Rewiring regulation on respiro-fermentative metabolism relieved Crabtree effects in *Saccharomyces cerevisiae*

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\textbf{ABSTRACT}

The respiro-fermentative metabolism in the yeast *Saccharomyces cerevisiae*, also called the Crabtree effect, results in lower energy efficiency and biomass yield which can impact yields of chemicals to be produced using this cell factory. Although it can be engineered to become Crabtree negative, the slow growth and glucose consumption rate limit its industrial application. Here the Crabtree effect in yeast can be alleviated by engineering the transcription factor Mth1 involved in glucose signaling and a subunit of the RNA polymerase II mediator complex Med2. It was found that the mutant with the \textit{MTH1}^{A81D} & \textit{MED2}^{422V} allele could grow in glucose rich medium with a specific growth rate of 0.30 h\textsuperscript{-1}, an ethanol yield of 0.10 g g\textsuperscript{-1} and a biomass yield of 0.21 g g\textsuperscript{-1}, compared with a specific growth rate of 0.40 h\textsuperscript{-1}, an ethanol yield of 0.46 g g\textsuperscript{-1} and a biomass yield of 0.11 g g\textsuperscript{-1} in the wild-type strain CEN.PK 113-5D. Transcriptome analysis revealed significant downregulation of the glycolytic process, as well as the upregulation of the TCA cycle and the electron transfer chain. Significant expression changes of several reporter transcription factors were also identified, which might explain the higher energy efficiencies in the engineered strain. We further demonstrated the potential of the engineered strain with the production of 3-hydroxypropionic acid at a titer of 2.04 g L\textsuperscript{-1}, i.e., 5.4-fold higher than that of a reference strain, indicating that the alleviated glucose repression could enhance the supply of mitochondrial acetyl-CoA. These results suggested that the engineered strain could be used as an efficient cell factory for mitochondrial production of acetyl-CoA derived chemicals.

1. Introduction

Under glucose rich conditions yeast *Saccharomyces cerevisiae* predominantly consumes glucose via the fermentative pathway, even in the presence of oxygen. In *S. cerevisiae*, the respiration pathway can produce 18 molecules of ATP from 1 molecule of glucose with an estimated P/O ratio of 1.2, while the fermentative pathway can only produce 2 molecules of ATP [1]. The respiro-fermentative metabolism, also referred to as the Crabtree effect, has been studied for decades, in which the carbon flux overflow to fermentation is coupled with the respiration repression. The underlying mechanism involves complex regulatory networks to simultaneously control the fermentation and respiration [2–4]. From an evolutionary perspective, the Crabtree positive yeast seems to favor an increased ATP production rate over the high ATP yield and the ‘make-accumulate-consume’ strategy seems to benefit survival in natural environments [5–8]. However, the low energy efficiency of respiro-fermentation not only limits biomass production, but also restricts synthesis of non-ethanol chemicals, especially when their synthesis pathways involve higher consumption of energy or redox power than those supplied from fermentation [9]. Previous studies on chemostat cultivations have found that the biomass yield of *S. cerevisiae* can reach up to 0.5 g g\textsuperscript{-1} glucose with a high respiration rate when the dilution rate is below the critical value, normally in the range of 0.2–0.3 h\textsuperscript{-1}. As the dilution rate increases above the value, the respiration rate decreases while fermentation starts to dominate, and the biomass yield decreases [10–12].
Attempts on pathway engineering have been made to convert \textit{S. cerevisiae} to Crabtree negative in order to achieve high energy efficiency, high biomass yield or product yield, including knocking out hexose transporter genes (HXTs), blocking ethanol production pathways, and introducing heterologous acetyl-CoA synthesis pathways, accompanied with adaptive laboratory evolutions (ALEs) \cite{8,13-17}.

However, the engineered strains usually grow with specific growth rates below 0.2 h$^{-1}$ and glucose consumption rates below 0.63 g L$^{-1}$ g$^{-1}$ h$^{-1}$, compared with the wild-type strain with 0.4 h$^{-1}$ and 2.88 g L$^{-1}$ g$^{-1}$ h$^{-1}$, respectively. The significantly reduced growth and glucose consumption rates limit their industrial application.

ALE studies of pyruvate decarboxylase (pdc) deletion strains have revealed several mutations of regulators that may play important roles in the respiro-fermentative metabolism shift. Studies find \textit{MTH1} mutations can repress the transcription of several hexose transporter genes to reduce the glucose transport, and hereby reduce the metabolic burden from the carbon flux overflow \cite{16,18-21}. Meanwhile, the MED2*432Y mutation has been identified with the global regulatory impact for improved cell growth \cite{13}. It is thus interesting to investigate whether their combination can result in synergistic effects to reinforce alleviation of the Crabtree effect while rescue cell growth.

The metabolism shift from fermentation to respiration may direct more carbon flux to the mitochondria and enhance the energy and the mitochondrial acetyl-CoA synthesis, thereof increase production on acetyl-CoA derived products. As an acetyl-CoA derived chemical, 3-hydroxypropionic acid (3-HP) (3-HP) is synthesized from acetyl-CoA by two enzymes acetyl-CoA carboxylase (ACC) and malonyl-CoA reductase (MCR), in which acetyl-CoA is converted to malonyl-CoA by ACC, and then malonyl-CoA is converted to 3-HP by MCR with consumption of NADPH. The enhanced acetyl-CoA supply has been demonstrated to be crucial for 3-HP synthesis in several previous studies \cite{22,23}, and it is thus interesting to investigate whether their combination can result in synergistic effects to enhance alleviation of the Crabtree effect while rescue cell growth.

Therefore, in this work we studied the effects of \textit{MTH1} and \textit{MED2} alleles in a wild-type strain \textit{CEN.PK} 113-5D, focusing on fermentation and respiration pathways, as previous studies performed in pdc deletion strains could not reflect their effects on ethanol production. We also performed transcriptome analysis on these engineered mutants to reveal their regulation on cell metabolism. Then we investigated 3-HP production in the mutants by targeting MCR into the yeast mitochondria. This work shed lights for harnessing yeast mitochondria for production of acetyl-CoA derived chemicals.

2. Materials and methods

2.1. Construction of plasmids and yeast strains

Plasmids and strains constructed in this study could be found in Table 1, and primers used in Supplementary Table 1. The yeast \textit{CEN.PK} 113-5D was used as the reference strain. Genomic engineering of yeast strains was performed using the GTR-CRISPR system according to previous reports \cite{24}. Mutations of \textit{MTH1}^\Delta13, \textit{MTH1}^\Delta855, \textit{MTH1}^\Delta113, \textit{BD11} and \textit{MED2}^*432Y were introduced using a 2-step method. First, the spacer sequence (1st spacer) containing the target mutation point was replaced with donor DNAs of different promoters. Then the 2nd spacer sequence was replaced using donor DNA and the Cas9 plasmid harboring gRNA targeting 2nd spacer to generate the mutations. The first step of \textit{MTH1} mutation used pCas9_M226 and donor amplified with primers ds226s1 and ds226s1. The second step of \textit{MTH1} mutation used pCas9_M226s and donor amplified with primers ds226_1 and ds226_81 (or ds226_85). The first step of \textit{MED2} mutation used pCas9_MED2 and donor amplified with primers med2_s1.1 and med2_s1.2. The partial deletion mutant of \textit{MTH1} was achieved using pCas9_M226 and donor DNA amplified with ds226_ID1s and ds226_ID2.

The promoter replacement of \textit{MTH1} was achieved by pCas9_MTH1p and donor DNAs of different promoters.

To harness yeast mitochondria for 3-HP production, the mitochondrial localization sequence (MLS) CAT2m was fused to the N-terminal of CaMCR from \textit{Chloroflexus aurantiacus} \cite{25}. The intact CaMCR and dissected CaMCR with improved activity \cite{26} were cloned into the vector pUGG1 using Golden Gate assembly, resulting in pMCR1 and pMCR2, respectively. In pMCR1, the \textit{pTDH3} fragment amplified with primers P124–P125 from pYC1 were assembled into pUGG1. In pMCR-C, the \textit{pTDH3-CAT2m} fragment amplified from pMCR1 was constructed by assembling donor DNAs of different promoters.

\begin{table}[h]
\centering
\begin{tabular}{|c|c|c|}
\hline
\textbf{Plasmid/Strain} & \textbf{Description} & \textbf{Reference} \\
\hline
\textit{pCas9} & 2\text{u}, ampR, \textit{TEF1p} CaMCR, \textit{SNR52} & [24] \\
\textit{pScURA} & 2\text{u}, ampR, \textit{gRNA, UR3A, SNR52p, U6G} & [24] \\
\textit{pCas9_M226} & pCas, \textit{Ura3}, \textit{pSNR52B_M226step1, gRNA}\_\textit{SNR52} & This study \\
\textit{pCas9_M226s} & pCas, \textit{Ura3, pSNR52B_M226step2, gRNA}\_\textit{SNR52} & This study \\
\textit{pCas9_MED2} & pCas, \textit{Ura3, pSNR52B_Med2step1, gRNA}\_\textit{SNR52} & This study \\
\textit{pCas9_MED2_N} & pCas, \textit{Ura3, pSNR52B_Med2step2, gRNA}\_\textit{SNR52} & This study \\
\textit{pCas9_MTH1p} & pCas, \textit{Ura3, pSNR52B_MTH1p, gRNA}\_\textit{SNR52} & This study \\
\textit{pUGG1} & ori, AmpR, \textit{2u, Ura3} & [28] \\
\textit{pYCI} & ori, AmpR, \textit{2u, Ura3, pTEF11, CaMCR, CYC1} & [22] \\
\textit{pMCRI} & pUGG1, \textit{pTDH3-CAT2m, CaMCR, CYC1} & This study \\
\textit{pMCR2} & pUGG1, \textit{pPGK1-CAT2m, (MCR-N)-ADH1, pTDH3-CAT2m, (MCR-C)-CYC1} & This study \\
\textit{CEN.PK 113-5D} & \textit{MATa SUC2 MAL8C ura3-52} & [11] \\
\textit{ZS_mth1} & \textit{CEN.PK 113-5D, MTH1}^\Delta11D & This study \\
\textit{ZS_mth1_2} & \textit{CEN.PK 113-5D, MTH1}^\Delta855 & This study \\
\textit{ZS_mth1_3} & \textit{CEN.PK 113-5D, MTH1}^\Delta113,\textit{BD11} & This study \\
\textit{ZS_mth1_4} & \textit{CEN.PK 113-5D, MTH1-\DeltaT} & This study \\
\textit{ZS_med2} & \textit{CEN.PK 113-5D, MED2}^*432Y & This study \\
\textit{ZS_mm} & \textit{CEN.PK 113-5D, MED2}^*432Y & This study \\
\textit{MTH1_HXT1p} & \textit{CEN.PK 113-5D, MTH1p::HXT1p} & This study \\
\textit{mth1_HXT1p} & \textit{ZS_mth1, MTH1p::HXT1p} & This study \\
\textit{MTH1_PGK1p} & \textit{CEN.PK 113-5D, MTH1p::PGK1p} & This study \\
\textit{mth1_PGK1p} & \textit{ZS_mth1, MTH1p::PGK1p} & This study \\
\textit{5D_mCR} & \textit{CEN.PK 113-5D, pMCR1} & This study \\
\textit{5D_3HP} & \textit{CEN.PK 113-5D, pMCR2} & This study \\
\textit{mm_3HP} & \textit{ZS_mm, pMCR2} & This study \\
\textit{mm_3HP} & \textit{ZS_mm, pMCR2} & This study \\
\hline
\end{tabular}
\caption{Plasmids and strains used and constructed in this study.}
\end{table}
purification and plasmid construction were purchased from Omega Bio-Tek.

2.2. Medium and cultivation conditions

LB medium was used for *E. coli* DH5α cultivation, and ampicillin was supplemented to LB medium to a final concentration of 80 mg L⁻¹ when needed. YPD medium and SC-URA medium were used for yeast cultivation [28]. All plates contained 15 g L⁻¹ agar.

The minimal synthetic medium for yeast fermentation in shake flasks was composed of 5 g L⁻¹ (NH₄)₂SO₄, 14.4 g L⁻¹ KH₂PO₄, 0.5 g L⁻¹ MgSO₄·7H₂O, trace metal solution, vitamin solution, 40 mg L⁻¹ uracil and 20 g L⁻¹ glucose, with adjusted pH of 6.5. The compositions of trace metal solution and vitamin solution was previously described in Ref. [29]. Yeast cells were pre-cultured in 5 mL YPD medium at 30 °C overnight, then inoculated with an initial OD₆₀₀ of 0.01, and cultured in the 20 mL minimal medium in 250 mL shake flasks at 30 °C. For the mitochondria activity analysis, a mitochondria pyruvate transporter inhibitor UK-5099 (Sigma, USA) dissolved in DMSO was supplemented into medium with a final concentration of 200 nM when OD₆₀₀ reached around 1. The minimal synthetic medium for yeast fermentation in 1L bioreactors (DasGip, Germany) was composed of 5 g L⁻¹ (NH₄)₂SO₄, 3 g L⁻¹ KH₂PO₄, 0.5 g L⁻¹ MgSO₄·7H₂O, 40 mg L⁻¹ uracil, 20 g L⁻¹ glucose, trace metal solution and vitamin solution, as described in Ref. [30]. For bioreactor batch fermentation, after pre-cultured in shake flasks to exponential phase (OD₆₀₀ ~1), cells were inoculated in bioreactor with an initial OD₆₀₀ of 0.01. The fermentation was performed at 30 °C, with pH controlled at 5 by 2 M KOH and the dissolved oxygen concentration controlled above 30% by adjusting the stirring speed (800–1500 rpm) and the aeration rate (1–1.5vvm).

All the chemical used for cell cultivations were purchased from Sinopharm Chemical Reagent Corporation, China, if not specifically annotated.

2.3. Measurement of biomass and extracellular metabolites

The biomass was quantified by OD₆₀₀ (optical density at 600 nm) using the spectrophotometer (GENESYS 30 Visible, Thermo Electron Scientific) and cell dry weight (CDW) calculated by the weight of washed cell collected from 5 mL culture. Cells were harvested to a pre-weighed dry microporous filter sheet (Aquo system, 0.22 μm, 50 mm) and washed with sterile water twice using vacuum filtration. Then the sheet with cells folded inside was first dried in the microwave at 100 W for 3 min, and then dried in a glass dryer with allochroic Silica Gel blue for more than 3 days.

Cell samples were centrifuged at 15,000 g for 3 min at 4 °C and the supernatants were then filtered through 0.22 μm polyethersulfone (PES) filters (JIN TENG, China) and collected for extracellular metabolite analysis on a HPLC system (Shimadzu LC-20AT, Japan) with Aminex HPX-87H column (Bio-Rad, USA) with a UV detector and a RID detector at 65 °C. Measurements of glucose, ethanol, pyruvate, succinate, acetate and glycerol were performed as described in Ref. [15]. Measurements of 3-HP were performed using 0.5 mM H₂O₂ as eluent for 20 min with a flow rate at 0.5 mL min⁻¹. The volume for sample injection was 10 μL.

2.4. Transcriptome analysis

Samples were collected when the OD₆₀₀ reached 0.9–1.2 in triplicate. Around 10 mL broth was transferred into 50 mL falcon tubes with 30 mL crushed ice and centrifuged at 4000 rpm at 4 °C for 3 min. Cell pellets were then quenched in liquid nitrogen immediately and stored at –80 °C before RNA extraction. Total RNA was extracted using TRIzol Reagent (Invitrogen, USA) based on the manufacturer’s instructions. The library preparation and sequencing were carried out on the HiSeq instrument (Illumina, USA) by GENEWIZ Inc. Principal Component Analysis (PCA) was plotted according to variance stabilizing normalization and differential gene expression was analyzed according to Benjamini-Hochberg method using the DEseq2 package in the R studio [31]. Reporter Gene Ontology (GO) term analysis was performed using the PIANO (Integrative Analysis of Omics) package in R studio [32]. The functional enrichment analysis of KEGG pathways was performed using the Database for Annotation, Visualization and Integrated Discovery (DAVID) [33].

2.5. Cellular respiration measurement

Cellular respiration was characterized using the oxygen consumption rate (OCR), which can be quantified using Seahorse XFe96 Extracellular Flux Analyzer (Agilent, USA) [34,35]. The OCR assays for cells responding to two different concentrations of glucose, 1 g L⁻¹ and 10 g L⁻¹, were performed in six biological replicates. Cells were incubated overnight in YPD medium to exponential growth phase and washed once with sterile water, and diluted by XF base medium (Agilent, USA) with glucose of two different concentrations, including 1 g L⁻¹ and 10 g L⁻¹ glucose, respectively. Then, 175 μL cell suspensions were transferred into XFe96 microplates with a final concentration of 2 × 10⁵ cells per well. The plates were centrifuged at 500 rpm for 3 min and incubated for 30 min at 30 °C without CO₂, and the OCR measurements were performed at an interval of 6.5 min for three timepoints for basal values. Prior to the assays, the Seahorse sensor cartridges were hydrated overnight and the XFe96 microplates were coated with 25 μL⁻¹ poly-l-lysine (25 μg mL⁻¹) for 30 min, and then evaporated to dryness at room temperature.

3. Results

3.1. Effects of the mutations MTH1ΔA81D and MED2ΔI85Y on cell growth and metabolism

Previous studies identified several MTH1 mutations in glucose sensitive strains, that could relieve the Crabtree effect [15,18–21,36,37]. Among these alleles, MTH1ΔA81D, MTH1ΔI85Y and MTH1ΔT were investigated as these mutations were positioned in a highly conserved module to form a putative alpha helix [15,21]. The mutagenesis of MTH1 was performed using the GTR-CRISPR system, generating four mutant strains with MTH1ΔA81D, MTH1ΔI85Y and MTH1ΔT, respectively. When cultured in minimal medium in shake flasks, growth profiles of the mutants showed that MTH1ΔA81D and MTH1ΔT resulted in reduced cell growth, glucose consumption and ethanol production, while MTH1ΔI85Y did not have a significant effect on cell growth compared with the WT strain (Supplementary Fig. 1). The mutants with MTH1ΔA81D and MTH1ΔT accumulated 6 g L⁻¹ ethanol within 27 h, and their specific growth rates were both around 0.22 h⁻¹, while the wildtype strain accumulated 8.3 g L⁻¹ ethanol within 21 h, with the specific growth rate around 0.38 h⁻¹. Further characterization in bioreactor cultivations revealed that the strain with the MTH1ΔA81D allele consumed glucose slower with less ethanol produced (Fig. 1B), i.e., its specific growth rate decreased by 50%, the biomass yield increased by 36%, and the ethanol yield decreased by 73.9%, compared with the wildtype strain (Fig. 1A).

Next, the mutagenesis of MED2 was performed in the wild-type strain, generating the mutant strain with MED2ΔI85Y allele. When cultured in minimal medium in bioreactors, the mutant exhibited a slightly higher biomass yield of 0.14 gDCW gGlc⁻¹ and specific growth rate of 0.42 h⁻¹, but a lower specific glucose consumption rate of 3.13 g L⁻¹ h⁻¹ (Fig. 1C, Table 2), suggesting a higher energy efficiency of cell metabolism.

We thus combined the two alleles, and the strain with MTH1ΔA81D&MED2ΔI85Y showed a faster cell growth with the specific growth rate of 0.30 h⁻¹ and less ethanol accumulated (Fig. 1D, Table 2). Meanwhile, the biomass yields increased 40% compared with the strain with MTH1ΔA81D. Moreover, the strain with MTH1ΔA81D&MED2ΔI85Y showed a much shorter ethanol phase (Fig. 1D), which might be resulted from
from the altered respiro-fermentative metabolism. Although the strain with \textit{MTH1}^{432Y} \& \textit{MED2}^{432Y} consumed glucose faster than the strain with \textit{MTH1}^{A81D} (Fig. 1B&D), the specific glucose consumption rates of the two strains with \textit{MTH1}^{432Y} \& \textit{MED2}^{432Y} were almost the same (Table 2), which might be subjected to the restricted glucose uptake rates.

### 3.2. Transcriptional profiles of the mutant strains

To gain further insights of how the \textit{MED2}^{432Y} and \textit{MTH1}^{A81D} alleles resulted in altered respiro-fermentative metabolism of the engineered strains, cells of the WT strain and the mutants with \textit{MTH1}^{A81D}, \textit{MED2}^{432Y} and \textit{MTH1}^{A81D} \& \textit{MED2}^{432Y} were sampled at exponential phase for transcriptome sequencing in triplicate experiments. Principal component analysis of the transcriptome dataset treated with variance stabilization transformation showed strong reproducibility of biological triplicates (Fig. 2A), and revealed the variances among the strains, which was consistent with the physiological data described above (Table 2).

Differential gene expression analysis (Supplementary Table 2) identified a number of genes with significantly differential expression in the mutants compared with the WT strain (p-adj < 0.01), as shown in Fig. 2B. The \textit{MED2}^{432Y} allele resulted in 348 genes significantly regulated, while \textit{MTH1}^{A81D} resulted in 3882 genes significantly regulated. The \textit{MED2}^{432Y} \& \textit{MTH1}^{A81D} allele resulted in 3650 genes significantly regulated, in which 3230 genes were shared with those resulted from \textit{MTH1}^{A81D} and 245 genes were identified as commonly differentially expressed among all mutant strains. Gene set enrichment analysis using DAVID bioinformatics resources [33] revealed that 245 commonly differentially regulated genes were significantly enriched in 19 gene sets (p-adj < 0.05) (Fig. 2C), which included glycolytic process, gluconeogenesis, pentose-phosphate shunt, amino acid biosynthesis process, transposition, DNA biosynthetic process, RNA phosphodiester bond hydrolysis, oxidation-reduction process, and transmembrane transport.

Furthermore, reporter GO term analysis using PIANO R package [32] was performed on genes with significantly differential expressions (p-adj < 0.01) for each mutant in comparison with the WT strain (Supplementary Table 3). The mutations of \textit{MED2}^{432Y}, \textit{MTH1}^{A81D} and \textit{MED2}^{432Y} \& \textit{MTH1}^{A81D} resulted in 25, 176 and 166 GO terms significantly regulated, which suggested different metabolic regulation involving \textit{MED2}^{432Y} and \textit{MTH1}^{A81D} (Fig. 2D). Top 4 reporter GO terms identified included the downregulated glycolytic process and gluconeogenesis, as well as the upregulated electron transfer chain and heme biosynthesis (Fig. 2E). Meanwhile, other biological processes that were also significantly regulated included amino acid biosynthesis, DNA biosynthetic process, transposition oxidation-reduction process and transmembrane transport.

The underlying mechanism for the altered respiro-fermentative metabolism might involve a number of perturbations at the transcriptional level, as well as the post-transcriptional and post-translational...

### Table 2

Physiological parameters of the wild-type strain (WT) and the mutants with \textit{MED2}^{432Y} (ZS_med2), \textit{MTH1}^{A81D} (ZS_mth1), and \textit{MTH1}^{A81D} \& \textit{MED2}^{432Y} (ZS_mm), respectively. The cultivations were performed in minimal medium with 2% glucose in bioreactors in replicate. Error bars represent ±standard errors.

|            | \( \mu_{\text{max}} \) g L\(^{-1}\) h\(^{-1} \) | \( T_{\text{ex}} \) g CDW g Glc \(^{-1} \) | \( q_{\text{GLU}} \) g L\(^{-1}\) h\(^{-1} \) | \( V_{\text{Glu\text{th}}} \) g Glc g Glc \(^{-1} \) h\(^{-1} \) | \( q_{\text{Eth}} \) g L\(^{-1}\) h\(^{-1} \) |
|------------|---------------------------------------------|---------------------------------------------|---------------------------------------------|---------------------------------------------|---------------------------------------------|
| WT         | 0.40 ± 0.01                                | 0.11 ± 0.02                                 | 3.32 ± 0.01                                 | 0.46 ± 0.01                                 | 1.52 ± 0.04                                 |
| ZS_med2    | 0.42 ± 0.01                                | 0.14 ± 0.00                                 | 3.13 ± 0.10                                 | 0.45 ± 0.01                                 | 1.39 ± 0.08                                 |
| ZS_mth1    | 0.21 ± 0.00                                | 0.15 ± 0.01                                 | 1.46 ± 0.08                                 | 0.13 ± 0.03                                 | 0.18 ± 0.03                                 |
| ZS_mm      | 0.30 ± 0.01                                | 0.21 ± 0.01                                 | 1.47 ± 0.07                                 | 0.10 ± 0.01                                 | 0.15 ± 0.01                                 |

Fig. 1. Growth, glucose and ethanol profiles of the wild-type strain (A) and the mutants with \textit{MTH1}^{A81D} (B), \textit{MED2}^{432Y} (C) and \textit{MTH1}^{A81D} \& \textit{MED2}^{432Y} (D), respectively. The cultivations were performed in minimal medium with 2% glucose in bioreactors in duplicate and error bars represent ±standard errors.
Fig. 2. Transcriptional analysis of the mutant strains. (A) Principal component analysis (PCA) plot of the transcription data in triplicates. (B) Venn diagram of significantly regulated genes of mutant strains in comparison with the WT strain (p-adj < 0.01). (C) Gene set enrichment analysis of the 245 commonly regulated genes. Fold enrichment indicated the magnitude of enrichment against the genome background of the strain S288C analyzed via DAVID. (D) Venn diagram of highly scored (p-value < 0.01) reporter GO terms through analyzing significantly regulated genes of the mutants compared with the WT strain (p-adj < 0.05). (E) The common high scored reporter GO terms in distinct-directional up and down class presented by their significance in the mutants. (F) The high scored reporter transcription factors (TFs) in distinct-directional up and down class presented by their significance. (G) Expression levels of reporter TFs in the mutant strains compared with the wildtype strain.
levels. Therefore, we performed reporter TFs analysis, and found several high-scored TFs with their target genes directionally upregulated or downregulated. These reporter TFs and their expression levels were highly relevant for glucose derepression (Fig. 2F&G), including upregulation of SNF1, CAT8, ADR1, HAP1/2/3/4/5, and downregulation of MIG1, MIG2, MTH1, GRR1, GCR1, suggesting the global transcriptional responses resulted from the MTH1 and MED2 allele.

3.3. Carbohydrate metabolism and respiration analysis

We further investigated the differentially expressed genes related to carbohydrate metabolism (Fig. 3) and mitochondrial electron transport chains (Fig. 4A). Gene expression levels in the strain with MED2*432Y were much more different from those in other two mutants, but more similar with those of strain WT except that the genes related with glycolysis were downregulated, which might support its faster growth as glycolytic enzymes constituted the majority of the cellular proteome [38]. In the two mutants with MTH1*431D, most genes related to carbohydrate metabolism were regulated in the same direction but with variations in expression levels, probably due to the global regulation imposed by MED2*432Y for a faster growth rate. The major isoforms of hexose transporter encoding genes HXT1, HXT3 and HXT4 were downregulated, and other glucose transporter genes with high and moderate glucose affinity HXT6, HXT7, and HXT10 were upregulated. Most genes of glycolysis, glycerol production, and ethanol production were downregulated, including PGK1, PFK1/2, FBA1, TDH1/2/3, PGK1, GPM1, ENO1/2, CDC19, GDH1/2, GPD1/2 and ADH1 et al. Most genes involved in glycolgen biosynthesis, glycerol consumption, ethanol consumption, TCA cycle, glyoxylate cycle and fatty acid metabolism were upregulated, which were normally repressed under high glucose conditions (Fig. 3).

The electron transfer chain was coupled with the TCA cycle to completely oxidize acetyl-CoA to CO2, that consumes oxygen and yields ATP. Reporter GO analysis revealed upregulation of electron transfer chain in all the mutants (Fig. 2E). Therefore, we further investigated differentially expressed genes related to the electron transfer chain in mutant strains, with expression levels presented in Fig. 4A. All genes in the electron cycle were upregulated in the strains with MTH1*431D, and around 30% genes were upregulated in the strain with MED2*432Y. Moreover, the respiration activities were analyzed through oxygen consumption rate (OCR) measured under two glucose levels, 10 g L-1 and 1 g L-1, when the enzymes involved in the TCA cycle and respiration were normally repressed [39]. The mutants showed higher respiration activities compared with the WT strain (Fig. 4B), which was consistent.

![Fig. 3. Transcriptional representation of genes involved in carbohydrate metabolism. All data was made in comparisons with the WT strain.](image-url)
with their transcriptional results. The OCR values of the two mutants were about 2-fold higher than those of the WT strain, under both 10 g L\(^{-1}\) and 1 g L\(^{-1}\) glucose conditions.

Furthermore, to quantify the effects of the elevated mitochondrial activities on cell metabolism, a mitochondrial pyruvate transporter inhibitor UK-5099 was supplemented in the early exponential growth (OD\(_{600}\) around 1.0), as inhibited pyruvate transport would thereof restrict the following TCA cycle and electron transport chain. In WT strain and the mutant with \(MED2^{*432Y}\), UK-5099 slowed down the growth and ethanol consumption during the ethanol phase (Fig. 5 A&C). In the strains with \(MTH1^{A81D}\), UK-5099 slowed down the glucose consumption and ethanol accumulation during the glucose phase, as well as growth and ethanol consumption during the ethanol phase (Fig. 5 B&D). The results suggested that the respiration pathway subjected to mitochondrial pyruvate transport was derepressed earlier in the strains with \(MTH1^{A81D}\), indicating that the TCA cycle and electron transfer chain were derepressed at relatively high glucose.

3.4. Improved 3-HP production in strain ZS_mm with alleviated respiro-fermentative metabolism

The higher enzymatic activities of the TCA cycle and electron transfer chain in the strain with \(MED2^{*432Y} \& MTH1^{A81D}\) would probably enhance the mitochondrial acetyl-CoA supply, and thereof improve synthesis of its derived chemicals [22,23]. Therefore, we investigated 3-HP production by expressing CaMCR (MCR from \(C. aurantiacus\)) in the mitochondria, as mitochondrial acetyl-CoA could be converted to malonyl-CoA by a mitochondrial acetyl-CoA carboxylase Hfa1 [40].

The mitochondrial localization signal of \(CAT2\) gene CATm was fused at the N terminals of the intact and dissected CaMCR genes to target them into the mitochondria. The intact and dissected CaMCR with CATm were cloned into the episomal plasmids, yielding pMCR1 and pMCR2, respectively. When the two versions of CaMCR expressed in the mitochondria of the WT strain, the dissected one obtained a 3-HP titer of 0.45 g L\(^{-1}\), while the intact one showed a titer of 0.22 g L\(^{-1}\) (Supplementary Fig. 2), suggesting a higher enzymatic activity of the dissected

Fig. 4. Effects of \(MTH1\) and \(MED2\) mutations on cell respiration. (A) Expression fold changes of the genes involved in mitochondrial electron transport chains. All data were made in comparisons with WT strain. (B) Oxygen consumption rates (OCRs) of the wild-type strain (WT) and mutant strains under two different glucose levels using a Seahorse XF96 analyzer. The measurements were performed in replicate and error bars represent \(\pm\) standard errors.

Fig. 5. The growth, glucose and ethanol profiles of the wild-type strain (A) and the mutants ZS_mth1(B), ZS_med2 (C) and ZS_mm (D) with (red line) and without (black line) mitochondrial pyruvate transporter inhibitor UK-5099. The cultivations were performed in minimal medium with 2% glucose in shake flasks in triplicate and error bars represent \(\pm\) standard errors.
CaMCR.

Then the dissected CaMCR was used for further investigation in the strains of WT and ZS_mm. Briefly, plasmid pMCR2 was transformed into the two strains, generating the strains 5D_3HP and mm_3HP, respectively. The constructed strains were first evaluated for respiration capacity by cultivations –OCR values than those of the 5D_3HP strain under both conditions (Supplementary Fig. 3), confirming its higher respiration activities. Then the strains were evaluated in the minimal medium with 20 g L\(^{-1}\) glucose for 3-HP production. As shown in Fig. 6A, the strain 5D_3HP produced 0.35 g L\(^{-1}\) 3HP, and more than 75% of 3-HP were accumulated in ethanol phase, as mitochondrial acetyl-CoA supply was restricted due to glucose repression. In the mm_3HP strain, 3-HP was accumulated much earlier, and reached to 1.69 g L\(^{-1}\) when glucose exhausted, comprising almost 83% of 3-HP titer. The final 3-HP reached as high as 2.04 g L\(^{-1}\) at 60 h. These results suggested that the altered respiro-fermentative metabolism of ZS_mm not only could provide higher energy efficiency for biomass formation, and support efficient 3-HP synthesis in the mitochondria, which might be resulted from the derepressed TCA cycle and electron transfer chain.

4. Discussions

Previous studies showed that MTH1 alleles could improve growth of pdc deletion strains by limiting the glycolysis flux by reducing its degradation [15,16,21], indicating the possibility to regulate glucose uptake rates by manipulating MTH1 expression levels. Based on FPKM (fragments per kilobase million) values from transcription data (Supplementary Fig. 4), HXT1p and PKG1p were selected as weaker and stronger promoters compared with MTH1p, respectively. Indeed, the strain with reduced Mth1 expression exhibited a fast cell growth, glucose consumption and ethanol accumulation, to similar levels as in the wild-type strain, while the strain with elevated Mth1 expression resulted in a much slower growth, glucose consumption and ethanol accumulation (Supplementary Fig. 5). Yet, the reduced specific growth and glucose uptake rate would hinder its application for production of non-ethanol biochemicals.

The MED2\(^{432Y}\) allele was previously reported to improve cell growth of pdc deletion strain by its global regulation on carbon metabolism in pdc deletion strain [13], and in this study it could also improve cell growth of the wild-type and the mutant with MTH1\(^{AB1D}\) allele, which might mainly result from regulation on glycolysis and glucose repression as revealed in our transcriptional analysis. A previous study revealed that glucose transport could impose high control on the glycolytic flux if the capacity reduced below 56% of the wild-type strain [17], which was consistent with our results in which the specific glucose consumption rate of the strain with MED2\(^{432Y}\) & MTH1\(^{AB1D}\) was 43.7% of the WT strain. The lower specific ethanol production rate in the strain with MTH1\(^{AB1D}\)&MED2\(^{432Y}\) suggested less carbon flux overflow into fermentation and possibly more carbon flux into respiration, and thereof improved cell growth.

The transcription results also revealed the reduced overflow flux to fermentation, and triggered upregulation of the TCA cycle and electron transfer chain. It was noteworthy that several high scored reporter TFs were involved in glucose derepression. Specially, Mig1 and Mig2 functioned as repressors with Snf1 [41–43], while Cat8 and Adr1 functioned as Snf1 activators in glucose derepression associated with gluconeogenesis, glycerol utilization, ethanol utilization, fatty acid utilization and glyoxylate cycle [44–47]. Thus, significant downregulation of Mig1 and Mig2 and upregulation of CAT8 and ADR1 might result in the glucose derepression. Similarly, upregulation of HAP1/2/3/4/5, as transcription activators for respiratory genes [48], could increase mitochondrial biogenesis and the respiratory capacity [49,50]. The upregulation of MTH1 and downregulation of GRR1, required for HXTs induction, indicated that HXTs might be regulated by both intracellular and extracellular glucose availability. Downregulation of GCR1, as an activator of glycolysis genes [51–53], was highly related with significantly downregulation of genes involved in glycolysis process. Deep mining of key players in transcription factors involved in carbohydrate metabolism and respiration could be useful to further improve biomass yield and reduce ethanol yield.

The strain with MTH1\(^{AB1D}\)&MED2\(^{432Y}\) could grow with a specific growth rate of 0.30 h\(^{-1}\) and specific glucose consumption rate of 1.45 g L\(^{-1}\) g\(_{dw}\) h\(^{-1}\), which was higher than pdc deletion strains [13,21,26]. Furthermore, the depression of the mitochondrial activities under high glucose levels would make the yeast mitochondria robust compartments for non-ethanol chemical synthesis, as revealed in 3-HP production. The transcriptional results revealed that several genes related to the glyoxylate cycle were also upregulated in the mutant, indicating that peroxisomes could be explored as well. As several studies harnessed yeast mitochondria and peroxisomes for non-ethanol chemical production [56–58], strategies used in this study would be useful to enhanced activities of these compartments. Yet, the specific glucose consumption seemed to be limited by the reduced glucose uptake rate in the strains with the MTH1\(^{AB1D}\) allele. It would be optimal to optimize the glucose uptake rate to increase cell growth while reducing ethanol accumulation for industrial application for non-ethanol production.

5. Conclusions

The MTH1\(^{AB1D}\) and MED2\(^{432Y}\) alleles were investigated in wild-type strain for the alleviated Crabtree effects and 3-HP production. The MTH1\(^{AB1D}\) mutation resulted in lower maximum growth rates and ethanol yields accompanied with higher biomass yields. The MED2\(^{432Y}\) mutation resulted in improved cell growth rates and biomass yields as

![Fig. 6](image-url) The growth, glucose, ethanol and 3-HP profiles of the strains without (A) and with (B) the mutations of MTH1\(^{AB1D}\)&MED2\(^{432Y}\). The cultivations were performed in minimal medium with 2% glucose in shake flasks in triplicate and error bars represent ±standard errors.
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...as well as lower ethanol yields. The transcriptome analysis identified several differentially expressed genes and GO terms, which involved downregulation of glycolytic process and gluconeogenesis and upregulation of electron transfer chain and heme biosynthesis. The strain with...
