymes, and enabled a titre of 6.3 mg l<sup>-1</sup> FAEEs after expressing it in the engineered yeast strain.

In order to establish a stable expression system, the heterologous wax ester synthase gene (*ws2*) was integrated into the yeast chromosomes in multiple copies, resulting in an increase in FAEE production of up to 34 mg l<sup>-1</sup> [74]. Subsequently, the endogenous acyl-CoA binding protein and a bacterial NADP<sup>+</sup>-dependent GAPN were overexpressed in the integration strain to enhance the precursor and cofactor supply, which enabled a further 40% increase in FAEE production. During the synthesis of FAEEs in yeast, the concentrations of ethanol and acyl-CoA influence the yield of the final product. Thus, the carbon flux was redirected towards acetyl-CoA, the precursor of acyl-CoA, by overexpressing the alcohol dehydrogenase (*ADH2*), acetaldehyde dehydrogenase (*ALD6*) and ACS encoded by heterologous gene *acs*<sup>L641P</sup><sub>SE</sub>, together with the integrated *ws2*, resulting in a threefold improvement [39]. Then, the PHK pathway was introduced to enhance acetyl-CoA supply by heterologous expression of *xpkA* and either *ack* from *Aspergillus nidulans* or *pta* from *Bacillus subtilis*. Both PHK pathways helped to generate around 5.0 mg g<sup>-1</sup> cell dry weight FAEEs, an up to a 1.7-fold improvement. Besides, reducing the competition of other pathways for acyl-CoA also permits an overproduction of FAEEs. Therefore,

by eliminating the formation of steryl esters (SEs) and triacylglycerols (TAGs), a threefold increase in FAEE production was achieved [75]. The heterologous expression of a type I FAS from *Brevibacterium ammoniagenes* coupled with WS/DGAT yielded a 6.3-fold increase in FAEE production compared with a strain not containing the heterologous FAS [45]. Additionally, the alternative carbon source, glycerol, with the advantage of being a low-price and highly reduced substrate was used to produce FAEEs in *S. cerevisiae* [76]. The titre of FAEEs reached 0.52 g l<sup>-1</sup> after increasing the ethanol formation from glycerol, blocking the glycerol export route and adding exogenous fatty acids, which is the highest reported FAEE production to date in yeast.

## 5. Perspectives

Progress in developing more advanced biotechnology tools has led to more efficient engineering of microbes. An example is the clustered regularly interspaced short palindromic repeats (CRISPR)/Cas technology, which allows fast multiplex genome editing and has significantly shortened the time required for strain construction. Even though there has been much progress on engineering yeast for production of advanced biofuels in the laboratory, it is still challenging to meet the titre, rate and yield (TRY) requirements for commercial production of low-value fatty acid-derived products (table 1). In order to meet the commercial requirements, the yields and productivities of laboratory-scale processes need to approach around 85% of the theoretically possible yield and the fermentation has to be scaled up drastically [82]. The scientific progress, however, lays the basis for further development in case some of the key barriers can be passed [5].

Owing to the relatively low value of many fatty acid-derived chemicals, improving the utilization of the carbon source is a promising strategy that contributes to high TRY metrics. Therefore, attempts towards the utilization of single carbon feedstock such as CO<sub>2</sub> and methane have attracted much attention, and were shown to be a feasible alternative with high carbon- and energy-conversion efficiency [83]. However, there are many challenges that need to be conquered before the application of this concept in industrial production can be realized. For example, the carbon atom from CO<sub>2</sub> possesses a high oxidation state that requires large amounts of reducing power in microbes to efficiently remove the oxygen atoms for it to be used for hydrocarbon synthesis. When considering methane-based metabolism, the activation of the C–H bond in methane is a costly and extremely inefficient process, which indicates that a more feasible and efficient design is needed for methane utilization by microorganisms [84].

In many cases, the poor performance of the key enzymes involved in the different biosynthetic pathways is the major obstacle for improving product formation. This is, for example, the case for ADO, which has low catalytic activity even in its native host [85]. In addition, expression of heterologous membrane proteins is usually challenging owing to the distinct membrane structures between organisms [86,87], and poor expression is often observed for the members of the superfamily of cytochrome P450 enzymes such as OleT with one of potential reasons being cofactor (haem) deficiency [65]. Protein engineering of these enzymes can serve as a feasible strategy to enable further enhancement of final product levels through a

**Table 1.** Comparison of biofuel production from different organisms. FAAs, free fatty acids; OCFAs, odd chain fatty acids; FOHs, fatty alcohols; SCAs, short chain alkanes; FAEs, fatty acid ethyl esters; VLCFOHs, very-long-chain fatty alcohols; YNB, yeast nitrogen base; SD, synthetic defined.

microorganisms	product	titre (g l <sup>-1</sup> )	yield <sup>a</sup> (g g <sup>-1</sup> )	medium	cultivation condition	reference
<i>S. cerevisiae</i>	FAAs	1.0	0.05	MM <sup>b</sup>	shake flask	[19]
<i>E. coli</i>	FAAs	3.9	N.C. <sup>c</sup>	MK <sup>d</sup>	fed-batch	[77]
<i>E. coli</i>	FAAs	1.2	0.06	MM	shake flask	[53]
<i>Y. lipolytica</i>	OCFAs	0.75	N.C.	YNB	fed-batch	[78]
<i>S. cerevisiae</i>	FOHs	0.12	0.004	MM	shake flask	[19]
<i>S. cerevisiae</i>	VLCFOHs	0.084	0.0028	MM	shake flask	[52]
<i>S. cerevisiae</i>	FOHs	0.1	0.005	MM	shake flask	[16]
<i>E. coli</i>	FOHs	1.7	0.028	MM	batch	[79]
<i>S. cerevisiae</i>	1-alkenes	0.035	0.0011	MM	shake flask	[63]
<i>Y. lipolytica</i>	alkanes	0.023	N.C.	YNB	shake flask	[80]
<i>E. coli</i>	SCAs	0.58	N.C.	MR <sup>d</sup>	fed-batch	[81]
<i>S. cerevisiae</i>	FAEs	0.034	N.C.	SD medium	shake flask	[74]
<i>Y. lipolytica</i>	FAEs	0.14	N.C.	YNB	shake flask	[80]

<sup>a</sup>Yield was defined as carbon source conversion rate to biofuel production.

<sup>b</sup>MM, minimal medium.

<sup>c</sup>Unable to calculate because of the complex composition of the medium.

<sup>d</sup>Minimal medium with extra yeast extract.

more efficient metabolite conversion. Thus, with protein engineering strategies such as rational design based on known or simulated protein structures or random approaches through directed evolution, the cofactor, substrate or product specificity can be altered and benefit the overall improvement of enzyme activities. A successful example is the engineering of a type I FAS to generate more S/MCFAs in yeast, which offers the possibility to alter the fatty acid chain length and enables a larger diversity of biofuels produced by engineered strains [47]. However, gaining additional knowledge on enzymes and their structures as well as the development of advanced tools used for *in silico* analysis of proteins require more effort and attention in future studies.

With still relatively low yields and rates in biofuel production, the tolerance against toxic chemicals and fermentation processes needs to be further improved in order to achieve high TRY metrics to meet the requirements of commercial application [82,88]. Several successful attempts have shown that identifying respective transporters can be a feasible approach to releasing growth inhibition in microbial cell factories [59,63,72]. In addition, ALE towards toxic chemicals and inhibitory conditions has proven to be a promising method that can benefit the production of biofuels and other molecules, for instance resistance against high temperature or oxidative stress could be obtained by ALE [89,90]. If product formation can be linked to cell growth or survival, it is also possible to evolve strains for increased production. In this way, carotenoid production was improved by hydrogen peroxide-challenged adaptive evolution [91].

In addition, a number of biosensors have been developed to detect specific molecules and have been recently employed to facilitate the production of some valuable compounds, such as using a malonyl-CoA sensor to improve the production of 3-hydroxypropionic acid and fatty acids [77,92–94]. Nevertheless, the number of metabolites detectable as well as the properties (e.g. specificity) of existing biosensors need to be improved for additional applications [95]. Recently, the concept of synthetic product addiction facilitated by biosensors was proposed as a promising solution that would benefit high-yield bio-manufacturing [96]. Through linking production of the desired metabolite to the expression of non-conditionally essential genes, the product-addicted strains with biosynthetic capacity in the population will be selected without constraining the medium, thus providing production stability over many generations. With the help of such advanced biotechnology tools, efforts in the coming years will focus on how to improve the TRY metrics in order to meet the commercial requirement for lower value products and the production of higher value molecules using microorganisms.

**Data accessibility.** This article does not contain any additional data.

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## References

- Herrera S. 2006 Bonkers about biofuels. *Nat. Biotechnol.* **24**, 755–760. (doi:10.1038/nbt0706-755)
- Fortman JL, Chhabra S, Mukhopadhyay A, Chou H, Lee TS, Steen E, Keasling JD. 2008 Biofuel alternatives to ethanol: pumping the microbial well. *Trends Biotechnol.* **26**, 375–381. (doi:10.1016/j.tibtech.2008.03.008)



3. OECD/FAO. 2018 *OECD-FAO agricultural outlook 2018-2027*. Paris, France/Rome, Italy: OECD Publishing/Food and Agriculture Organization of the United Nations. (doi:10.1787/agr\_outlook-2018-en)
4. Aditiya HB, Mahlia TMI, Chong WT, Nur H, Sebayang AH. 2016 Second generation bioethanol production: a critical review. *Renew. Sustain. Energy Rev.* **66**, 631–653. (doi:10.1016/j.rser.2016.07.015)
5. Zhou YJ, Kerckhoven EJ, Nielsen J. 2018 Barriers and opportunities in bio-based production of hydrocarbons. *Nat. Energy* **3**, 925–935. (doi:10.1038/s41560-018-0197-x)
6. Zhang Y, Nielsen J, Liu Z. 2018 Metabolic engineering of *Saccharomyces cerevisiae* for production of fatty acid-derived hydrocarbons. *Biotechnol. Bioeng.* **115**, 2139–2147. (doi:10.1002/bit.26738)
7. Liao JC, Mi L, Pontrelli S, Luo S. 2016 Fuelling the future: microbial engineering for the production of sustainable biofuels. *Nat. Rev. Microbiol.* **14**, 288–304. (doi:10.1038/nrmicro.2016.32)
8. Goldemberg J. 2007 Ethanol for a sustainable energy future. *Science* **315**, 808–810. (doi:10.1126/science.1137013)
9. Bothast RJ, Schlicher MA. 2005 Biotechnological processes for conversion of corn into ethanol. *Appl. Microbiol. Biotechnol.* **67**, 19–25. (doi:10.1007/s00253-004-1819-8)
10. Adrio JL. 2017 Oleaginous yeasts: promising platforms for the production of oleochemicals and biofuels. *Biotechnol. Bioeng.* **114**, 1915–1920. (doi:10.1002/bit.26337)
11. Mattanovich D, Sauer M, Gasser B. 2014 Yeast biotechnology: teaching the old dog new tricks. *Microb. Cell Fact.* **13**, 34. (doi:10.1016/j.juro.2007.05.072)
12. Nielsen J, Larsson C, van Maris A, Pronk J. 2013 Metabolic engineering of yeast for production of fuels and chemicals. *Curr. Opin. Biotechnol.* **24**, 398–404. (doi:10.1016/j.copbio.2013.03.023)
13. Tehlivets O, Scheuringer K, Kohlwein SD. 2007 Fatty acid synthesis and elongation in yeast. *Biochim. Biophys. Acta Mol. Cell Biol. Lipids* **1771**, 255–270. (doi:10.1016/j.bbalip.2006.07.004)
14. Zhou YJ, Buijs NA, Zhu Z, Gómez DO, Boonsombuti A, Siewers V, Nielsen J. 2016 Harnessing yeast peroxisomes for biosynthesis of fatty-acid-derived biofuels and chemicals with relieved side-pathway competition. *J. Am. Chem. Soc.* **138**, 15 368–15 377. (doi:10.1021/jacs.6b07394)
15. Hassan SN, Sani YM, Abdul Aziz AR, Sulaiman NMN, Daud WMAW. 2015 Biogasoline: an out-of-the-box solution to the food-for-fuel and land-use competitions. *Energy Convers. Manag.* **89**, 349–367. (doi:10.1016/j.enconman.2014.09.050)
16. Runguphan W, Keasling JD. 2014 Metabolic engineering of *Saccharomyces cerevisiae* for production of fatty acid-derived biofuels and chemicals. *Metab. Eng.* **21**, 103–113. (doi:10.1016/j.ymben.2013.07.003)
17. Shi S, Valle-Rodríguez JO, Khoomrung S, Siewers V, Nielsen J. 2012 Functional expression and characterization of five wax ester synthases in *Saccharomyces cerevisiae* and their utility for biodiesel production. *Biotechnol. Biofuels* **5**, 7. (doi:10.1186/1754-6834-5-7)
18. Schirmer A, Rude MA, Li X, Popova E, Del Cardayre SB. 2010 Microbial biosynthesis of alkanes. *Science* **329**, 559–562. (doi:10.1126/science.1187936)
19. Zhou YJ, Buijs NA, Zhu Z, Qin J, Siewers V, Nielsen J. 2016 Production of fatty acid-derived oleochemicals and biofuels by synthetic yeast cell factories. *Nat. Commun.* **7**, 11709. (doi:10.1038/ncomms11709)
20. Schadeweg V, Boles E. 2016 N-Butanol production in *Saccharomyces cerevisiae* is limited by the availability of coenzyme A and cytosolic acetyl-CoA. *Biotechnol. Biofuels* **9**, 44. (doi:10.1186/s13068-016-0456-7)
21. Chen Y, Daviet L, Schalk M, Siewers V, Nielsen J. 2013 Establishing a platform cell factory through engineering of yeast acetyl-CoA metabolism. *Metab. Eng.* **15**, 48–54. (doi:10.1016/j.ymben.2012.11.002)
22. Ter Schure EG, Flikweert MT, Van Dijken JP, Pronk JT, Verrips CT. 1998 Pyruvate decarboxylase catalyzes decarboxylation of branched-chain 2-oxo acids but is not essential for fusel alcohol production by *Saccharomyces cerevisiae*. *Appl. Environ. Microbiol.* **64**, 1303–1307.
23. Van Maris AJA, Geertman JMA, Vermeulen A, Groothuizen MK, Winkler AA, Piper MDW, Van Dijken JP, Pronk JT. 2004 Directed evolution of pyruvate decarboxylase-negative *Saccharomyces cerevisiae*, yielding a C2-independent, glucose-tolerant, and pyruvate-hyperproducing yeast. *Appl. Environ. Microbiol.* **70**, 159–166. (doi:10.1128/AEM.70.1.159-166.2004)
24. Nielsen J. 2014 Synthetic biology for engineering acetyl coenzyme A metabolism in yeast. *MBio* **5**, e02153. (doi:10.1128/mBio.02153-14)
25. Oud B, Flores CL, Gancedo C, Zhang X, Trueheart J, Daran JM, Pronk JT, van Maris AJA. 2012 An internal deletion in MTH1 enables growth on glucose of pyruvate-decarboxylase negative, non-fermentative *Saccharomyces cerevisiae*. *Microb. Cell Fact.* **11**, 131. (doi:10.1186/1475-2859-11-131)
26. Chen Y, Zhang Y, Siewers V, Nielsen J. 2015 Ach1 is involved in shuttling mitochondrial acetyl units for cytosolic C2 provision in *Saccharomyces cerevisiae* lacking pyruvate decarboxylase. *FEMS Yeast Res.* **15**, 1–8. (doi:10.1093/femsyr/fov015)
27. Kozak BU *et al.* 2014 Engineering acetyl coenzyme a supply: functional expression of a bacterial pyruvate dehydrogenase complex in the cytosol of *Saccharomyces cerevisiae*. *MBio* **5**, e01696-14. (doi:10.1128/mBio.01696-14)
28. Zhang Y, Dai Z, Krivoruchko A, Chen Y, Siewers V, Nielsen J. 2015 Functional pyruvate formate lyase pathway expressed with two different electron donors in *Saccharomyces cerevisiae* at aerobic growth. *FEMS Yeast Res.* **15**, 24. (doi:10.1093/femsyr/fov024)
29. Yu T, Zhou YJ, Huang M, Liu Q, Pereira R, David F, Nielsen J. 2018 Reprogramming yeast metabolism from alcoholic fermentation to lipogenesis. *Cell* **174**, 1549–1558. (doi:10.1016/j.cell.2018.07.013)
30. Bergman A, Siewers V, Nielsen J, Chen Y. 2016 Functional expression and evaluation of heterologous phosphoketolases in *Saccharomyces cerevisiae*. *AMB Express* **6**, 115. (doi:10.1186/s13568-016-0290-0)
31. Kozak BU, van Rossum HM, Benjamin KR, Wu L, Daran JMG, Pronk JT, Van Maris AJA. 2014 Replacement of the *Saccharomyces cerevisiae* acetyl-CoA synthetases by alternative pathways for cytosolic acetyl-CoA synthesis. *Metab. Eng.* **21**, 46–59. (doi:10.1016/j.ymben.2013.11.005)
32. Ghosh A *et al.* 2016 13C metabolic flux analysis for systematic metabolic engineering of *S. cerevisiae* for overproduction of fatty acids. *Front. Bioeng. Biotechnol.* **4**, 76. (doi:10.3389/fbioe.2016.00076)
33. Shi S, Chen Y, Siewers V, Nielsen J. 2014 Improving production of malonyl coenzyme A-derived metabolites by abolishing Snf1-dependent regulation of Acc1. *MBio* **5**, e01130-14. (doi:10.1128/mBio.01130-14)
34. Chen X, Yang X, Shen Y, Hou J, Bao X. 2018 Screening phosphorylation site mutations in yeast acetyl-CoA carboxylase using malonyl-CoA sensor to improve malonyl-CoA-derived product. *Front. Microbiol.* **9**, 47. (doi:10.3389/fmicb.2018.00047)
35. Choi JW, Da Silva NA. 2014 Improving polyketide and fatty acid synthesis by engineering of the yeast acetyl-CoA carboxylase. *J. Biotechnol.* **187**, 56–59. (doi:10.1016/j.jbiotec.2014.07.430)
36. Nissen TL, Anderlund M, Nielsen J, Villadsen J, Kielland-Brandt MC. 2001 Expression of a cytoplasmic transhydrogenase in *Saccharomyces cerevisiae* results in formation of 2-oxoglutarate due to depletion of the NADPH pool. *Yeast* **18**, 19–32. (doi:10.1002/1097-0061(200101)18:1<19::AID-YEA650>3.0.CO;2-5)
37. Guo Z, Zhang L, Ding Z, Shi G. 2011 Minimization of glycerol synthesis in industrial ethanol yeast without influencing its fermentation performance. *Metab. Eng.* **13**, 49–59. (doi:10.1016/J.YMBEN.2010.11.003)
38. Moreira Dos Santos M, Raghevedran V, Kötter P, Olsson L, Nielsen J. 2004 Manipulation of malic enzyme in *Saccharomyces cerevisiae* for increasing NADPH production capacity aerobically in different cellular compartments. *Metab. Eng.* **6**, 352–363. (doi:10.1016/j.ymben.2004.06.002)
39. de Jong BW, Shi S, Siewers V, Nielsen J. 2014 Improved production of fatty acid ethyl esters in *Saccharomyces cerevisiae* through up-regulation of the ethanol degradation pathway and expression of the heterologous phosphoketolase pathway. *Microb. Cell Fact.* **13**, 39. (doi:10.1186/1475-2859-13-39)
40. Brody S, Oh C, Hoja U, Schweizer E. 1997 Mitochondrial acyl carrier protein is involved in lipoic acid synthesis in *Saccharomyces cerevisiae*. *FEBS Lett.* **408**, 217–220. (doi:10.1016/S0014-5793(97)00428-6)
41. Hiltunen JK, Schonauer MS, Autio KJ, Mittelmeier TM, Kastaniotis AJ, Dieckmann CL. 2009 Mitochondrial fatty acid synthesis type II: more than just fatty acids. *J. Biol. Chem.* **284**, 9011–9015. (doi:10.1074/jbc.R800068200)

42. Lomakin IB, Xiong Y, Steitz TA. 2007 The crystal structure of yeast fatty acid synthase, a cellular machine with eight active sites working together. *Cell* **129**, 319–332. (doi:10.1016/j.cell.2007.03.013)
43. Jenni S, Leibundgut M, Maier T, Ban N. 2006 Architecture of a fungal fatty acid synthase at 5 Å resolution. *Science* **311**, 1263–1267. (doi:10.1126/science.1123251)
44. Jenni S, Leibundgut M, Boehringer D, Frick C, Mikolásek B, Ban N. 2007 Structure of fungal fatty acid synthase and implications for iterative substrate shuttling. *Science* **316**, 254–261. (doi:10.1126/science.1138248)
45. Eriksen DT, Hamedirad M, Yuan Y, Zhao H. 2015 Orthogonal fatty acid biosynthetic pathway improves fatty acid ethyl ester production in *Saccharomyces cerevisiae*. *ACS Synth. Biol.* **4**, 808–814. (doi:10.1021/sb500319p)
46. Fernandez-Moya R, Leber C, Cardenas J, Da Silva NA. 2015 Functional replacement of the *Saccharomyces cerevisiae* fatty acid synthase with a bacterial type II system allows flexible product profiles. *Biotechnol. Bioeng.* **112**, 2618–2623. (doi:10.1002/bit.25679)
47. Zhu Z, Zhou YJ, Krivoruchko A, Grninger M, Zhao ZK, Nielsen J. 2017 Expanding the product portfolio of fungal type I fatty acid synthases. *Nat. Chem. Biol.* **13**, 344–345. (doi:10.1038/nChemBio.2301)
48. Gajewski J, Pavlovic R, Fischer M, Boles E, Grninger M. 2017 Engineering fungal de novo fatty acid synthesis for short chain fatty acid production. *Nat. Commun.* **8**, 14650. (doi:10.1038/ncomms14650)
49. Denic V, Weissman JS. 2007 A molecular caliper mechanism for determining very long-chain fatty acid length. *Cell* **130**, 663–677. (doi:10.1016/j.CELL.2007.06.031)
50. Kihara A. 2012 Very long-chain fatty acids: elongation, physiology and related disorders. *J. Biochem.* **152**, 387–395. (doi:10.1093/jb/mvs105)
51. Oh CS, Toke DA, Mandala S, Martin CE. 1997 ELO2 and ELO3, homologues of the *Saccharomyces cerevisiae* ELO1 gene, function in fatty acid elongation and are required for sphingolipid formation. *J. Biol. Chem.* **272**, 17 376–17 384. (doi:10.1074/jbc.272.28.17376)
52. Yu T, Zhou YJ, Wenning L, Liu Q, Krivoruchko A, Siewers V, Nielsen J, David F. 2017 Metabolic engineering of *Saccharomyces cerevisiae* for production of very long chain fatty acid-derived chemicals. *Nat. Commun.* **8**, 15587. (doi:10.1038/ncomms15587)
53. Steen EJ, Kang Y, Bokinsky G, Hu Z, Schirmer A, McClure A, Del Cardayre SB, Keasling JD. 2010 Microbial production of fatty-acid-derived fuels and chemicals from plant biomass. *Nature* **463**, 559–562. (doi:10.1038/nature08721)
54. Teerawanichpan P, Qiu X. 2010 Fatty acyl-CoA reductase and wax synthase from *Euglena gracilis* in the biosynthesis of medium-chain wax esters. *Lipids* **45**, 263–273. (doi:10.1007/s11745-010-3395-2)
55. Akhtar MK, Turner NJ, Jones PR. 2013 Carboxylic acid reductase is a versatile enzyme for the conversion of fatty acids into fuels and chemical commodities. *Proc. Natl Acad. Sci. USA* **110**, 87–92. (doi:10.1073/pnas.1216516110)
56. Kudo H, Nawa R, Hayashi Y, Arai M. 2016 Comparison of aldehyde-producing activities of cyanobacterial acyl-(acyl carrier protein) reductases. *Biotechnol. Biofuels* **9**, 234. (doi:10.1186/s13068-016-0644-5)
57. Metz JG. 2000 Purification of a jojoba embryo fatty acyl-coenzyme A reductase and expression of its cDNA in high erucic acid rapeseed. *Plant Physiol.* **122**, 635–644. (doi:10.1104/pp.122.3.635)
58. d'Espaux L *et al.* 2017 Engineering high-level production of fatty alcohols by *Saccharomyces cerevisiae* from lignocellulosic feedstocks. *Metab. Eng.* **42**, 115–125. (doi:10.1016/j.ymben.2017.06.004)
59. Hu Y, Zhu Z, Nielsen J, Siewers V. 2018 Heterologous transporter expression for improved fatty alcohol secretion in yeast. *Metab. Eng.* **45**, 51–58. (doi:10.1016/j.ymben.2017.11.008)
60. Henritzi S, Fischer M, Grninger M, Oreb M, Boles E. 2018 An engineered fatty acid synthase combined with a carboxylic acid reductase enables de novo production of 1-octanol in *Saccharomyces cerevisiae*. *Biotechnol. Biofuels* **11**, 150. (doi:10.1186/s13068-018-1149-1)
61. Foo JL, Susanto AV, Keasling JD, Leong SSS, Chang MW. 2017 Whole-cell biocatalytic and de novo production of alkanes from free fatty acids in *Saccharomyces cerevisiae*. *Biotechnol. Bioeng.* **114**, 232–237. (doi:10.1002/bit.25920)
62. Kang MK, Zhou YJ, Buijs NA, Nielsen J. 2017 Functional screening of aldehyde decarboxylases for long-chain alkane production by *Saccharomyces cerevisiae*. *Microb. Cell Fact.* **16**, 74. (doi:10.1186/s12934-017-0683-z)
63. Zhou YJ, Hu Y, Zhu Z, Siewers V, Nielsen J. 2018 Engineering 1-alkene biosynthesis and secretion by dynamic regulation in yeast. *ACS Synth. Biol.* **7**, 584–590. (doi:10.1021/acssynbio.7b00338)
64. Herman NA, Zhang W. 2016 Enzymes for fatty acid-based hydrocarbon biosynthesis. *Curr. Opin. Chem. Biol.* **35**, 22–28. (doi:10.1016/j.cbpa.2016.08.009)
65. Liu Y, Wang C, Yan J, Zhang W, Guan W, Lu X, Li S. 2014 Hydrogen peroxide-independent production of  $\alpha$ -alkenes by OleTJE P450 fatty acid decarboxylase. *Biotechnol. Biofuels* **7**, 28. (doi:10.1186/1754-6834-7-28)
66. Mendez-Perez D, Begemann MB, Pflieger BF. 2011 Modular synthase-encoding gene involved in  $\alpha$ -olefin biosynthesis in *Synechococcus* sp. strain pC 7002. *Appl. Environ. Microbiol.* **77**, 4264–4267. (doi:10.1128/AEM.00467-11)
67. Sorigué D *et al.* 2017 An algal photoenzyme converts fatty acids to hydrocarbons. *Science* **357**, 903–907. (doi:10.1126/science.aan6349)
68. Buijs NA, Zhou YJ, Siewers V, Nielsen J. 2015 Long-chain alkane production by the yeast *Saccharomyces cerevisiae*. *Biotechnol. Bioeng.* **112**, 1275–1279. (doi:10.1002/bit.25522)
69. Rottensteiner H, Theodoulou FL. 2006 The ins and outs of peroxisomes: co-ordination of membrane transport and peroxisomal metabolism. *Biochim. Biophys. Acta Mol. Cell Res.* **1763**, 1527–1540. (doi:10.1016/j.bbamcr.2006.08.012)
70. Zhu Z, Zhou YJ, Kang MK, Krivoruchko A, Buijs NA, Nielsen J. 2017 Enabling the synthesis of medium chain alkanes and 1-alkenes in yeast. *Metab. Eng.* **44**, 81–88. (doi:10.1016/j.ymben.2017.09.007)
71. Ling H, Chen B, Kang A, Lee JM, Chang MW. 2013 Transcriptome response to alkane biofuels in *Saccharomyces cerevisiae*: identification of efflux pumps involved in alkane tolerance. *Biotechnol. Biofuels* **6**, 95. (doi:10.1186/1754-6834-6-95)
72. Chen B, Ling H, Chang MW. 2013 Transporter engineering for improved tolerance against alkane biofuels in *Saccharomyces cerevisiae*. *Biotechnol. Biofuels* **6**, 21. (doi:10.1186/1754-6834-6-21)
73. Chen B, Lee D-Y, Chang MW. 2015 Combinatorial metabolic engineering of *Saccharomyces cerevisiae* for terminal alkene production. *Metab. Eng.* **31**, 53–61. (doi:10.1016/j.ymben.2015.06.009)
74. Shi S, Valle-Rodríguez JO, Siewers V, Nielsen J. 2014 Engineering of chromosomal wax ester synthase integrated *Saccharomyces cerevisiae* mutants for improved biosynthesis of fatty acid ethyl esters. *Biotechnol. Bioeng.* **111**, 1740–1747. (doi:10.1002/bit.25234)
75. Valle-Rodríguez JO, Shi S, Siewers V, Nielsen J. 2014 Metabolic engineering of *Saccharomyces cerevisiae* for production of fatty acid ethyl esters, an advanced biofuel, by eliminating non-essential fatty acid utilization pathways. *Appl. Energy* **115**, 226–232. (doi:10.1016/j.apenergy.2013.10.003)
76. Yu KO, Jung J, Kim SW, Park CH, Han SO. 2012 Synthesis of FAEs from glycerol in engineered *Saccharomyces cerevisiae* using endogenously produced ethanol by heterologous expression of an unspecific bacterial acyltransferase. *Biotechnol. Bioeng.* **109**, 110–115. (doi:10.1002/bit.23311)
77. Xu P, Li L, Zhang F, Stephanopoulos G, Koffas M. 2014 Improving fatty acids production by engineering dynamic pathway regulation and metabolic control. *Proc. Natl Acad. Sci. USA* **111**, 11 299–11 304. (doi:10.1073/pnas.1406401111)
78. Park Y-K, Dulermo T, Ledesma-Amaro R, Nicaud J-M. 2018 Optimization of odd chain fatty acid production by *Yarrowia lipolytica*. *Biotechnol. Biofuels* **11**, 158. (doi:10.1186/s13068-018-1154-4)
79. Liu A, Tan X, Yao L, Lu X. 2013 Fatty alcohol production in engineered *E. coli* expressing *Marinobacter* fatty acyl-CoA reductases. *Appl. Microbiol. Biotechnol.* **97**, 7061–7071. (doi:10.1007/s00253-013-5027-2)
80. Xu P, Qiao K, Ahn WS, Stephanopoulos G. 2016 Engineering *Yarrowia lipolytica* as a platform for synthesis of drop-in transportation fuels and oleochemicals. *Proc. Natl Acad. Sci. USA* **113**, 10 848–10 853. (doi:10.1073/pnas.1607295113)
81. Choi YJ, Lee SY. 2013 Microbial production of short-chain alkanes. *Nature* **502**, 571–574. (doi:10.1038/nature12536)
82. Peralta-Yahya PP, Zhang F, Del Cardayre SB, Keasling JD. 2012 Microbial engineering for the production of advanced biofuels. *Nature* **488**, 320–328. (doi:10.1038/nature11478)

83. Shinnar R. 1991 Future of chemical engineering. *Chem. Eng. Prog.* **87**, 80–85. (doi:10.1126/science.aag0804)
84. Balasubramanian R, Smith SM, Rawat S, Yatsunyk LA, Stemmler TL, Rosenzweig AC. 2010 Oxidation of methane by a biological dicopper centre. *Nature* **465**, 115–119. (doi:10.1038/nature08992)
85. Wang Q, Bao L, Jia C, Li M, Li J-J, Lu X. 2017 Identification of residues important for the activity of aldehyde-deformylating oxygenase through investigation into the structure-activity relationship. *BMC Biotechnol.* **17**, 31. (doi:10.1186/s12896-017-0351-8)
86. Newstead S, Kim H, von Heijne G, Iwata S, Drew D. 2007 High-throughput fluorescent-based optimization of eukaryotic membrane protein overexpression and purification in *Saccharomyces cerevisiae*. *Proc. Natl Acad. Sci. USA* **104**, 13 936–13 941. (doi:10.1073/pnas.0704546104)
87. Schlegel S, Löfblom J, Lee C, Hjelm A, Klepsch M, Strous M, Drew D, Slotboom DJ, de Gier J-W. 2012 Optimizing membrane protein overexpression in the *Escherichia coli* strain Lemo21(DE3). *J. Mol. Biol.* **423**, 648–659. (doi:10.1016/J.JMB.2012.07.019)
88. Rodionova MV *et al.* 2017 Biofuel production: challenges and opportunities. *Int. J. Hydrogen Energy* **42**, 8450–8461. (doi:10.1016/j.ijhydene.2016.11.125)
89. Caspeta L, Chen Y, Ghiaci P, Feizi A, Buskov S, Hallström BM, Petranovic D, Nielsen J. 2014 Altered sterol composition renders yeast thermotolerant. *Science* **346**, 75–78. (doi:10.1126/science.1258137)
90. Çakar ZP, Seker UOS, Tamerler C, Sonderegger M, Sauer U. 2005 Evolutionary engineering of multiple-stress resistant *Saccharomyces cerevisiae*. *FEMS Yeast Res.* **5**, 569–578. (doi:10.1016/j.femysr.2004.10.010)
91. Reyes LH, Gomez JM, Kao KC. 2014 Improving carotenoids production in yeast via adaptive laboratory evolution. *Metab. Eng.* **21**, 26–33. (doi:10.1016/j.ymben.2013.11.002)
92. Li S, Si T, Wang M, Zhao H. 2015 Development of a synthetic malonyl-coA sensor in *Saccharomyces cerevisiae* for intracellular metabolite monitoring and genetic screening. *ACS Synth. Biol.* **4**, 1308–1315. (doi:10.1021/acssynbio.5b00069)
93. David F, Nielsen J, Siewers V. 2016 Flux control at the malonyl-CoA node through hierarchical dynamic pathway regulation in *Saccharomyces cerevisiae*. *ACS Synth. Biol.* **5**, 224–233. (doi:10.1021/acssynbio.5b00161)
94. Li H, Chen W, Jin R, Jin J-M, Tang S-Y. 2017 Biosensor-aided high-throughput screening of hyper-producing cells for malonyl-CoA-derived products. *Microb. Cell Fact.* **16**, 187. (doi:10.1186/s12934-017-0794-6)
95. Adeniran A, Sherer M, Tyo KEJ. 2015 Yeast-based biosensors: design and applications. *FEMS Yeast Res.* **15**, 1–15. (doi:10.1111/1567-1364.12203)
96. Rugbjerg P, Sarup-Lytzen K, Nagy M, Sommer MOA. 2018 Synthetic addiction extends the productive life time of engineered *Escherichia coli* populations. *Proc. Natl Acad. Sci. USA* **115**, 2347–2352. (doi:10.1073/pnas.1718622115)