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2025, 9, 3838

Engineering a hypoxia-tolerant *Saccharomyces cerevisiae* for rapid ethanol production via co-utilization of glucose and acetic acid and redox-enhanced flocculation†

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Enhancing the robustness of microbial cell factories is essential for improving both first- and second-generation bioethanol production. During fermentation, *Saccharomyces cerevisiae* produces acetic acid as a by-product under certain conditions, which inhibits cellular functions and reduces fermentation efficiency. Additionally, pretreatment of lignocellulosic biomass releases acetic acid, further exacerbating fermentation stress toward the yeast. Hypoxic fermentation, combined with metabolic engineering, offers an alternative strategy to mitigate these challenges. To address this, we used CRISPR-Cas9 gene editing to sequentially delete NADH-dependent glycerol-3-phosphate dehydrogenase 1 (*GPD1*), cytosolic aldehyde dehydrogenase (*ALD6*), and mitochondrial external NADH dehydrogenase isoforms (*NDE1* and *NDE2*), while integrating an empty plasmid into the *LEU2* locus to generate control strains C1 to C5. Notably, strain C5 (*GPD1Δ ALD6Δ NDE1Δ NDE2Δ*), exhibited a 150% increase in the fermentation rate compared to strain C1 when fermenting a minimal medium containing 10% glucose and 0.4% acetic acid under hypoxic conditions. To further enhance acetic acid utilization and ethanol production, we integrated a plasmid containing acetylating acetaldehyde dehydrogenase from *Salmonella enterica* (*SeEutE*) into the *LEU2* locus, generating EutE strains E1 to E5. Strain E5 (*GPD1Δ ALD6Δ NDE1Δ NDE2Δ* [*SeEutE*]) exhibited a 200% increase in fermentation rate compared to strain C5, with 75% ethanol-induced flocculation. Strain E5 consumed approximately 25% of the supplemented acetic acid and achieved near-theoretical ethanol yields from the total consumed glucose and acetic acid. Furthermore, strain E5 exhibited a 9% improvement in the fermentation rate under hypoxic conditions compared to hyperoxic conditions. These enhancements together represent an overall improvement of more than 343% compared to the parent strain. Thus, by integrating quadruple deletion (*GPD1Δ ALD6Δ NDE1Δ NDE2Δ*) with the heterologous expression of *SeEutE* integration, we introduce a novel strategy to construct a hypoxia and acetate tolerant *S. cerevisiae* strain. This engineered strain achieves rapid, redox-balanced fermentation and ethanol-induced flocculation, offering a significant advance by overcoming limitations in glucose fermentation rate, redox imbalance, and weak acetate tolerance.

Received 18th February 2025
Accepted 27th May 2025

DOI: 10.1039/d5se00258c

rsc.li/sustainable-energy

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† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d5se00258c>

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1. Introduction

Addressing climate change, which has a significant impact on the biosphere and ecosystems, is a global priority. The United Nations Sustainable Development Goals (SDGs) emphasize the need to balance environmental, economic, and social requirements, making the sustainable production of biofuels both essential and highly valued.¹ *Saccharomyces cerevisiae*, a model microbial cell factory, plays a key role in first-generation bioethanol production, contributing to an annual yield exceeding 100 billion liters.² However, acetic acid is produced as a by-product during bioethanol fermentation, inhibiting cell growth and reducing both fermentation rate and ethanol yield, particularly under conditions that lead to elevated acetic acid



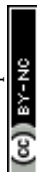
accumulation. While excessive acetic acid can be inhibitory, its toxicity varies depending on strain tolerance and environmental conditions.^{3,4} High concentrations of acetic acid induce apoptosis in yeast cells through mechanisms involving mitochondrial dysfunction, reactive oxygen species (ROS) accumulation, and cytochrome c release. The target of rapamycin (TOR) pathway plays a regulatory role in stress responses and metabolic adaptation to acetic acid toxicity, influencing processes such as amino acid and nucleotide biosynthesis, protein turnover, carbohydrate metabolism, cell wall and membrane integrity, signaling, and cell cycle regulation.⁵

In second-generation bioethanol production, *S. cerevisiae* strains face additional challenges, including reduced robustness and efficiency in handling inhibitors released from lignocellulosic biomass, such as furfural, 5-hydroxy methyl furfural (5-HMF), and acetic acid. Acetic acid, commonly released during the deacylation of hemicellulose and lignin in lignocellulosic hydrolysates, exacerbates these challenges and poses a significant obstacle to bioethanol production from lignocellulosic biomass.⁶ The co-utilization of acetic acid as an electron acceptor by *S. cerevisiae* was first demonstrated using the heterologous expression of *E. coli* acetylating acetaldehyde dehydrogenase (*EcMhpF*) along with double deletion of both NADH-dependent glycerol-3-phosphate dehydrogenases (*GPD1* and *GPD2*) under anaerobic conditions. This strategy successfully prevented glycerol biosynthesis, enabled acetic acid co-

utilization, and enhanced ethanol production. However, the engineered strain exhibited a 2.6-fold reduction in fermentation speed and showed reduced tolerance to higher sugar concentrations.⁷ In a subsequent report, recovery of tolerance was achieved through evolutionary engineering.⁸ The conversion of acetic acid to ethanol requires the transport of acetic acid into the cell, its conversion to acetyl CoA by ATP-dependent acetyl-CoA synthase (*ACS1*, *ACS2*), and subsequent reductions to acetaldehyde and ethanol by *EcMhpF* and alcohol dehydrogenase isoforms (*ADHs*), respectively.⁷ A key limitation for acetic acid co-utilization in this pathway is the availability of NADH during glucose fermentation. Heterologous expression of NADPH-dependent alcohol dehydrogenase from *Entamoeba histolytica* (*EhADH*) in industrial diploid *S. cerevisiae* strains increased NADH levels, supporting the activity of two copies of acetylating acetaldehyde dehydrogenase from *Bifidobacterium adolescentis* (*BaAdhE*). This modification replaced *GPD1* and *GPD2* with *BaAdhE*, thereby preventing glycerol biosynthesis and increasing the co-utilization of acetate.⁹ Replacing the NADP⁺-dependent 6-phosphogluconate dehydrogenase (*GND*) with a NAD⁺-dependent variant, combined with substituting glycerol biosynthesis with *E. coli* *EutE*, improves acetic acid co-utilization with glucose and enhances ethanol production, though it results in a lower fermentation rate.¹⁰ Similarly, heterologous replacement of the native NADH-dependent *ScGPD1* with an archaeal NADPH-preferring gene (*GpsA*) and



Fig. 1 Diagram of the fermentative metabolic pathways in *S. cerevisiae* with a focus on glucose and acetic acid. Native pathways and native enzymes are written in black. NAD-dependent glycerol-3-phosphate dehydrogenase 1 (*GPD1*), cytosolic aldehyde dehydrogenase 6 (*ALD6*), and mitochondrial external NADH dehydrogenases 1 (*NDE1*) and 2 (*NDE2*), which were sequentially deleted in this study, are shown in the yellow ovals and marked with red crosses. The heterologous enzyme, NADH-dependent putative acetaldehyde dehydrogenase from *Salmonella enterica* (*SeEutE*), is shown in red within a yellow oval. The acetate futile cycle is highlighted in the blue square. "Created in BioRender. Mohammed, M. (2025) <https://BioRender.com/snzfdcy>".



heterologous replacement of the native *ScGPD2* with *EcEutE*, along with the deletion of cytosolic aldehyde dehydrogenase (*ALD6*), successfully blocked glycerol biosynthesis and increased ethanol production and acetic acid co-utilization under hyperosmotic conditions.¹¹ However, this also led to a significant reduction in fermentation rates.

Xylose, the second most abundant sugar in lignocellulosic biomass after glucose, provides additional NADH through the cofactor recycling between xylose reductase (XR) and xylitol dehydrogenase (XDH). The redox imbalance between XR and XDH has been utilized to improve acetic acid co-utilization with xylose, confirming that NADH deficiency constrains acetic acid co-utilization. Co-fermentation of xylose and acetic acid also helps reduce xylitol accumulation.¹² Optimization of the acetic acid reduction pathway, including expression of three copies of codon-optimized *EcAdhE* and *Salmonella enterica* acetyl-CoA synthetase (*SeACS*), has significantly improved anaerobic xylose fermentation and acetate co-utilization.¹³ Under aerobic conditions without glucose, when the target product does not involve oxidoreductase reactions, acetic acid was co-consumed with xylose at a 1:4 ratio, underscoring the importance of NADH availability and the inhibitory effect of glucose on acetic acid co-consumption.¹⁴

Despite these advances, most existing redox engineering strategies present significant trade-offs—such as reduced fermentation speed, impaired osmoadaptation, or diminished strain robustness—especially under hypoxic or industrially relevant conditions. Designs based on full deletion of *GPD1/GPD2*, overexpression of NADH-consuming enzymes, or anaerobic xylose–acetate co-fermentation often result in slow ethanol production, excessive glycerol accumulation, and poor tolerance to process-related stresses. Furthermore, ethanol-induced flocculation—an important trait for stress protection and biomass recovery—has not been previously integrated into acetate-utilizing yeast strains.

In this study, we addressed these limitations by developing a modular, stepwise metabolic engineering strategy. This involved deleting mitochondrial external NADH dehydrogenases (*NDE1* and *NDE2*)—enzymes typically active under normoxic or hyperoxic conditions—and introducing redox balancing *via* heterologous expression of *Salmonella enterica* acetylating acetaldehyde dehydrogenase (*SeEutE*), as illustrated in Fig. 1. To our knowledge, this is the first report to link ethanol-induced flocculation with enhanced acetate co-utilization.

Specifically, we constructed a hypoxia- and acetate-tolerant *S. cerevisiae* strain through sequential deletion of *GPD1*, *ALD6*, *NDE1*, and *NDE2* (Fig. 1), which significantly accelerated glucose fermentation under hypoxia. Subsequent integration of *SeEutE* improved NADH recycling, promoted acetate assimilation, enhanced ethanol production, and triggered flocculation in response to ethanol accumulation. This strategy led to more than a threefold increase in fermentation rate, near-theoretical ethanol yield, and superior performance under hypoxic *versus* hyperoxic conditions. Collectively, this work establishes a robust platform for efficient glucose–acetate co-fermentation and scalable bioethanol production under oxygen-limited industrial settings.

2. Materials and methods

2.1 Media components

YPD medium (10 g yeast extract, 20 g peptone, and 20 g glucose per liter) was used for routine cultivation and preservation of yeast strains. Agar was added at 1.8% to solidify the medium for agar plates. For positive selection in the Clustered Regularly Interspaced Short Palindromic Repeats (CRISPR) system, YPD medium was supplemented with geneticin (G418), resulting in YPD_{G418} medium. Positive colonies with chromosomal integration at the *LEU2* locus were selected on Yeast Nitrogen Base (YNB) medium without amino acids (Invitrogen™), supplemented with 6.7 g L⁻¹ YNB, 20 g L⁻¹ glucose and 20 mg L⁻¹ histidine, designated as YNB_{D20}.¹⁵ The composition of YNB is listed in Table S1.†

For fermentations with glucose as the sole carbon source, YNB was supplemented with 100 g L⁻¹ glucose, 4 g L⁻¹ acetic acid, and 0.2 g L⁻¹ histidine, with the pH adjusted to 5 using NaOH (referred to as YNB_{D100-AC4}). To prevent interference from degradation products during autoclaving, all liquid media used in this study were filter-sterilized by TPP vacuum filtration (rapid-Filtermax, 0.2 µm PES, Switzerland). *Escherichia coli* NEB® 10-beta (New England Biolabs) was cultivated in Luria-Bertani (LB) medium, containing 1% tryptone, 1% sodium chloride (NaCl), and 0.5% yeast extract. Ampicillin (150 µg mL⁻¹) was added to the LB medium to select for *E. coli* carrying ampicillin-resistant plasmids, while kanamycin (75 µg mL⁻¹) was used for selecting *E. coli* harboring the pCas9 multiplex plasmid, as previously described.^{16,17}

2.2 Primers, cassettes, and plasmids constructions

All primers were designed based on the *S. cerevisiae* S288C sequence from the *Saccharomyces* Genome Database (Table S2†). Primers were synthesized by FASMAC, Japan.

2.2.1 Construction of multiplex pCas-gRNA-CRISPR plasmid. The multiplex pCas-gRNA-CRISPR plasmids were constructed following the protocol developed by Ryan *et al.*,¹⁸ with modifications as reported previously.^{4,16,17} Briefly, the online tool <https://chopchop.cbu.uib.no/> was used to design guide gRNA sequences with high knockout efficiency, located 20 bp upstream of the protospacer-adjacent motif (PAM). These sequences were used to design the primers for knocking out *ALD6*, *NDE1*, and *NDE2* (Table S2†). To avoid common PCR-induced mutations and the difficulties of amplifying large plasmids (pCas, 8743 bp), a two-step PCR process was employed to generate universal scaffolds. In the first round of PCR, two separate fragments were synthesized: one containing the upstream region and the other the downstream region of the gRNA scaffold. These PCR products were purified using a 2% agarose gel (50–800 bp, 75 min.) and a FastGene Gel/PCR extraction kit (Nippon Genetics, Tokyo, Japan) to obtain high-purity DNA fragments. These fragments were then used as templates for a second round of PCR to overlap with the 20-nucleotide guide sequence, constructing the complete gRNA scaffold. KOD-plus neo and pCas-scaffold forward and reverse primers were used with 6 pg of each template DNA segment.



The DNA scaffold was cleaved with *SmaI/PstI* and cloned into a multiplex plasmid cut with the same enzymes, yielding a new multiplex pCas-gRNA plasmid. This process was repeated to construct multiplex pCas-gRNA plasmids targeting *ALD6*, *NDE1*, and *NDE2* (Table S3†). The plasmids were introduced into NEB 10-beta *E. coli* (New England BioLabs, Tokyo, Japan) using the heat shock method, following the manufacturer's instructions. After overnight incubation, the plasmids were extracted using the QIAprep Spin miniprep kit (QIAGEN, Hilden, Germany) and the sequences were confirmed by sequencing.

2.2.2 Construction of *SeEutE*-YEpM4 plasmid. The YEpM4 plasmid was recovered from *E. coli* stocks.¹⁵ It was cleaved with *Sall* and dephosphorylated using shrimp alkaline phosphatase (Takara, Japan). The glyceraldehyde-3-phosphate dehydrogenase promoter (*TDH3p*) and a mutated *DIT1* terminator (*DIT1*_{d22t})¹⁹ were amplified by PCR from the *TDH3-DIT1*_{d22t}-*URA3* plasmid¹⁶ using primers (Table S2†). *TDH3p* was cleaved with *Sall/NotI*, while *DIT1*_{d22t} was cleaved with *BamHI/XhoI* and *NotI/XhoI*. Codon-optimized *SeEutE* was synthesized (Twist Bioscience, Japan), amplified by PCR, and cleaved with *NotI/BamHI*. A one-step ligation of *TDH3p*, *SeEutE*, and *DIT1*_{d22t} into the dephosphorylated *Sall* YEpM4 plasmid produced the *SeEutE*-YEpM4 plasmid (Table S3†). The complete sequence of the cassette is provided (Table S4†). The control plasmid, YEpM4-TD, was constructed by cloning *TDH3p* and *DIT1*_{d22t} into the dephosphorylated *Sall* YEpM4 plasmid. The plasmids were then introduced into *E. coli*, recovered, and confirmed by PCR and sequencing. The DNA concentration was measured using a NanoPhotometer UV/Vis spectrophotometer (IMPLEN, Germany). All plasmids and DNA fragments were stored on ice during the experiments and at −20 °C for long term storage.

2.3 Engineering the recombinant strains

The engineering strategy is summarized in Fig. S1.† The haploid D452-2 strain (*MATα leu2 his3 ura3 can1*) served as the parental strain (D).²⁰ The *URA3* locus in the D strain was restored in a previous report,¹⁶ resulting in the base strain B1 (Table 1). Strain B2, with *GPD1* deleted,⁴ was used for sequential deletions of *ALD6*, *NDE1*, and *NDE2*, following a method reported by Ryan *et al.*¹⁸ with a few modifications.^{16,17}

A repair cassette composed of *TDH3p-DIT1*_{d22t} was generated by PCR from the YEpM4-TD plasmid, incorporating homologous regions (50–60 bp) flanking the *ALD6* promoter and terminator. The repair cassette and a CRISPR-Cas9 multiplex plasmid were introduced into competent cells using a lithium acetate transformation protocol detailed previously.¹⁷ Transformants were selected on YPD_{G418} medium, and colonies were screened for successful *ALD6* deletions *via* PCR verification. At least four positive colonies were re-cultured on YPD_{G418} plates to eliminate false positives, and their reproducibility was confirmed through fermentation experiments. Strain B3 was propagated in YPD medium with shaking to remove the CRISPR-Cas9 plasmid and was subsequently used to generate strains B4, C3, and E3 (Table 1).

Similar procedures were applied to delete *NDE1* and *NDE2*, resulting in strains B4 and B5. The plasmid YEpM4-TD was

Table 1 Native (D), base (B), control (C), and EutE (E) strains used in this study, and their relative genotypes

Strains	Relevant genotype	Reference
D452-2 (D)	<i>MATα leu2 his3 ura3 can1</i>	20
D _{URA3} (B1)	D, <i>URA3::TDH3</i> promoter and <i>DIT1</i> _{d22t} terminator	16
B2	B1, <i>ΔGPD1::TDH3p-DIT1</i> _{d22t}	4
B3	B2, <i>ΔALD6::TDH3p-DIT1</i> _{d22t}	This study
B4	B3, <i>ΔNDE1::TDH3p-DIT1</i> _{d22t}	This study
B5	B4, <i>ΔNDE2::TDH3p-DIT1</i> _{d22t}	This study
C1	B1, <i>LEU2::TDH3p-DIT1</i> _{d22t}	This study
C2	B2, <i>LEU2::TDH3p-DIT1</i> _{d22t}	This study
C3	B3, <i>LEU2::TDH3p-DIT1</i> _{d22t}	This study
C4	B4, <i>LEU2::TDH3p-DIT1</i> _{d22t}	This study
C5	B5, <i>LEU2::TDH3p-DIT1</i> _{d22t}	This study
E1	B1, <i>LEU2::TDH3p-SeEutE-DIT1</i> _{d22t}	This study
E2	B2, <i>LEU2::TDH3p-SeEutE-DIT1</i> _{d22t}	This study
E3	B3, <i>LEU2::TDH3p-SeEutE-DIT1</i> _{d22t}	This study
E4	B4, <i>LEU2::TDH3p-SeEutE-DIT1</i> _{d22t}	This study
E5	B5, <i>LEU2::TDH3p-SeEutE-DIT1</i> _{d22t}	This study

linearized with *ClaI* and introduced into B1–B5 strains to generate control strains (C1–C5) (Fig. S1†). Concurrently, the linearized YEpM4-*SeEutE* plasmid was integrated into the same locus in B1–B5 strains, generating EutE-expressing strains (E1–E5) (Fig. S1†).

Transformants were selected on YNB_{D20} agar plates, and the first 12 colonies were transferred to and re-cultivated on fresh YNB_{D20} plates to prevent parental strain overgrowth. Four colonies were randomly selected to verify fermentation reproducibility, and all strains and their genotypes are listed in Table 1.

2.4 Pre-culture and fermentation conditions

Under sterile conditions, strains were pre-cultured in 10 mL of YPD medium in 50 mL TPP® TubeSpin bioreactor tubes (Switzerland) with orbital shaking at 180 rpm (EYELA, LTI-601SD, Japan) for 18 hours at an approximately 45° sitting angle during the orbital shaking. The cells were harvested by centrifugation at 5800 g, washed twice with sterile Milli-Q water, and resuspended to an OD₆₀₀ of 2 in 50 mL of YNB_{D100-4AC} (pH 5) in a 100 mL Erlenmeyer flask under hypoxic conditions [volume of yeast culture (*V_c*)/volume of flask (*V_f*) at a ratio of 50/100] to initiate the fermentation process. Fermentation was conducted at 150 rpm with 1 mL samples taken at intervals to measure growth and other fermentation characteristics.

2.5 Procedures of fermentation analysis

Fermentation analyses were conducted following the method described by Khattab and Watanabe (2021). Briefly, 100 μL samples were diluted with 900 μL of Milli-Q water, mixed, and centrifuged at 15 300 g for 5 minutes using a Tomy centrifuge (Japan). The supernatants were filtered through a hydrophilic FILTSTAR PTFE 0.45 μm syringe filter (Hawach Scientific, China) and transferred to 2 mL HPLC glass vials using a 1 mL syringe. An autosampler injected the samples into an HPLC system equipped with an Aminex HPX-87H column and



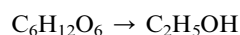
a refractive index detector. The mobile phase was 5 mM H₂SO₄ in aqueous solution and analyses were run at a flow rate of 0.6 mL min⁻¹. Concentrations of glucose, acetic acid, glycerol, ethanol, and other intermediate metabolites such as acetoin, 2,3-BDO, succinic acid, pyruvate, and acetaldehyde were quantified using high-grade standards.

2.6 Calculations of fermentation efficiency

Fermentation efficiency was calculated based on the theoretical yield (TY) using the following equations:

TY = molar mass of product/molar mass of substrate (glucose and/or acetic acid)

TY of bioethanol from glucose:



$$1 \text{ g} \rightarrow 0.51 \text{ g}$$

TY of bioethanol from acetic acid:



$$1 \text{ g} \rightarrow 0.77 \text{ g}$$

TY of bioethanol from glucose (g L⁻¹) = [initial concentration (IC) of glucose (g L⁻¹) – residual concentration (RC) of glucose (g L⁻¹)] × 0.51

TY of bioethanol from acetic acid (g L⁻¹) = [IC of acetic acid (g L⁻¹) – RC of acetic acid (g L⁻¹)] × 0.77

2.7 Enzyme activity

Cell pellets were harvested for enzyme activity measurements at 15 h of fermentation by centrifugation at 3 000 g for 5 min at 4 °C. The cell lysate was prepared as previously described:²¹ cell pellets were resuspended in 100 mM potassium phosphate (pH 7.5) containing 1 mM MgCl₂ and 10 mM 2-mercaptoethanol, followed by vortexing with approximately 400 mg of glass beads. The lysing process involved six cycles of vigorous shaking, each followed by 30 seconds of chilling on ice. The cell-free extract (CFE) was separated by centrifugation at 20 400 g for 5 min at 4 °C and further clarified by additional centrifugation. Protein concentrations were determined using the Pierce BCA Protein Assay Kit (Thermo Fisher Scientific) using bovine serum albumin standards and measurements were made at an absorbance of 562 nm on an Infinite M200 PRO plate reader (Tecan, Switzerland).

The specific activity of *SeEutE* was assayed as described by Eftance *et al.*,²² in a solution of 50 mM citrate buffer (pH 6.0), 100 μM zinc acetate, 240 μM NADH, and 140 μM acetyl-CoA. A 10 μL aliquot of the crude extract was used to monitor the decrease in NADH concentration over time at 340 nm. Background rates were subtracted in the calculations. Enzyme activity was expressed as μmol of NADH oxidized per min per mg of CFE.

2.8 Flocculation assay

Flocculation was measured as previously described,²³ with modifications. YNB_{D100-AC4} cell pellets were harvested after 21 h of fermentation by centrifugation at 2 300 g for 5 min at 25 °C and washed twice with 0.1 M EDTA to remove any residue that could interfere with flocculation. The washed cells were resuspended in a de-flocculating buffer (50 mM sodium acetate and 0.1 M EDTA), and the optical density was adjusted to an OD₆₀₀ of approximately 2.0 (initial optical density, A). 10 mL of this suspension was centrifuged, and the pellet was washed with deionized water and flocculating buffer (50 mM sodium acetate and 0.1% CaCl₂). The cells were then resuspended in 10 mL of flocculating buffer, vortexed for 5 min, and allowed to rest for 5 min. The OD₆₀₀ of the supernatant (final optical density, B) was then measured. Flocculation percentages were calculated as follows:

$$\text{Flocculation percentage} = (1 - B/A) \times 100\%$$

2.9 Varying the fermentation conditions from hyperoxic to hypoxic in strains E4 and E5

Oxygen availability was varied by adjusting the V_c relative to the V_f in 100 mL Erlenmeyer flasks. Fermentation of YNB_{D100-AC4} was performed with V_c/V_f ratios of 10/100 (hyperoxic conditions), 30/100, and 70/100 (hypoxic conditions).

2.10 Ethanol-induced engineered yeast flocculation

To evaluate ethanol-induced flocculation, the standard fermentation procedures were used with 50 mL of YNB_{D100-AC4} medium and ethanol was added to the medium to a final concentration of 3%, and 4% after three hours fermentation. Flocculation was observed four hours post-addition and compared with a control experiment without ethanol supplementation.

2.11 Analysis of cellular NADH/NAD⁺ ratio

Intracellular NADH and NAD⁺ levels were determined photometrically using the NAD⁺/NADH Assay Kit-WST (N509, Dojindo, Japan), following the manufacturer's instructions with modifications to the initial sample preparation step. At 9 hours of fermentation in YNB_{D100-AC4} medium, cells corresponding to 10 mL at OD₆₀₀ = 6 were harvested and lysed as described in Section 2.7. The supernatant was subsequently processed using Amicon Ultra centrifugal filters (10 kDa molecular weight cutoff) to isolate low-molecular-weight metabolites, including NADH and NAD⁺. A 10% aliquot of the filtrate was used for the quantitative determination of total NADH and NAD⁺ concentrations.

2.12 Statistical analysis

A completely randomized design (CRD) was used with three replications for all fermentation characteristics, except for the enzyme activity of *SeEutE*, which had four replications. Analysis of variance (ANOVA) was performed using the Proc Mixed



procedure in the SAS software package version 9.2.²⁴ Means were compared using Duncan's multiple range test at a 5% level of probability.²⁵

3. Results and discussions

3.1 Selection basis for target gene deletion in this study

Studies on anaerobic fermentation using engineered *S. cerevisiae* with double deletions of *GPD1* and *GPD2* have successfully demonstrated the oxidation of excess NADH through the introduction of acetylating acetaldehyde dehydrogenase, facilitating the co-utilization of acetic acid with glucose.^{7–9} While this strategy increases NADH availability for enzymes like *SeEutE*, it also eliminates G-3P, which is crucial for glycerophospholipid formation. This disruption can affect various cellular processes, including cell wall integrity, signaling, and regulation.^{26,27} G-3P is normally dephosphorylated to glycerol, serving as an osmoadaptive regulator,²⁸ and its absence exacerbates the inhibitory effects of acetic acid.⁴ The double substitution of *ScGPD1* with *GpsA* and *ScGPD2* with *EcEutE*, coupled with the deletion of *ScALD6*, effectively halted glycerol biosynthesis and enabled the co-utilization of acetic acid and glucose under osmotic fermentation conditions. However, this modification resulted in a significant decline in the fermentation rate.^{10,11} *GPD1* is well known for being expressed under aerobic, and *GPD2* under anaerobic conditions.²⁸ Furthermore, gene expression profiles under anaerobic conditions reveal that 140 genes have transcript levels more than threefold higher than under aerobic conditions, while 219 genes are more expressed under aerobic conditions.²⁹ These metabolic shifts may limit the fermentation rate, indicating a need for alternative metabolic engineering strategies tailored to hypoxic conditions. Indeed, under aerobic conditions, the exclusive co-conversion of acetic acid and xylose into triacetic acid lactone or vitamin A demonstrated that a 1 : 4 ratio between acetic acid and xylose was optimal. This serves as a strong example of proper design where ATP requirements, reducing equivalents, substrate co-utilization, and feeding patterns were optimized.¹⁴

In the presence of oxygen which serves as the terminal electron acceptor in the electron transport chain, *NDE1* and *NDE2* play a crucial role in oxidizing cytosolic NADH, facilitating ATP generation (Fig. 1). It has been observed that *NDE1* compensates for the loss of *NDE2* (*NDE2Δ*). Additionally, the deletion of both *NDE1* and *NDE2* (*NDE1Δ NDE2Δ*) results in more than a sevenfold increase in glycerol biosynthesis.³⁰ Removing these genes also enhances the enzyme activity of mitochondrial glycerol-3-phosphate dehydrogenase (*GUT2*) and decreases its apparent K_m for G-3P. G-3P is the substrate of *GUT2*, and also a product of *GPD* isoforms.³¹ Deleting *GPD1* in the base strain D452-2 (B2) reduced glycerol biosynthesis by 79% under hypoxic conditions.⁴ Based on these findings, we selected *GPD1* for deletion while leaving *GPD2* to balance the need for osmoadaptation with the provision of NADH for *SeEutE*. Deleting *GPD1*, *NDE1*, and *NDE2* genes decreases oxidative phosphorylation and respiration, generating hypoxia-tolerant cells. Hypoxia is known for spatial reorganization and accelerating fermentation rate under certain conditions.^{32,33}

Additionally, during co-utilization with glucose, acetic acid biosynthesis can form a futile cycle, despite its essential role in the pyruvate dehydrogenase (PDH) bypass pathway for supplying acetyl-CoA (Fig. 1). To disrupt this futile cycle, *ALD6* was selected for deletion in this study. Previous studies have shown that *ALD6Δ* can be effectively combined with the xylose-fermenting yeast strain SR8 to enhance acetic acid co-utilization.^{12,13} Also, deletion of *ScALD6* improved anaerobic co-fermentation of acetic acid with 1 M glucose in strain IMX901, which had its native NADH-dependent *ScGPD1* replaced with the archaeal NADPH-preferring *GpsA*, and *ScGPD2* substituted by *EcEutE*.¹¹ Therefore, combining *NDE1Δ*, *NDE2Δ* with *GPD1Δ* could potentially provide additional NADH for the heterologous expression of *SeEutE*. Based on this rationale, *GPD1*, *ALD6*, *NDE1*, and *NDE2* were selected for sequential deletion in this study (Fig. S1†). The sequential deletion of *GPD1*, *ALD6*, *NDE1*, and *NDE2* was implemented not only to redirect NADH toward acetate reduction but also to establish a controlled redox and respiratory rewiring under hypoxic-like conditions. Specifically, the deletion of *NDE1* and *NDE2*, which encode the primary mitochondrial external NADH dehydrogenases functional under normoxic or hyperoxic environments, restricts cytosolic NADH oxidation *via* the electron transport chain, thereby simulating an O_2 -independent, hypoxia-mimicking state. This genetic intervention is expected to suppress oxidative phosphorylation and ATP-coupled respiration, thereby promoting fermentative metabolism to compensate for the energetic deficit, particularly in the presence of acetic acid, while simultaneously generating a strong redox driving force for acetate reduction. Notably, hypoxia-induced metabolic adaptation in *S. cerevisiae* has been shown to involve the spatial reorganization of glycolytic enzymes, such as enolase, into cytoplasmic foci—a phenomenon that enhances glucose conversion to pyruvate and oxaloacetate, ultimately boosting fermentative carbon flux under conditions of restricted respiration.³² Furthermore, the deletion of *GPD1* reduces glycerol biosynthesis and impacts osmoadaptation by limiting glycerol-3-phosphate availability, potentially increasing

Table 2 Specific enzyme activities of a putative acetaldehyde dehydrogenase (*SeEutE*) in the CFE^a of control (C1–C5) and EutE strains (E1–E5)

Strain name	Specific activity ^b
C1 (native)	ND ^c E
E1 (EutE)	0.063 ± 0.01 ^D
C2 (<i>GPD1Δ</i>)	ND ^c E
E2 (<i>GPD1Δ</i> + EutE)	0.146 ± 0.02 ^B
C3 (<i>GPD1Δ</i> + <i>ALD6Δ</i>)	ND ^c E
E3 (<i>GPD1Δ</i> + <i>ALD6Δ</i> + EutE)	0.074 ± 0.01 ^C
C4 (<i>GPD1Δ</i> + <i>ALD6Δ</i> + <i>NDE1Δ</i>)	ND ^c E
E4 (<i>GPD1Δ</i> + <i>ALD6Δ</i> + <i>NDE1Δ</i> + EutE)	0.072 ± 0.01 ^{C&D}
C5 (<i>GPD1Δ</i> + <i>ALD6Δ</i> + <i>NDE1Δ</i> + <i>NDE2Δ</i>)	ND ^c E
E5 (<i>GPD1Δ</i> + <i>ALD6Δ</i> + <i>NDE1Δ</i> + <i>NDE2Δ</i> + EutE)	0.158 ± 0.02 ^A

^a CFE were obtained from cells at 15 h of fermentation in YNB_{D100-Ac4} medium, pH 5. Error values represent standard deviation from the mean ($n = 4$). ^b Indicate $\mu\text{mole}/\text{min mg}$ CFE. ^c Not detected. Statistical analysis indicated that means sharing the same letter (A, B, C, D) are not significantly different from each other ($P < 0.05$).



membrane stress under acetate exposure. Deletion of *ALD6*, a key enzyme in the acetate-producing PDH bypass, prevents futile cycling of acetaldehyde to acetate, thereby conserving reducing equivalents and reinforcing the redox shift. Together, these deletions not only redirect metabolic flux but also alleviate oxidative and membrane-associated stress, enhancing robustness under hypoxic fermentation.

3.2 Expression of *SeEutE* in the parent strain and its effects on fermentation

The codon-optimized *SeEutE* was successfully expressed in *S. cerevisiae* for the first time, achieving significant specific activity

of 0.063 ± 0.01 U in the CFE for strain E1 ($P < 0.05$), while the control strain C1 showed no detectable activity (Table 2). Previously, *SeEutE* expressed in *E. coli* demonstrated reversible activities, with acetaldehyde dehydrogenase activity at 68.1 ± 2 U and acetyl-CoA reductase activity at 49.2 ± 3 U for the purified protein.³⁴

Incorporation of the *SeEutE* gene into the parent strain (E1) led to an 11% increase in the glucose consumption rate (1.99 ± 0.05 g (L⁻¹ h⁻¹) vs. 1.8 ± 0.04 g (L⁻¹ h⁻¹) in Fig. 2A) and a 4% increase in ethanol yield ratio (0.47 ± 0.01 g g⁻¹ vs. 0.45 ± 0.01 g g⁻¹ in Fig. 2B) compared to the control strain C1. Meanwhile, acetic acid and glycerol production decreased by 6% (AcAP of 0.67 ± 0.02 g L⁻¹ vs. 0.71 ± 0.01 g L⁻¹ in Table 3) and 19% (GP of 0.8 ± 0.02 g L⁻¹ vs. 0.99 ± 0.02 g L⁻¹ in Table 3), respectively. Cell growth increased slightly by 7% (CG of 9 ± 0.72 OD vs. 8.4 ± 0.40 OD) (Table 3, and Fig. 2 and 3). Duncan's multiple range test at the $P < 0.05$ level revealed a significant difference between E1 and C1 only in glycerol production, with no significant change observed in other parameters (Table S5B†).

Although the integration of *SeEutE* resulted in the improvement of ethanol production, competition with native NADH-dependent pathways with enzymes such as *GPD* and *NDE* limited the extent of the enhancements, as expected. In our recent study, the heterologous expression of other NADH oxidases, such as water-forming *LlNoxE* from *Lactococcus lactis*, in the same base strain B1 under hypoxic conditions led to an increase in acetic acid biosynthesis.⁴ This outcome underscores the advantage of using *SeEutE* over *LlNoxE* for reducing acetic acid production.

3.3 Integration of *SeEutE* with *GPD1Δ* in strain E2

To assess the effect of *GPD1* deletion on the activity of *SeEutE*, we engineered strain E2. This strain exhibited a significantly larger specific activity than E1 in the CFE ($P < 0.05$); activity of E2 (0.146 ± 0.02 U) is 2.3 times higher than that of strain E1 (0.063 ± 0.01 U) (Table 2). The differences in specific activities of *SeEutE* between strains remain unclear. This enhancement in activity led to a further decrease in acetic acid concentration in E2 (AcAP of 0.42 ± 0.12 g L⁻¹) by 37% compared to E1 (AcAP of 0.67 g L⁻¹ ± 0.02 g L⁻¹) (Table 3).

Although the decrease in AcAP was not significant at $P < 0.05$ level as well as the growth (Table S5B†), the ethanol production rate in E2 is significantly higher than C2 by 40% (0.91 ± 0.03 g (L⁻¹ h⁻¹ vs. 0.65 ± 0.01 g (L⁻¹ h⁻¹, Fig. 2A). Ethanol yield ratio in E2 was enhanced by 9% compared to C2 (0.47 ± 0.01 vs. 0.43 ± 0.01 g g⁻¹, Fig. 2B), although no further improvement was seen over E1 (Fig. 2).

3.4 Abolishing the acetic acid futile cycle in strain E3 and beyond

Acetic acid biosynthesis serves as an intermediate step in the formation of acetyl-CoA via the PDH bypass pathway, which involves pyruvate decarboxylase *PDC* isoforms 1, 5, and 6, *ALD6*, *ACS1*, and *ACS2*.^{35–37} The *S. cerevisiae* genome contains five alternative isoforms for *ALD* associated with acetic acid production. Acetic acid production in the C3 strain (*GPD1Δ*



Fig. 2 Fermentation characteristics in YNBD_{100-AC4} medium after 24 hours by engineered *S. cerevisiae* strains under hypoxic fermentation conditions ($V_c/V_f = 50/100$). (A) Glucose consumption and ethanol production rates shown in g (L⁻¹ h⁻¹). Blue bars represent glucose consumption and red bars represent ethanol production. (B) Ethanol yield ratios for the same strains, expressed as g g⁻¹ (orange bars), cmol cmol⁻¹ (black bars), and mol mol⁻¹ (green bars). Error bars represent the standard deviation from the mean ($n = 3$).



Table 3 A comparison of fermentation characteristics on YNB_{D100-AC4} medium using strains generated in this study at 24 h

Strains	Fermentation parameters							
	GC ^a (g L ⁻¹)	AcAC ^b (g L ⁻¹)	AcAP ^c (g L ⁻¹)	EP ^d (g L ⁻¹)	EP/T ^e (%)	GP ^f (g L ⁻¹)	TPR ^g (g ³ /g ⁵)	CG ^h (OD)
C1 (native)	43.5 ± 1.99	0.0 ± 0.00	0.71 ± 0.01	19.6 ± 0.18	88.0 ± 0.1	0.99 ± 0.01	0.49 ± 0.01	8.4 ± 0.40
E1 (EutE)	48.3 ± 0.86	0.0 ± 0.00	0.67 ± 0.10	22.7 ± 0.3	92.2 ± 0.2	0.80 ± 0.02	0.50 ± 0.01	9.0 ± 0.72
C2 (GPD1Δ)	35.99 ± 0.77	0.0 ± 0.00	0.67 ± 0.02	15.6 ± 0.22	85.0 ± 0.2	0.19 ± 0.01	0.46 ± 0.10	8.3 ± 0.23
E2 (GPD1Δ + EutE)	46.7 ± 0.19	0.0 ± 0.00	0.42 ± 0.12	21.9 ± 0.7	91.9 ± 0.1	0.19 ± 0.01	0.48 ± 0.01	8.7 ± 0.94
C3 (GPD1Δ + ALD6Δ)	29.8 ± 1.10	0.0 ± 0.00	0.02 ± 0.04	12.7 ± 0.6	83.0 ± 0.2	0.13 ± 0.01	0.43 ± 0.19	7.9 ± 0.23
E3 (GPD1Δ + ALD6Δ + EutE)	27.5 ± 0.13	0.26 ± 0.02	0.00 ± 0.00	13.1 ± 0.02	92.9 ± 0.2	0.16 ± 0.01	0.48 ± 0.01	9.0 ± 0.72
C4 (GPD1Δ + ALD6Δ + NDE1Δ)	31.0 ± 0.34	0.00 ± 0.00	0.33 ± 0.03	13.6 ± 0.47	86.0 ± 0.2	0.26 ± 0.01	0.46 ± 0.01	8.1 ± 0.23
E4 (GPD1Δ + ALD6Δ + NDE1Δ + EutE)	98.0 ± 0.25	1.04 ± 0.10	0.00 ± 0.0	50.0 ± 0.5	98.5 ± 0.3	0.50 ± 0.01	0.50 ± 0.01	10.2 ⁱ ± 0.35
C5 (GPD1Δ + ALD6Δ + NDE1Δ + NDE2Δ)	45.6 ± 0.70	0.0 ± 0.00	0.03 ± 0.01	21.2 ± 0.07	91.4 ± 0.18	0.46 ± 0.02	0.48 ± 0.01	8.6 ^j ± 0.20
E5 (GPD1Δ + ALD6Δ + NDE1Δ + NDE2Δ + EutE)	98.7 ± 0.96	0.83 ± 0.07	0.00 ± 0.0	49.8 ± 0.34	97.7 ± 0.3	0.54 ± 0.04	0.51 ± 0.0	10.7 ^j ± 0.23

^a Glucose consumption. ^b Acetic acid production. ^c Ethanol production. ^d Ethanol product/theoretical. ^e Glycerol production. ^f Total products ratio ($\frac{g^{products}}{g^{substrates}}$). ^g Cell growth. Values represent the averages ± SD, $n = 3$. ^h CG was measured after de-flocculation using a buffer containing 50 mM sodium acetate and 0.1 M EDTA.



Fig. 3 Intracellular NADH/NAD⁺ ratio in engineered *S. cerevisiae* strains E4 and E5 compared to control strain C1. The ratio was measured after 9 hours of fermentation in YNB_{D100-AC4} medium under hypoxic conditions ($V_c/V_f = 50/100$). Data represent mean ± SD of triplicate.

ALD6Δ) was very low (AcAP of 0.02 ± 0.01 g L⁻¹, Table 3), confirming the predominance of *ALD6* in acetic acid production.

Additionally, strain E3 (*GPD1Δ ALD6Δ [SeEutE]*) further consumed acetic acid by 0.26 g L⁻¹ (AcAC, Table 3). These significant changes in the AcAP ($P < 0.05$, Table S5B†) clearly demonstrated that the deletion of *ALD6Δ*, in combination with incorporation of *SeEutE* can eliminate the futile cycle of acetic acid production during the co-utilization of acetic acid and glucose (Table 3 and Fig. S2 and S3†).

In strain E3, bioethanol production significantly increased to 92.9% of the theoretical yield, compared to 88% in strain C1 (EP/T, Table 3). However, glucose consumption in E3 was significantly reduced at $P < 0.05$ to 27.5 ± 0.13 g L⁻¹ compared to C1 (43.5 ± 1.99) at 24 h (GC, Table 3, Fig. S2 and S3, S5B†). Thus, while the elimination of the acetic acid futile cycle is promising, it is not sufficient on its own for effective co-utilization.

3.5 Triple and quadruple deletion of *GPD1*, *ALD6*, and *NDE1* (and *NDE2*) with *SeEutE* expression: achieving ethanol-induced flocculation and fermentation robustness

3.5.1 Effects of *GPD1*, *ALD6*, and *NDE1* triple deletion and subsequent integration of *SeEutE*. Building on previous engineering steps, we implemented the triple deletion of *GPD1*, *ALD6*, and *NDE1* in strain C4 and expressed *SeEutE* in strain E4 to enhance fermentation performance and acetic acid co-utilization. Notably, all fermentation parameters measured in YNB_{D100-AC4} medium were significantly improved in strain E4 compared to C4 ($P < 0.05$, Table S5†). The cell growth (OD₆₀₀) of E4 increased by 21% compared to C1 at 24 h (CG of 10.2 ± 0.35 vs. 8.4 ± 0.40 OD, Table 3).

Strain E4 consumed 98.0 ± 0.25 g L⁻¹ of glucose within 24 h, compared to 31.0 ± 0.34 g L⁻¹ consumed by strain C4 and 43.5 ± 1.99 g L⁻¹ by strain C1 (GC, Table 3), indicating the increased vigour of strain E4. The glucose consumption rates for C1, C4 and E4 were 1.8 ± 0.04 g (L⁻¹ h⁻¹), 1.3 ± 0.01 g (L⁻¹ h⁻¹), and 4.1 ± 0.01 g (L⁻¹ h⁻¹) (Fig. 2A), respectively, with corresponding

bioethanol production rates of $0.82 \pm 0.01 \text{ g (L}^{-1} \text{ h}^{-1})$, $0.57 \pm 0.01 \text{ g (L}^{-1} \text{ h}^{-1})$, and $2.08 \pm 0.02 \text{ g (L}^{-1} \text{ h}^{-1})$ (Fig. 2A). Furthermore, the ethanol product/theoretical of E4 reached $98.5 \pm 0.3\%$, compared to $86.0 \pm 0.2\%$ for C4 and $88.0 \pm 0.1\%$ for C1 (EP/T, Table 3). Glycerol production was $0.99 \pm 0.01 \text{ g L}^{-1}$ in C1, $0.26 \pm 0.01 \text{ g L}^{-1}$ in C4, and $0.50 \pm 0.01 \text{ g L}^{-1}$ in E4 (GP, Table 3). Acetic acid consumption by strain E4 increased to $1.04 \pm 0.10 \text{ g L}^{-1}$ compared to $0.26 \pm 0.02 \text{ g L}^{-1}$ by strain E3 (AcAC, Table 3 and Fig. S2 and S3†).

3.5.2 Effects of quadruple deletion of *GPD1*, *ALD6*, *NDE1*, and *NDE2*, and subsequent integration of *SeEutE*. Fig. 2 illustrates the markedly enhanced fermentation performance of strain E5, including a more than twofold increase in both glucose consumption and ethanol production rates under hypoxic conditions, compared to control strains. In line with the promising results from strain E4, strain E5 demonstrated enhanced fermentation robustness. Strain E5 exhibited 24% greater growth compared to C5 (CG, 10.7 ± 0.23 vs. 8.6 ± 0.2 , Table 3). Notably, C5 consumed glucose much faster than C1, completing glucose consumption within 48 h, while C1 required 72 h (Fig. S2A†). E5 consumed $98.7 \pm 0.98 \text{ g L}^{-1}$ of glucose within 24 h, whereas C5 consumed $45.6 \pm 0.7 \text{ g L}^{-1}$ (GC, Table 3). This raises the glucose consumption rate in E5 to $4.1 \pm 0.04 \text{ g (L}^{-1} \text{ h}^{-1})$, compared to $1.9 \pm 0.02 \text{ g (L}^{-1} \text{ h}^{-1})$ in C5 and $1.8 \pm 0.04 \text{ g (L}^{-1} \text{ h}^{-1})$ in C1 (Fig. 2A).

Ethanol production by C5 at 48 h reached $46.3 \pm 0.2 \text{ g L}^{-1}$, compared to $36.3 \pm 0.2 \text{ g L}^{-1}$ produced by strain C1 (Fig. S2D†). Strain E5 produced ethanol even more efficiently, reaching $49.8 \pm 0.34 \text{ g L}^{-1}$ in just 24 h (EP, Table 3). The ethanol production rate in E5 rose to $2.07 \pm 0.01 \text{ g (L}^{-1} \text{ h}^{-1})$, compared to $0.88 \pm 0.0 \text{ g (L}^{-1} \text{ h}^{-1})$ in C5 and $0.82 \pm 0.01 \text{ g (L}^{-1} \text{ h}^{-1})$ in C1 (Fig. 2B). C5 did not consume any acetic acid, whereas E5 consumed $0.83 \pm 0.07 \text{ g L}^{-1}$ (AcAC, Table 3). Glycerol production in C5 and E5 increased to $0.46 \pm 0.02 \text{ g L}^{-1}$ and $0.54 \pm 0.04 \text{ g L}^{-1}$, respectively, compared to $0.26 \pm 0.01 \text{ g L}^{-1}$ in C4 and $0.50 \pm 0.01 \text{ g L}^{-1}$ in E4 (GP, Table 3). These enhancements in the fermentation characteristics of E5 compared to C1 and C5 were further validated through statistical analysis ($P < 0.05$; Table S5†). The increased acetic acid consumption and glycerol production in E5 compared to C5 suggest a balance between *ScGPD2* and *SeEutE* for NADH oxidation. This balance enabled the generation of G-3P and glycerol, along with the co-utilization of glucose and acetic acid. This balance between *ScGPD2* and *SeEutE* was demonstrated for the first time. In earlier studies, both *ScGPD1* and *ScGPD2* were deleted to fully direct NADH toward acetylating acetaldehyde dehydrogenase.^{7,9} However, this approach inhibited G-3P and glycerol biosynthesis, leading to deficiencies in cell wall viability, integrity, signaling, and regulation.^{26,27} These factors may explain the sluggish fermentation rate observed in previous studies that deleted both *GPD1* and *GPD2* under anaerobic conditions. Our alternative approach with hypoxic conditions, which retains *GPD2*, has produced both superior fermentation rates and efficiencies compared to previous reports.

The intracellular NADH/NAD⁺ ratio is a critical indicator of redox balance during fermentation. As shown in Fig. 3, the control strain C1 exhibited a high NADH/NAD⁺ ratio 0.59 ± 0.03 ,

indicating significant cytosolic NADH accumulation under hypoxic conditions. In contrast, both engineered strains E4 and E5 showed markedly reduced NADH/NAD⁺ ratios 0.3 ± 0.002 and 0.25 ± 0.003 , respectively (Fig. 3). This significant decline ($P < 0.05$) demonstrates the effectiveness of the introduced modifications in rebalancing the redox state.

Specifically, the deletions of *GPD1* and *ALD6* reduce competing NADH-consuming pathways (glycerol and acetate formation), while deletion of *NDE1* and *NDE2* restricts mitochondrial NADH oxidation, creating a more reduced cytosolic environment. Introduction of *SeEutE*, provides an alternative NADH oxidation route directly linked to ethanol production. Together, these modifications redirect excess NADH toward ethanol synthesis, improving fermentation efficiency and reducing the need for byproduct formation. The enhanced redox homeostasis in E5, as reflected by the lowest NADH/NAD⁺ ratio, correlates well with its superior ethanol productivity (Fig. 2). Taken together, the engineered acetate co-utilization pathway in strain E5 combines (i) enhanced NADH availability via *GPD1*, *NDE1*, and *NDE2* deletions, (ii) disruption of the acetic acid futile cycle through *ALD6* deletion, and (iii) redox-balanced reduction of acetyl-CoA to ethanol via *SeEutE*. This configuration enables robust acetate assimilation under hypoxic conditions, overcoming the limitations of previous systems.

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3.5.3 Ethanol-induced flocculation and stress tolerance of strains E4, C5, and E5. Interestingly, the triple deletion of *GPD1*, *ALD6*, and *NDE1*, combined with *SeEutE* expression, was associated with a significant onset of flocculation in E4 approximately 15 h into the fermentation under hypoxic conditions in YNB_{D100-AC4} medium (Fig. 4). In contrast, the control strain C4 exhibited a phenotype similar to C1, with no notable flocculation (Fig. 4). The cell wall flocculation assay showed that flocculation in E4 was 7.6 times greater than that in C1 ($50.8 \pm 3.6\%$ vs. $6.7 \pm 5.1\%$, Fig. 4), suggesting significant changes in cell wall hydrophobicity in strain E4, as has been observed previously.⁶

In *S. cerevisiae*, flocculin genes (*FLO1*, *FLO5*, *FLO8*, *FLO9*, *FLO10*, and *FLO11*) play key roles in enhancing cell wall



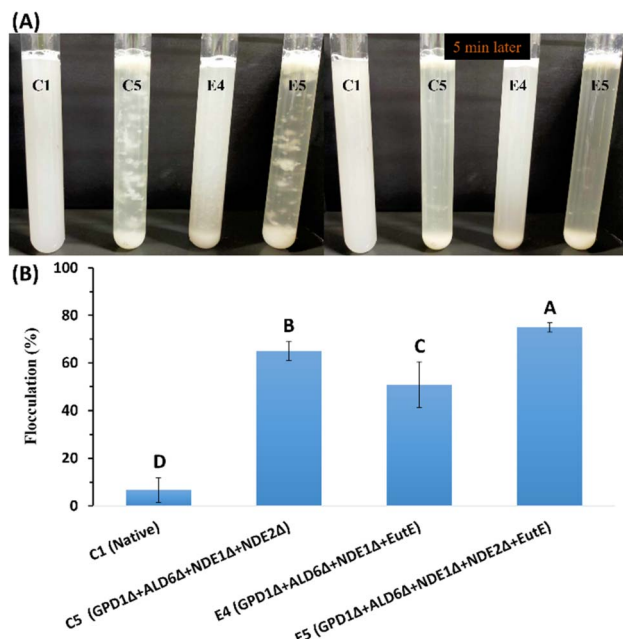


Fig. 4 Flocculation of strains C5, E4, and E5 compared to the non-flocculating original strain C1 after 21 hours of fermentation in YNB_{D100-AC4} medium under hypoxic conditions (V_c/V_f ratio of 50/100). (A) The upper left picture was taken immediately after transfer from the fermentation flasks and mixing, while the upper right image was taken after 5 minutes of gravity settling. (B) The percentage of flocculated cells in strains C1, C5, E4, and E5 (see section 2.8 for details). Error bars represent the standard deviations from the mean of ($n = 3$, SD). Statistical analysis indicated that means sharing the same letter are not significantly different from each other ($P < 0.05$).

hydrophobicity, flocculation, and cell adhesion. These traits facilitate yeast cell separation, which is advantageous in industrial processes.^{38–43} Moreover, flocculation provides protection under stress conditions by shielding cells from inhibitors.^{6,44} A transcriptional repressor and activator MOT3 has regulates cell wall hydrophobicity by modulating the expression of the hydrophobic cell wall protein YPG1. This regulation enhances resistance to inhibitory chemical compounds (ICCs), even in the absence of acetic acid co-utilization.⁶

With the quadruple deletions of *GPD1*, *ALD6*, *NDE1*, and *NDE2* in C5, flocculation reached $65.0 \pm 4.1\%$ after 21 h of fermentation (Fig. 4). Glucose consumption in C5 was enhanced by 150% compared to C1 (Fig. 2A). In E5, the expression of *SeEutE* alongside these quadruple deletions further increased the flocculation to $75.0 \pm 2.0\%$ (Fig. 4), with larger flocs observed compared to those formed by E4 (Fig. S4†). E5 exhibited over 200% higher glucose consumption and ethanol production rates than C5, as well as slightly faster fermentation compared to E4 (Fig. S2 and S3†). We hypothesize that the more pronounced flocculation in E5 is associated with enhanced protection against multiple stresses, including ethanol, as previously reported.^{40,43,44}

In E5, flocculation began 15 h into the fermentation, after ethanol production reached $30.16 \pm 0.47 \text{ g L}^{-1}$. It was reported that the exposure to 3% ethanol induces 70% flocculation in



Fig. 5 Induction of flocculation in strain E5 by adding two different concentrations of ethanol (3% and 4%) after three hours of fermentation in YNB_{D100-AC4} under hypoxic conditions (V_c/V_f ratio of 50/100). Flocculation assay of strain E5 at the same time points after adding 3% and 4% ethanol, compared to normal fermentation without ethanol addition. Error bars represent the standard deviations from the mean ($n = 3$, SD). Means sharing the same letter are not significantly different from each other ($P < 0.05$).

strains overexpressing *FLO1* and 28% in strains overexpressing *FLO5*.⁴⁰ Ethanol can trigger flocculation through controlled *FLO1* expression regulated by the trehalose-6-phosphate synthase 1 (*TPS1*) promoter, which responds to ethanol accumulation during fermentation. Ethanol-induced flocculation has been shown to enhance biomass recovery.⁴³ These observations led us to hypothesize that ethanol promoted flocculation. To test this, we supplemented the medium with 3% and 4% ethanol after 3 h of initial fermentation using strain E5. As a result, the flocculation increased by 80% and 210%, respectively, compared to the control without ethanol supplementation (Fig. 5). These results support our hypothesis and are consistent with previous reports demonstrating that 3% ethanol induces flocculation under certain conditions as reported elsewhere.^{40,43}

Moreover, the integration and overexpression of biofilm-forming genes (*FLO5*, *FLO8*, and *FLO10*) in *S. cerevisiae* have been shown to improve biofilm formation. Engineered strains 1308-FLO5 and 1308-FLO10 demonstrated significant increases in adhesion and ethanol production compared to the wild type. These strains also reduced broth turbidity and enhanced membrane separation efficiency, increasing ethanol flux through the membrane by 36.3% during biofilm-immobilized fermentation.⁴²

Interestingly, our engineering approach did not involve the overexpression of any *FLO* genes by design, suggesting that ethanol-induced flocculation in E5 may be driven by an indirect mechanism, warranting further investigation. In future studies, we will conduct molecular characterization of key regulatory genes associated with accelerated fermentation rates, as well as flocculation-related genes such as *FLO1*, *FLO5*, *FLO8*, *FLO10*, and *YPG1*. We will also examine their transcriptional responses to ethanol exposure to elucidate the molecular basis of this phenotype.

3.5.4. Oxygen limitation and fermentation efficiency. Previous research showed that *FLO1* overexpression induced



flocculation and reduced the growth rate by fourfold, with flocculent cells upregulating strictly anaerobic and starvation-related genes.⁴⁴ These findings, along with the triple deletion of oxidative phosphorylation-related genes (*GPD1*, *NDE1*, and *NDE2*) prompted us to investigate the effects of oxygen limitation on E4 and E5. Interestingly, E5 demonstrated a preference for hypoxic conditions over hyperoxic conditions, with a 9% increase in fermentation speed at V_c/V_f ratios of 30/100 and 70/100 compared to 10/100 (Fig. 6). Under hyperoxic conditions at a V_c/V_f ratio of 10/100, E5 consumed $88.05 \pm 0.26 \text{ g L}^{-1}$ of glucose and $1.0 \pm 0.02 \text{ g L}^{-1}$ of acetic acid. At a V_c/V_f ratio of 70/100, glucose consumption increased to $96.6 \pm 0.5 \text{ g L}^{-1}$ and acetic acid consumption decreased to $0.6 \pm 0.02 \text{ g L}^{-1}$, resulting in $48.9 \pm 0.23 \text{ g L}^{-1}$ ethanol production within 21 h. Statistical analysis ($P < 0.05$) confirmed that hyperoxic conditions are significantly different from hypoxic conditions (Table S6†).



Fig. 6 Time course fermentation in YNBD₁₀₀-AC₄ medium at pH 5 by strain E5 under hyperoxic (V_c/V_f ratio of 10/100) and hypoxic (V_c/V_f ratios of 30/100 and 70/100) conditions. (A) Glucose consumption; (B) glycerol production; (C) acetic acid concentration; and (D) ethanol production. Error bars represent the standard deviations from the mean ($n = 3$, SD).

Acetic acid consumption was lower under the 70/100 V_c/V_f ratio conditions, but glycerol production increased to $1.3 \pm 0.01 \text{ g L}^{-1}$, approximately 2.4 times higher than under hyperoxic conditions (Fig. 6). This suggests further activation of GPD2 activity under near-anaerobic conditions (V_c/V_f 70/100), consistent with previous studies highlighting the role of GPD2 under anaerobic conditions.⁴⁵ On the other hand, glycerol production slightly increased in E4 at a 70/100 V_c/V_f ratio, suggesting that NDE2 may play a role in sharing NADH oxidation with GPD2 (Fig. S3B and Table S7†). The increased glycerol production in E5 under stronger hypoxia may reflect a redox-balancing response in which GPD2 compensates for impaired mitochondrial respiration due to the deletion of NDE1 and NDE2. As *SeEutE* and GPD2 both consume NADH, their co-function under hypoxic stress likely contributes to redox homeostasis. This cooperative NADH oxidation enables E5 to maintain higher fermentation efficiency and cell viability, whereas complete GPD deletions in previous studies often led to fermentation defects. This indicates a distinct metabolic response between strains E4 and E5 under varying oxygen levels, with E5 showing a more robust adaptation to hypoxic conditions. These findings underscore the importance of preserving GPD2-mediated redox flexibility in engineered strains, as its activity appears to complement *SeEutE* under hypoxic stress to sustain efficient fermentation.

3.6 Cross-study comparison and application context

To further contextualize the potential of acetate co-utilization and address strain performance relative to the literature, we compared the sugar consumption rates, ethanol production rates, and yield efficiencies of our current strain (E5) with previously engineered *S. cerevisiae* strains (Table 4). A defining feature of our strategy is the induction of O_2 -independent hypoxia *via* triple deletion of *GPD1*, *NDE1*, and *NDE2*, along with *ALD6* deletion. The sequential deletion of *GPD1*, *ALD6*, *NDE1*, and *NDE2*, combined with *SeEutE* expression, significantly accelerated glucose fermentation under hypoxic conditions in this study, resulting in a 3.42-fold increase in fermentation rate compared to the control strain C1 (Fig. S2 and S3†). Strain E5 reached 97.7% of the theoretical ethanol yield within 24 h when fermenting 100 g L^{-1} glucose supplemented with 4 g L^{-1} acetate, representing a significant improvement over the 88% yield observed in C1. Ethanol-induced flocculation was also triggered in E5 when ethanol concentration reached approximately 3%.

We respectfully highlight the study by Papapetridis *et al.*,¹¹ in which a *GPD1Δ GPD2Δ ALD6Δ* strain co-expressing *AfGpsA* and *EcEutE* co-utilized 4.86 g L^{-1} acetate with 180 g L^{-1} glucose under anaerobic conditions, achieving a 92.3% ethanol yield over 61 h. In contrast, our hypoxia-tolerant strain E5—retaining GPD2 to preserve osmoadaptation—achieved a higher yield in less than half the time (Table 4). These findings suggest that future integration of *AfGpsA* into E5 may further enhance acetate assimilation efficiency. Acetate co-utilization has also been validated in glucose-xylose co-fermentation systems. For example, integration of three copies of *EcAdhE* and *SeACS*

Table 4 Comparative literature data on acetic acid co-conversion to ethanol versus the performance of the engineered strain E5 in this study

Carbons consumption rate ($\text{g L}^{-1} \text{ h}^{-1}$)	Ethanol production rate ($\text{g L}^{-1} \text{ h}^{-1}$)	Acetic acid co-utilized (g L^{-1})	Fermentation time (h)	Efficiency of ethanol yield (%) ^a	Key genetic modifications	Media and conditions	Reference
0.53 ^b	0.48 ^b	0.50 ^b	38 ^b	89.2 ^b	GPD1Δ, GPD2Δ + <i>EcMhpF</i>	20 g glucose + 2 g acetic acid; anaerobic	7
0.99	0.46	5.30	120	87.8	GPD1Δ, GPD2Δ, <i>EhADH1</i> , <i>ACS2</i> , <i>ZWF1</i> , 2 × <i>BcAdhE</i>	117 g glucose + 8 g acetic acid; anaerobic	9
0.95 ^b	0.45 ^b	0.80 ^b	23 ^b	90.9 ^b	GPD1Δ, GPD2Δ, <i>ALD6Δ</i> , <i>GND2Δ</i> , <i>GND1Δ</i> + <i>MfGNDa</i> , <i>EcEutE</i>	20 g glucose + 3 g acetic acid; anaerobic	10
3.03	1.45	4.86 ^b	61 ^b	92.3 ^b	GPD1Δ, GPD2Δ, <i>ALD6Δ</i> + <i>AfGpsA</i> , <i>EcEutE</i>	180 g glucose + 3 g acetic acid (maintained $>1.5 \text{ g L}^{-1}$); anaerobic	11
1.22 ^b	0.5*	1.70 ^b	88	82.3 ^b	<i>ALD6Δ</i> , <i>XR</i> , <i>XDH</i> , <i>KK</i> (2 copies), adapted strain + <i>EcEutE</i>	80 g xylose + 20 g glucose + 2 g acetic acid; anaerobic	12
1.42	0.60	7.1	72	80.4	<i>ALD6Δ</i> , <i>XR</i> , <i>XDH</i> , <i>KK</i> (2 copies), adapted strain + 3 × <i>EcAdhE</i> + <i>SeACS</i>	80 g xylose + 20 g glucose + 8 g acetic acid; anaerobic	13
4.15	2.07	0.83	24	97.7	Δ <i>GPD1</i> , Δ <i>ALD6</i> , Δ <i>NDE1</i> , Δ <i>NDE2</i> + <i>SeEutE</i>	100 g glucose + 4 g acetic acid; hypoxic	E5 (this study)

^a Efficiency was calculated based on the total consumption of sugars and acetate. The theoretical maximum yield is 0.51 g g^{-1} for glucose and xylose, and 0.77 g g^{-1} for acetate. ^b Experimental data were estimated approximately from figure visualizations.

enabled consumption of 7.1 g L^{-1} acetate alongside 80 g L^{-1} xylose, significantly improving ethanol yield.¹³ When NADH limitation was alleviated by expressing *EhADH1*, up to 5.3 g L^{-1} acetate was co-utilized with 114 g L^{-1} glucose.⁹ In a system targeting triacetic acid lactone (TAL)—a product whose biosynthesis does not require NADH—acetate-to-xylose co-utilization reached a 1:4 ratio.¹⁴ Notably, these acetate concentrations ($1\text{--}15 \text{ g L}^{-1}$) are consistent with levels typically found in lignocellulosic hydrolysates.¹²

Collectively, these findings confirm that acetate co-utilization enhances ethanol production across diverse substrate types (glucose, xylose, and mixtures), and the performance of E5 aligns well with industrially relevant acetate concentrations (Table 4). Its hypoxia and acetate tolerant phenotype make E5 a promising and robust chassis for scalable bioethanol production from lignocellulosic biomass.

3.7 Future direction and metabolic engineering

The metabolic engineering strategy applied in E5 (*GPD1Δ ALD6Δ NDE1Δ NDE2Δ [SeEutE]*) represents a novel approach for developing a *S. cerevisiae* strain that prefers hypoxic conditions associated with ethanol-induced flocculation and achieves unprecedented rates (342% faster than the native strain) of co-fermenting acetic acid with glucose under the tested conditions. These results were achieved with the integration of a single copy of *SeEutE*. An additional strategy to enhance performance involves the “tugging” approach, successfully applied in previous studies. Henningsen *et al.* employed two copies of *BaAdhE* to co-utilize acetic acid with glucose, followed by the introduction of NADPH-dependent alcohol dehydrogenase to supply additional NADH to *EhADH*.⁹ Therefore, further improvements could possibly be realized by introducing multiple copies of *SeEutE* or combining it with more efficient acetyl-CoA synthase enzymes.

Acetic acid metabolism is regulated by carbon catabolite repression (CCR) through the *ACS1* and *ACS2* genes in response to glucose.^{14,46} Zhang *et al.* optimized acetate reduction in the presence of glucose and xylose by expressing three copies of both mutated *SeACS* and codon-optimized *EcAdhE*, which enhanced xylose fermentation and reduced by-products such as xylitol and glycerol while enhancing acetate co-utilization. A similar approach could be applied to E5 to further enhance co-fermentation.¹³

4. Conclusions

A metabolically engineered *S. cerevisiae* strain, E5, was developed for glucose fermentation and acetic acid co-utilization under hypoxic conditions. The quadruple deletion of *GPD1*, *ALD6*, *NDE1*, and *NDE2*, combined with *SeEutE* expression, enhanced flocculation triggered by ethanol production at $\sim 3\%$. E5 fermented 10% glucose and 0.4% acetic acid 342% faster than C1, achieving 98% of the theoretical ethanol yield—11% higher than C1. Additionally, it utilized 25% of the supplemented acetic acid and exhibited a 9% higher fermentation rate under hypoxic conditions. This engineering strategy



demonstrates significant potential for bioethanol and bio-based chemical production from glucose and acetic acid.

Data availability

All necessary data required to assess our findings are available in this manuscript or its ESI.† Further details related to this article may be requested from the authors.

Author contributions

Sadat M. R. Khattab: conceptualization, methodology, validation, formal analysis, investigation, resources, data curation, writing – original draft, writing – review & editing, supervision, project administration, funding acquisition. Mohamed O. Mohamed: methodology, validation, formal analysis, investigation, data curation, writing – review & editing. Takashi Nagata: validation, writing – review & editing, supervision. Takashi Watanabe: conceptualization, writing – review & editing, project administration, funding acquisition. Masato Katahira: conceptualization, data curation, writing – review & editing, supervision, project administration, funding acquisition.

Conflicts of interest

The authors declare that they have no financial interests or personal relationships that could have influenced the work presented in this paper.

Acknowledgements

This work was supported by a Mission 5–2 Research Grant from the Research Institute for Sustainable Humanosphere, Kyoto University [Project No. 5-2-3], the Joint Usage/Research Program on Zero-Emission Energy Research, Kyoto University [ZE2024A-01], e-ASIA Joint Research Program [e-ASIA JRP, JPMJSC18E1], the Research Unit for Realization of Sustainable Society (RURSS), Research Coordination Alliance, Kyoto University, and collaborative research with DAICEL Co. Ltd. We would like to thank Prof. Alhosein Hamada, Faculty of Agriculture and Food Science, King Faisal University, KSA, for his assistance in verifying the statistical analysis.

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