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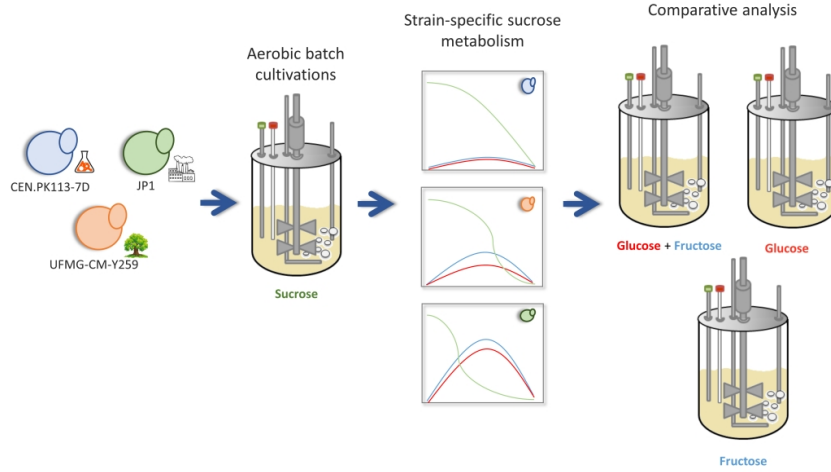
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1   **Aerobic growth physiology of *Saccharomyces cerevisiae* on sucrose is strain-dependent**

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9

10   **Abstract**

11   Present knowledge on the quantitative aerobic physiology of the yeast *Saccharomyces*
12   *cerevisiae* during growth on sucrose as sole carbon and energy source is limited to either adapted
13   cells or to the model laboratory strain CEN.PK113-7D. To broaden our understanding of this
14   matter and open novel opportunities for sucrose-based biotechnological processes, we
15   characterized three strains, with distinct backgrounds, during aerobic batch bioreactor
16   cultivations. Our results reveal that sucrose metabolism in *S. cerevisiae* is a strain-specific trait.
17   Each strain displayed a distinct extracellular hexose concentration and invertase activity profiles.
18   Especially, the inferior maximum specific growth rate (0.21 h^{-1}) of the CEN.PK113-7D strain,
19   with respect to that of strains UFMG-CM-Y259 (0.37 h^{-1}) and JP1 (0.32 h^{-1}), could be associated
20   to its low invertase activity (0.04 to 0.09 U mg$_{DM}^{-1}$). Moreover, comparative experiments with
21   glucose or fructose alone, or in combination, suggest mixed mechanisms of sucrose utilization by
22   the industrial strain JP1, and points out the remarkable ability of the wild isolate UFMG-CM-259
23   to grow faster on sucrose than on glucose in a well-controlled cultivation system. This work hints
24   to a series of metabolic traits that can be exploited to increase sucrose catabolic rates and
25   bioprocess efficiency.

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1   **INTRODUCTION**

2   Sucrose has long been used in the food industry as a substrate for the production of bakery

3   goods and beverages. In the last decades, however, it has also been considered a valuable

4   feedstock for the replacement of petrochemical derived materials, due to its low market price

5   (Polat and Linhardt 2001; Marques et al. 2016). Especially, sucrose feedstocks have a high land

6   efficiency and vast availability of sugar-rich crops. Furthermore, sucrose does not require any

7   pretreatment prior to its use in industrial fermentation. Whilst fuel ethanol is by far the prime

8   product of the nonfood sucrose-based industry (OECD-FAO 2018), with an annual production of

9   33.1 billion liters in Brazil alone in 2019 (Brazilian Sugarcane Industry Union (Unica)), the

10   potential use of this disaccharide goes beyond fuel manufacturing, with value-added chemicals

11   such as citric acid (Förster et al. 2007; Show et al. 2015), lactic acid (Lunelli et al. 2010; Wang et

12   al. 2012) and farnesene (E4tech et al. 2015) also being successfully commercialized. Several

13   additional chemical intermediates can be produced using sucrose as substrate, including 5-

14   hydroxymethylfurfural (Peters et al. 2010; E4tech et al. 2015), 1-2-propylene glycol (Peters et al.

15   2010; Rosales-Calderon and Arantes 2019), acrylic acid (E4tech et al. 2015), and succinic acid

16   (Chan et al. 2012; Jiang et al. 2014; E4tech et al. 2015; Rosales-Calderon and Arantes 2019).

17   The yeast *Saccharomyces cerevisiae*, the primary workhorse of the biotechnology industry

18   (Steensels et al. 2014; Steensels and Verstrepen 2014; Kavšček et al. 2015), naturally

19   metabolizes sucrose either through its hydrolysis in the periplasmic space or through direct

20   uptake via active transport of the disaccharide and its hydrolysis in the cytosol (Badotti et al.

21   2008) (**Figure 1**). In the first mechanism, *S. cerevisiae* needs to express an invertase-encoding

22   gene (*SUC2*, the most common one) and secrete the protein to the periplasmic space after

23   oligomerization and post-translational modification (Carlson and Botstein 1982). The hydrolysis'

24   products, namely glucose and fructose, can enter the cells by facilitated diffusion. In the other

25   mechanism, sucrose is directly transported to the cytosol via a proton-symport mechanism

26   mediated by the high-affinity ($K_M = 7.9 \pm 0.8$ mM) transporter Agt1p or the low-affinity ($K_M = 120 \pm 20$ mM) transporters encoded by *MALx1* genes (x denotes the locus number) (Stambuk et

28   al. 2000; Basso et al. 2011). In this case, ATP is invested to expel the imported proton with a

29   stoichiometry of 1:1, in order to keep cell's homeostasis and the proton motive force across the

30   plasma membrane. Consequently, cells are expected to achieve a higher glycolytic rate to

31   compensate for the lower energy efficiency in the overall metabolic process.

2

1   FIGURE 1

2   Aerobic sucrose metabolism by *S. cerevisiae* has been shown to be fast, with maximum

3   specific growth rates ranging from 0.38 to 0.57 h^{-1} (van Dijken et al. 2000; Beato et al. 2016),

4   depending on the strain, conditions and analytical methods employed. In spite of this successful

5   relationship between sucrose and *S. cerevisiae* (Marques et al. 2016), little attention has been

6   given to the specific effects of this carbon source on the quantitative aerobic physiology of this

7   yeast. The few studies involving a comprehensive quantitative aerobic physiological analysis

8   were performed with strains pre-evolved on sucrose for 200 to 250 generations (Orlowski and

9   Barford 1991; Barford et al. 1995; Mwesigye and Barford 1996), or with the model laboratory

10   strain CEN.PK113-7D (Herwig et al. 2001), which limits our understanding of relevant

11   phenomena. On the one hand, the use of pre-adapted strains changes their initial/natural

12   physiology. On the other hand, the results published on a non-adapted strain refer to one

13   particular laboratory strain, meaning that we still do not have a good overview of the physiology

14   of this yeast species during growth on sucrose and how this eventually varies among different

15   strains. Relevant biological questions remain to be elucidated: does the strain background

16   influence *S. cerevisiae*'s physiology on sucrose? Would a non-adapted fast sucrose-growing

17   strain rely on direct uptake as a preferred natural mechanism of sucrose utilization? How does

18   growth on sucrose compare to growth on glucose? A better understanding of *S. cerevisiae*'s

19   growth on sucrose will not only aid in answering those questions but also open novel

20   opportunities for the development of strain improvement strategies to enhance sucrose-based

21   industrial bioprocesses, in terms of the yields, productivities, and/or titers required for the

22   production of market goods in a competitive manner.

23   Here, we present the quantitative physiology of *S. cerevisiae* grown on sucrose as the sole

24   carbon and energy source in aerobic batch bioreactor cultivations. To access the effects of the

25   strain background on sucrose physiology, the experiments were performed with one laboratory

26   (CEN.PK113-7D), one industrial (JP1) and one wild isolate (UFMG-CM-Y259) strain. While the

27   CEN.PK113-7D strain serves as a reference for physiological studies (Van Dijken et al. 2000),

28   JP1 and UFMG-CM-Y259 were chosen due to their different behaviors on sucrose (Beato et al.

29   2016). JP1 was isolated from fermenters used to produce fuel ethanol from sugarcane in the

30   Northeast of Brazil and is a relatively thermotolerant strain (da Silva Filho et al. 2005; Della-

3

1 bianca and Gombert 2013), whereas UFMG-CM-Y259 presented the highest maximum specific

2 growth rate (μ_{MAX}) on sucrose in a screening test made with 18 different *S. cerevisiae* strains. Its

3 μ_{MAX} on sucrose was also ~20% higher than the corresponding value on glucose (Beato et al.

4 2016).

5 Additionally, in order to investigate the mechanisms underlying the diverging physiologies

6 observed during growth on sucrose, cultivations were also performed on an equimolar mixture of

7 glucose and fructose and on each one of these two monosaccharides separately.

8 **MATERIAL AND METHODOLOGY**

9 **Yeast strains, preservation and pre-cultures**

10 The *Saccharomyces cerevisiae* strains studied in this work comprise an indigenous strain,

11 named UFMG-CM-Y259, isolated from barks of the tree *Quercus rubra* (Northern Red Oak)

12 located in Santuário do Caraça (Minas Gerais, Brazil) within the Atlantic Forest biome (Beato et

13 al. 2016), kindly provided by Dr. Carlos A. Rosa (Federal University of Minas Gerais, Belo

14 Horizonte, Brazil); JP1, a Brazilian fuel ethanol industrial strain isolated from the Japungu

15 Agroindustrial sugarcane-based distillery located in Northeastern Brazil (da Silva Filho et al.

16 2005), kindly provided by Dr. Marcos Morais Jr. (Federal University of Pernambuco, Recife,

17 Brazil); and the laboratory strain CEN.PK113-7D (kindly provided by Dr. Petter Kötter,

18 EUROSCARF, Germany), which is largely employed in physiological studies by the scientific

19 community (van Dijken et al. 2000).

20 Stock cultures were prepared by growing cells until stationary phase in 500-ml Erlenmeyer

21 flasks containing 100 ml YPD medium (per liter: 10.0 g **Y**east extract, 20.0 g **P**eptone and 20.0 g

22 **D**extrose/Glucose), in an incubator shaker (Certomat BS-1, Braun Biotech International, Berlin,

23 Germany) operating at 30 °C and 200 rpm for 24 h. 20% (v/v, final concentration) glycerol was

24 added and 1 ml aliquots were stored in 2-ml cryogenic vials in an ultra-freezer (CryoCube HEF,

25 model F570h-86, Eppendorf, Hamburg, Germany) at -80 °C until further use.

26 The pre-culture medium was prepared according to (Verduyn et al. 1992) with a few

27 modifications. The medium consisted of (per liter): 3.0 g KH_2PO_4, 6.6 g K_2SO_4, 0.5 g

28 $MgSO_4.7H_2O$, 2.3 g urea, 1 ml trace elements solution, 1 ml vitamins solution and 10 g carbon

29 source (sucrose, glucose, fructose or an equimolar mixture of glucose and fructose). The initial

4

1 pH of the pre-culture medium was adjusted to 6.0 using 2 mol/l KOH. Sterilization of the pre-

2 culture medium occurred by filtration through 0.22 μm pore membranes (Millex-GV, Merck

3 Millipore, Massachusetts, USA).

4   To prepare the inoculum for bioreactor cultivations, the content of one stock cryogenic vial

5 was centrifuged at 867 g for 4 min, and cells were transferred to a 500-ml Erlenmeyer flask

6 containing 100 ml of the pre-culture medium. The pre-inoculum was left in a shaker (Certomat

7 BS-1, Braun Biotech International) set at 30 °C and 200 rpm for 24 h preceding the direct

8 transfer of 1 ml of its content to another shake-flask with fresh pre-culture medium. After a

9 second round of 24 h of growth in a shaker, operating at the same settings as before, an aliquot

10 sufficient to start the bioreactor batch cultivation with an optical density of 0.2 at 600 nm was

11 collected and washed. For the washing procedure, cells were centrifuged at room temperature

12 and 3500 g for 3 min, the supernatant was discarded and fresh pre-culture medium was added to

13 the cell pellet. This was then vortexed and centrifuged once again. At last, the cells were

14 resuspended with cultivation medium and transferred to a proper flask that allows for an aseptic

15 transfer to the bioreactor.

16   **Bioreactor batch cultivations**

17   A synthetic medium formulated as described in (Verduyn et al. 1992) was used in all

18 bioreactor batch cultivations. The medium contains per liter: 3.0 g KH_2PO_4, 5.0 g $(NH_4)_2SO_4$, 0.5

19 g $MgSO_4.7H_2O$, 1 ml trace elements solution, 1 ml vitamins solution and 20 g (or equivalent in

20 g_{GLCeq}) carbon and energy source ─ sucrose, glucose, fructose or an equimolar mixture of

21 glucose and fructose. The initial pH was adjusted to 5.0 using of 4 mol/l NaOH. Sterilization of

22 the medium occurred by filtration through 0.22 μm pore membranes (Millex-GV, Merck

23 Millipore, Massachusetts, USA).

24   A 2-liter bioreactor (Applikon Biotechnology B.V., Delft, The Netherlands), with a working

25 volume of 1.2 l was used throughout this work. Cells were grown at 30°C and 800 rpm stirring

26 speed. Aeration occurred with compressed air at 0.5 l min^{-1} flow rate using a mass flow

27 controller (Model 58505, Brooks Instrument, Hatfield, USA). The pH of the broth was controlled

28 at 5.0 by automatic addition of a 0.5 mol/l KOH solution. A 10% (v/v) antifoam C emulsion

29 (Sigma-Aldrich, Missouri, USA) was added manually to the broth upon necessity. Aliquots were

30 collected manually at different sampling times to be analysed for extracellular metabolites

5

1   concentrations, dry cell mass and invertase activity. For each collected sample, the exact mass

2   withdrawn was determined.

3   The gas flowing out of the bioreactor had its CO_2 and O_2 molar fractions determined using a

4   gas analyzer device (Rosemount NGA 2000, Emerson Electric Co., Ferguson, USA).

5   Determination of O_2 molar fraction was performed through a paramagnetic detector, while an

6   infrared detector was used to determine CO_2 molar fraction. Bioreactor volumetric rates were

7   calculated taking into account variations in pressure and volume (e.g. due to sampling, base and

8   antifoam addition).

9   Cultivations were finished when a decrease in the CO_2 molar fraction in the off-gas was

10   observed.

11   **Analytical Methods**

12   *Dry cell mass concentration*

13   Cells were harvested and filtered through a 0.45 μm nitrocellulose membrane (SO-Pak filters,

14   HAWP047S0 – Merck Millipore, Massachusetts, USA) that had been previously dried and

15   weighed (m_1). The cell pellet was washed twice with demineralized water. The filter containing

16   the pellet was dried in an oven at 70 °C for 48 h and then placed in a desiccator to cool down

17   prior to being weighed (m_2). The cell dry mass (X_{DM}) was calculated by dividing the difference

18   between the filter's mass after and before filtration by the sample volume filtered (V); $X_{DM} = (m_2$

19   $- m_1)/V$. The result is expressed in $g_{DM}.l^{-1}$.

20   *Concentration of extracellular metabolites*

21   Sampling for extracellular metabolites followed the procedure described elsewhere (Mashego

22   et al. 2003). Briefly, a defined volume of broth was rapidly collected in a syringe containing a

23   calculated amount of cold metal beads (-20°C) - enough to cool the collected broth to 1°C, and

24   filtered through a 0.45 μm PVFD membrane (Millex - HV, Merck Millipore) directly into a tube.

25   All sample tubes containing filtrate were immediately placed on ice and stored at - 80°C until

26   analysis.

27   The concentrations of residual sugars in the filtrate - sucrose, glucose and fructose - were

28   determined either enzymatically using the K-SURFG kit (Megazyme, Bray, Ireland), following

29   the manufacturer's instructions, or by means of ion chromatography coupled with pulsed

6

1   electrochemical detection at gold electrode (CarboQuad pulse, AgCl reference). The

2   chromatography system was a Dionex ICS – 5000 HPIC system with AS-AP sampler, SP pump

3   (Thermal Scientific, Waltham, USA) equipped with a Carbopac PA-20 3×150 mm column and

4   Aminotrap 3×30 mm precolumn at 30°C, eluted at 0.5 ml min^{-1} with 5% NaOH 200 mM for 15

5   min, followed by a cleaning step with 20% sodium acetate solution 0.5 M in 200 mM NaOH for

6   5 min, and reequilibration for 15 min with Milli-Q water (eluents C and D). The temperature of

7   the detector was kept at 15°C.

8   The concentrations of ethanol, glycerol and organic acids (lactate, succinate and acetate) were

9   always determined by high performance liquid chromatography (HPLC) using a Bio-Rad

10   Aminex HPX-87H column (Bio-Rad, USA), kept at 60°C and eluted with highly diluted

11   phosphoric acid (60 ml min^{-1}) at a pH between 2 and 3, which was preheated before use. The

12   samples were injected using an autosampler (Waters 717, USA). Detection of organic acids was

13   performed via a UV detector (Waters 2489), while detection of ethanol or glycerol was

14   performed using a refraction index detector (Waters 2414, USA). During cultivations without

15   sucrose, residual glucose and fructose were determined in the same HPLC run used for the other

16   metabolites, and the detection of these hexoses was performed using a refraction index detector

17   (Waters 2414, USA). HPLC data were processed using Empower software (Waters Corporation,

18   Milford, USA).

19   *Extracellular invertase activity*

20   Extracellular (or periplasmic) invertase activity was determined according to the approach

21   described in (Silveira et al. 1996), with a few modifications. Briefly, cells were centrifuged and

22   resuspended in distilled water such that a 20 g_{DM}.l^{-1} suspension was obtained. Next, the cells

23   were treated with Succinate-Tris buffer (pH 5.0) containing sodium fluoride, which is an

24   inhibitor of enolase. A sucrose solution was added to the reaction mixture and the glucose

25   formed at 30°C due to disaccharide hydrolysis was measured using an enzymatic kit (R-

26   Biopharm AG, Darmstadt, Germany). Invertase activity was reported as μmol of glucose

27   produced per minute per milligram of cell dry mass (U mg_{DM}^{-1}).

28   **Calculation of physiological parameters**

29   Prior to calculating physiological parameters, the experimental data points were treated as

30   follows. Extracellular metabolites concentrations and CO_2 and O_2 amounts were plotted against

7

1   time and a polynomial was fitted to the data. Concentrations or amounts were calculated for each

2   time point taking the polynomial equation. Glucose and fructose concentrations during

3   experiments performed with either sucrose or an equimolar mixture of glucose and fructose were

4   not adjusted to avoid concealing their actual consumption trend. Only data points within the

5   exponential growth phase (EGP) were considered for the calculation of physiological parameters.

6   The specific growth rate during the EGP (μ_{MAX}) was calculated as the slope of the straight line

7   adjusted to the linear region of an $\ln(X_{DM})$ *versus* time plot. The time span corresponding to this

8   linear region was considered to be the EGP. Biomass ($Y_{X/S}$) and product yield ($Y_{P/S}$) on

9   substrate, except for CO_2, were calculated as the absolute value of the slope of a biomass

10 concentration (X_{DM}) *versus* substrate concentration (S) plot, and of a product concentration (P)

11 *versus* substrate concentration (S) plot, respectively. CO_2 yield on substrate was derived from the

12 absolute value of the slope of an integrated CO_2 amount versus substrate amount plot. For

13 cultivations with sucrose or a mixture of glucose and fructose, substrate concentration (S) was

14 determined as the sum of the concentrations of all carbohydrates in g_{GLGeq} l^{-1}. For that, 1 g of

15 sucrose was considered to be equivalent to 1.0526 g of hexose.

16 The maximum specific substrate consumption rate (during the EGP) was calculated as the

17 ratio between the maximum specific growth rate and the biomass yield on substrate; $q_{S,MAX}$ =

18 $\mu_{MAX}/Y_{X/S}$.

19 The maximum specific product formation rate (q_P) was calculated taking the ratio between the

20 desired yield on substrate and biomass yield on substrate, and multiplying the result by the

21 maximum specific growth rate; $q_{P,MAX} = \mu_{MAX}*Y_{P/S}/Y_{X/S}$.

22 For determining the specific oxygen consumption rate (q_{O2}), the integrated amount in mmol of

23 oxygen consumed per gram of substrate consumed at each sampling time was first obtained by

24 the absolute slope of the integrated O_2 *versus* substrate amount plot. Following, this value was

25 divided by the biomass yield on substrate and multiplied by the maximum specific growth rate.

Calculation of the percentage of fermented sugar during the EGP

27 The specific rate of ethanol formation (q_{ETH}) was used to calculate the specific rate of CO_2

28 formed due to fermentation (q_{CO2_ferm}) in the EGP. This value was divided by 2 and considered to

29 be the specific rate of sugar that was fermented (e.g. q_{GLC_ferm}), as a result of the stoichiometry of

1 ethanolic fermentation ($C_6H_{12}O_6 \rightarrow 2\ C_2H_6O + 2\ CO_2$). This value, in turn, was divided by the

2 (total) maximum specific substrate consumption rate (e.g. $q_{GLC,MAX}$) and multiplied by 100,

3 resulting in the % fermented sugar.

4 **Sequencing of the *S. cerevisiae* JP1 and UFMG-CM-Y259 genomic DNA**

5 The *S. cerevisiae* JP1 and UFMG-CM-Y259 strains were sequenced at Serviço Nacional de

6 Aprendizagem Industrial (SENAI), Rio de Janeiro, Brazil. DNA extraction was performed using

7 the Wizard® Genomic DNA Purification kit (Promega) and sequencing libraries were prepared

8 using the Nextera DNA Flex kit (Illumina), always following the manufacturer's guidelines.

9 Sequencing was carried out on an Illumina NextSeq 550 machine, using a High Output cartridge,

10 producing 300 bp reads (2x 151 bp paired-end). Coverage (relative to a 12 Mbp haploid genome)

11 was 427x for the UFMG-CM-Y259 strain and 174x for the JP1 strain. The reads can be accessed

12 using the BioProject ID PRJNA699867 at NCBI.

13 **Bioinformatics pipeline**

14 The fastq files containing the sequencing results were analyzed using a pipeline recommended

15 by the Research Computing Faculty of Applied Sciences (RCFAS), Harvard University, which

16 included: 1) trimming with NGmerge (Gaspar, 2018), 2) alignment against the *S. cerevisiae*

17 S288c reference genome (yeastgenome.org, release R64) with Bwa (Li and Durbin, 2009), 3)

18 conversion of .sam to .bam files and validation of the final .bam file with Picard

19 (http://broadinstitute.github.io/picard), 4) haplotype calling with GATK HaplotypeCaller

20 (McKenna et al, 2010), 5) databasing with GATK GenomicsDBImport, and 6) genotyping with

21 GATK GenotypeGVCFs. The .bam files generated with Picard were used to manually inspect

22 the desired variants using the Integrated Genome Viewer (IGV, Robinson et al. 2011). The final

23 .vcf files generated with GATK were used to annotate variants using SnpEff (Cingolani et al.

24 2012).

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2   **RESULTS AND DISCUSSION**

3   *S. cerevisiae* **strains from different environments display different physiologies on**

4   **sucrose**

5   The physiology of the *S. cerevisiae* strains CEN.PK113-7D, UFMG-CM-Y259 and JP1

6   during growth on sucrose was assessed in controlled batch aerobic bioreactor cultivations

7   (**Figures 2 and 3; Table 1; Supplementary Figures S1-S5, Supplementary Tables S1-S3**).

8   The laboratory strain CEN.PK113-7D displayed the lowest maximum specific growth rate on

9   sucrose (μ_{MAX} = 0.21 ± 0.01 h^{-1}), which is 56.8 % and 65.6 % of the corresponding rates

10   presented by the indigenous strain UFMG-CM-Y259 and the industrial strain JP1, respectively

11   (**Table 1**). Strain CEN.PK113-7D was also found to present the lowest specific substrate

12   consumption rate (q_{SMAX} = -8.23 ± 0.37 mmol$_{GLCeq}$ g$_{DM}^{-1}$ h^{-1}), when compared to the values

13   displayed by UFMG-CM-Y259 (q_{SMAX} = -12.84 ± 0.21 mmol$_{GLCeq}$ g$_{DM}^{-1}$ h^{-1}) and by JP1 (q_{SMAX}

14   = -12.29 ± 0.12 mmol$_{GLCeq}$ g$_{DM}^{-1}$ h^{-1}).

15   Accordingly, the relative amount of substrate metabolized through the fermentative pathway

16   was higher for UFMG-CM-Y259 and JP1 strains (~ 80%; **Supplementary Figure S1**) than that

17   of CEN.PK113-7D (~ 70%; **Supplementary Figure S1**). This could also be deduced by the

18   respiratory quotient (**Table 1**). In addition, allied to such observations, the ethanol yield on

19   substrate was lower in the laboratory strain ($Y_{E/S}$ = 0.36 ± 0.01 g g$_{GLCeq}^{-1}$), as compared to the

20   indigenous ($Y_{E/S}$ = 0.41 ± 0.01 g g$_{GLCeq}^{-1}$) and the industrial strain ($Y_{E/S}$ = 0.41 ± 0.00 g g$_{GLCeq}^{-1}$).

21   One of the main traits of the yeast *Saccharomyces cerevisiae* is its capability to perform

22   aerobic fermentation (De Deken 1966). In other words, even in the presence of oxygen, a higher

23   relative amount of the substrate is subjected to fermentative catabolism rather than respiratory. A

24   phenomenon that constrains the use of the respiratory pathway in yeast under circumstances of

25   high sugar concentrations, as imposed in batch mode, is the so-called glucose (or carbon

26   catabolite) repression. High concentrations of this sugar trigger a signaling cascade that will

27   culminate in the repression of the transcription of genes encoding components of the electron

28   transport chain and other respiratory proteins (Gancedo 1992; Santangelo 2006; Belinchón and

29   Gancedo 2007a; Conrad et al. 2014). This strategy results in an energetically less efficient

30   metabolism, as the ATP yield from fermentation is lower than that from respiration; for instance,

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1   in *S. cerevisiae* approximately 18 ATP are produced per mole of glucose via respiration against

2   only 2 through fermentation (Verduyn et al. 1991; De Kok et al. 2012). A few hypotheses have

3   been proposed to explain how cells overcome the lower ATP yield from fermentation (Pfeiffer

4   and Morley 2014; Nilsson and Nielsen 2016; Niebel et al. 2019). The so-called rate/yield trade-

5   off hypothesis (RYT), for instance, proposes that the cells accelerate growth to rise energy

6   production rates (Pfeiffer and Morley 2014). Through another perspective, the use of

7   fermentation allows for a higher ATP yield per protein mass as compared to respiration (Nilsson

8   and Nielsen 2016). Our data somehow fit with these theories, in the sense that the lower growth

9   rate of CEN.PK113-7D on sucrose, as compared to the other two strains, correlates with a lower

10 relative amount of substrate channeled to fermentation in this laboratory strain.

11 The different ways in which the three yeast strains investigated here consume sucrose is likely

12 to reflect the ecological niche of each one individually, at least partially. Strain JP1 was isolated

13 from a sugarcane-based distillery where sugarcane juice — rather than molasses, which is not as

14 sucrose-rich — is used to prepare the fermentation medium (da Silva Filho et al. 2005), meaning

15 that it is adapted to an anaerobic environment, in which fermentative metabolism is the only

16 option, and to excess sucrose. On the other hand, CEN.PK113-7D is commonly employed in

17 fundamental laboratory research on physiology (van Dijken et al. 2000), where glucose is the

18 major substrate. The long-term exposure of a microbe to a specific environment may alter its

19 regulatory mechanisms, as well as cause adaptation, strategies used to cope with stress and to

20 enhance fitness. In this sense, JP1 holds great advantage over the CEN.PK113-7D strain when

21 sucrose is the carbon and energy source, due to its primary extensive contact with this sugar,

22 which may have changed sucrose regulation in this lineage, or triggered mutations in genes

23 encoding proteins that are essential to sucrose metabolism, resulting in an improved phenotype.

24 We compared the *SUC2* sequences of JP1, UFMG-CM-Y259 and CEN.PK113-7D to the

25 reference sequence of the S288c strain (**Supplementary Figure S6, Supplementary Table S4**).

26 The protein sequence of the CEN.PK113-7D strain is identical to the one of the S288c strain.

27 There are two amino acido changes shared by JP1 and UFMG-CM-Y259, namely N84H (Asn →

28 His) and Q88E (Gln → Glu). In the JP1 strain alone, there are also two additional amino acid

29 changes, with respect to the reference: A77T (Ala → Thr) and A409P (Ala → Pro). All other

30 nucleotide differences do not lead to amino acid changes. We also analysed the 1,000 bp region

31 upstream of the *SUC2* ORF (**Supplementary Table S5**) and verified that there are some

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1   differences when the JP1 and the UFMG-CM-Y259 sequences are compared to the reference.

2   Whether the amino acid changes in the *SUC2* protein and/or any differences in gene expression

3   due to nucleotide changes in the upstream region of *SUC2* in the JP1 and the UFMG-CM-Y259

4   strains are related to their faster growth phenotype on sucrose, when compared to the

5   CEN.PK113-7D strain, remains to be experimentally tested.

6   Whether CEN.PK113-7D cells could evolve on sucrose and achieve a fitness comparable (or

7   superior) to that of JP1 or UFMG-CM-Y259, remains to be explored. Previous laboratory

8   evolution studies have demonstrated the potential of *S. cerevisiae* CEN.PK113-7D cells to

9   improve its phenotype for maltose (Jansen et al. 2004), galactose (Hong et al. 2011) and mixed-

10   substrate consumption (Papapetridis et al. 2018). Especially when sucrose was the substrate of

11   choice, evolved engineered *S. cerevisiae* strains were shown to increase sucrose transport

12   capacity (Basso et al. 2011; Marques et al. 2018). Concerning the UFMG-CM-Y259 strain, it

13   was originally isolated from the barks of a tree that was brought by Europeans to Brazil, namely

14   *Quercus rubra* (Northern Red Oak tree) (Beato et al. 2016). The characteristics of this niche are

15   largely unknown. However, sucrose has been found in the barks of several other *Quercus* species

16   from where *Saccharomyces* yeasts, including *S. cerevisiae*, have been isolated (Sampaio and

17   Gonçalves 2008), suggesting that this disaccharide might also be present in Northern red oak

18   tree's bark. Also, another study showed that a yeast strain isolated from an oak tree performed

19   better than Ethanol Red (a widespread industrial strain used mainly in fuel ethanol production

20   from corn) under mimicked industrial conditions in the laboratory (Ruyters et al. 2015).

21   Taken together, these observations led us to believe that the historical background plays a key

22   role on the enhanced growth and sucrose consumption displayed by JP1 and UFMG-CM-Y259,

23   over CEN.PK113-7D cells.

24   FIGURE 2

25   FIGURE 3

26   **During cultivations of *S. cerevisiae* on sucrose, higher glucose and fructose accumulation**

27   **seem to correlate with higher growth rates**

28   The sugar consumption profiles during aerobic growth on sucrose of the three strains reveal

29   that the accumulation of the released monosaccharides in the cultivation broth occurs to different

12

1   extents (**Figure 3**). The highest concentrations, 6.0 $g_{GLCeq}.l^{-1}$ of glucose and 6.1 $g_{GLCeq}.l^{-1}$ of

2   fructose, were observed for the industrial strain JP1. For UFMG-CM-Y259, the levels of

3   extracellular glucose and fructose reached 3.2 $g_{GLCeq}.l^{-1}$ and 3.3 $g_{GLCeq}.l^{-1}$, respectively. On the

4   other hand, the accumulation of hexoses was minimal with the laboratory strain CEN.PK113-7D

5   and did not surpass 0.69 $g_{GLCeq}\ l^{-1}$ and 0.91 $g_{GLCeq}\ l^{-1}$ of glucose and fructose, respectively. Since

6   this was the strain with the lowest maximum specific growth rate on sucrose (**Table 1**), we

7   speculate that hexose accumulation during growth on this disaccharide contributes to faster

8   growth. This, in turn, could be caused by the regulatory mechanisms related to the extracellular

9   glucose concentration – such as the repression of the transcription of respiratory-related genes,

10   which is known to have pleiotropic effects on yeast's metabolism (Newcomb et al. 2003;

11   Santangelo 2006; Belinchón and Gancedo 2007b; Kayikci and Nielsen 2015).

12   Moreover, as the accumulation of either glucose or fructose in the cultivation broth is a result

13   of the difference between invertase activity and hexose transport rate, the observed different

14   profiles could indicate that (1) hexose transport occurs at different rates in the different strains

15   and/or (2) the enzyme invertase has different activity in each strain.

16   The existence of low- and high-affinity hexose transporter systems in the yeast *S. cerevisiae*

17   has been well documented and the expression of these proteins depends on the levels of their

18   substrates in the environment (Bisson et al. 1993; Ozcan and Johnston 1995; Kruckeberg 1996;

19   Özcan and Johnston 1999). For low-affinity hexose transporters, such as Hxt1p and Hxt3p, the

20   K_M for glucose ranges from 15 to 20 mM, while the $K_{M(glucose)}$ of high-affinity hexose

21   transporters, for instance Hxt2p and Hxt6p, is in the range of 1 to 2 mM (Özcan and Johnston

22   1999). Thus, one could assume that the cultivations on sucrose performed in this work triggered

23   the expression of hexose transporters that differ from case to case, according to the concentration

24   of glucose or fructose present in the broth. For instance, with the CEN.PK113-7D strain, for

25   which the lowest levels of hexoses were observed (**Figure 3**), it is expected that high-affinity

26   transporters are expressed. In a similar fashion, during cultivations with the UFMG-CM-Y259

27   and JP1 strains, low-affinity hexose transporters might be present. Transporter affinity and

28   maximum velocity (V_{max}) of these transport systems are inversely correlated in *S. cerevisiae*

29   (Walsh et al. 1994). This means that a high-affinity transporter system displays lower V_{max} as

30   compared to low-affinity, high-rate transporter systems. The hypothesis that hexose transport

13

1   occurs at different rates in the three scenarios investigated here is, therefore, supported by these

2   previously described properties of hexose transporters in *S. cerevisiae*.

3   **Invertase activity might constrain *S. cerevisiae* CEN.PK113-7D's growth rate on sucrose**

4   To investigate whether invertase activity contributes to the different hexose accumulation

5   levels in the broth, we determined the specific activity of the periplasmic form of the enzyme in

6   each of the three strains investigated here, during cultivations on sucrose, in the beginning, mid,

7   and late exponential phase of growth.

8   In all samples, the periplasmic invertase of *S. cerevisiae* CEN.PK113-7D displays lower

9   biomass specific activity, when compared to the UFMG-CM-Y259 and JP1 strains (**Figure 3**).

10   The biomass specific periplasmic invertase activity achieved with the laboratory strain ranged

11   from 0.04 ± 0.00 to 0.09 ± 0.00 U mg_{DM}^{-1}. In a previous study, Herwig and colleagues (Herwig et

12   al. 2001) reported biomass specific total invertase activity within 0.2 to 1.0 U mg_{DM}^{-1},

13   approximately, for the *S. cerevisiae* CEN.PK113-7D strain cultivated in aerobic batch

14   bioreactors at 30°C and pH 5.0, with sucrose as sole carbon and energy source. For this particular

15   strain, it is known that the cytoplasmic invertase displays much lower activity compared to that

16   of the periplasmic invertase (Basso et al. 2011; Marques et al. 2017), therefore the specific

17   activity of the periplasmic form of this enzyme can be assumed to correspond roughly to total

18   invertase activity. Hence, a remarkable difference (one order of magnitude) can be observed

19   between the above mentioned study and our data, obtained under similar conditions. This was

20   accompanied by much higher (approximately 10 times) levels of glucose and fructose in the

21   broth observed by those authors, which is probably a consequence of the higher invertase activity

22   achieved in their study. It is worth noticing that while our pre-cultivation was carried out with

23   sucrose as sole carbon and energy source, their pre-culture was grown on glucose. This shift in

24   the substrate from glucose in the pre-cultivation to sucrose in the cultivation itself likely explains

25   the higher invertase activity observed by Herwig and colleagues, and is presumably a

26   consequence of the cells being exposed to a sudden need for invertase.

27   The inferior biomass specific invertase activity combined with the lower levels of hexose

28   accumulation in the broth (**Figure 3**) and the lower specific sucrose consumption rate (**Table 1**),

29   suggests that sucrose hydrolysis is a constraint for CEN.PK113-7D's growth on this carbon

30   source. The two forms of the enzyme invertase (E.C. 3.2.1.2.6) present in *S. cerevisiae*'s cells are

14

1 translated from genes of the *SUC* family that contains a total of nine structural genes (*SUC1-*

2 *SUC5*, *SUC7-SUC10*) located in distinct loci of several chromosomes (Carlson et al. 1981;

3 Carlson and Botstein 1983; Naumov and Naumova 2010). The *SUC2* gene, located in the

4 subtelomeric region of chromosome IX (Carlson and Botstein 1983; Naumov and Naumova

5 2010), is found in all strains of the *Saccharomyces cerevisiae* species, as well as in other yeasts

6 from the same genus (Naumov and Naumova 2010). The observed differences in biomass

7 specific invertase activity among the strains here analyzed could be due to the mutations in the

8 coding sequence for the *SUC2* gene of the UFMG-CM-Y259 and JP1 strains, as discussed above.

9 We searched for the presence of additional *SUC* genes in the genomes of the JP1 and the UFMG-

10 CM-Y259 strains. The analysis performed (results not shown) indicated no evidence for the

11 presence of any *SUC* gene besides *SUC2*, and we could only find evidence for the presence of a

12 single *SUC2* copy in both JP1 and UFMG-CM-Y259.

13 **The enhanced glycolytic rate of *S. cerevisiae* JP1 on sucrose might be due to a combined**

14 **mechanism of sucrose utilization**

15 To investigate whether the mechanism of sucrose utilization is responsible for the distinct

16 substrate uptake rates observed in the sucrose-experiments, we carried out, under identical

17 operational conditions, a physiological analysis of the studied strains on an equimolar mixture of

18 glucose and fructose (**Figure 4; Table 1; Supplementary Figures S1-S5, Supplementary**

19 **Tables S1-S3**). We hypothesized that if sucrose was being utilized exclusively via periplasmic

20 hydrolysis, no difference in sugar uptake would be observed between the sucrose and the

21 glucose/fructose mixture growth conditions, as long as no limitations in the hydrolysis step

22 occurred.

23 Furthermore, since the metabolism of sucrose in the yeast *S. cerevisiae* differs from that of its

24 monomers (or a mixture of them) only in the step of disaccharide breakdown, when compared to

25 growth on glucose and/or fructose, it can be speculated that active transport of sucrose

26 accelerates the sugar uptake rate, and, consequently, the higher growth rate observed on the

27 disaccharide. In fact, Barford and co-workers (Orlowski and Barford 1991; Barford et al. 1993)

28 demonstrated that the superior growth of *S. cerevisiae* 248 UNSW 703100 — fully adapted to

29 the culture medium for 20 - 250 generations — on sucrose, in comparison to a mixture of its

15

1 monomers, was due to the direct uptake of sucrose molecules by actively growing yeast cells,

2 which is faster than the passive transport of hexoses.

3 From our experiments, *S. cerevisiae* CEN.PK113-7D cells were more efficient in consuming

4 the hexoses when they were provided directly ($q_{SMAX\_sucrose}$ = -8.23 ± 0.37 $mmol_{GLCeq}$ g_{DM}^{-1} h^{-1},

5 q_{SMAX_G+F} = -11.96 ± 0.02 $mmol_{GLCeq}$ g_{DM}^{-1} h^{-1}; **Table 1; Table S1**); the opposite was observed

6 with the industrial strain, JP1 ($q_{SMAX\_sucrose}$ = -12.29 ± 0.12 $mmol_{GLCeq}$ g_{DM}^{-1} h^{-1}, q_{SMAX_G+F} = -

7 10.08 ± 0.39 $mmol_{GLCeq}$ g_{DM}^{-1} h^{-1}; **Table 1; Table S3**), whereas the UFMG-CM-Y259 strain

8 displayed equivalent sugar uptake rates for the sucrose experiment, as compared to the one with

9 the glucose/fructose mixture ($q_{SMAX\_sucrose}$ = -12.84 ± 0.21 $mmol_{GLCeq}$ g_{DM}^{-1} h^{-1}, q_{SMAX_G+F} = -

10 12.19 ± 0.63 $mmol_{GLCeq}$ g_{DM}^{-1} h^{-1}; **Table 1; Table S2**).

11 The observation of immediate glucose and fructose formation in the broth as well as the

12 measured biomass specific periplasmic invertase activity indicate that extracellular hydrolysis of

13 sucrose is a mechanism of sucrose utilization in all experiments carried out on sucrose alone.

14 However, the hypothesis that sucrose is not being actively transported likely holds true for the

15 yeast strains UFMG-CM-Y259 and CEN.PK113-7D. In the latter case, as discussed above, there

16 is evidence for a growth limitation caused by insufficient invertase activity. Because sucrose-

17 grown JP1 cells displayed enhanced glycolytic rates, when compared to cells cultivated on the

18 glucose/fructose mixture — which is evidenced both by the specific rates of substrate

19 consumption and by the specific rates of products formation (**Table 1; Table S3**) —, we believe

20 that a combined mechanism of sucrose utilization is likely to take place when JP1 cells grow on

21 sucrose alone. In other words, both periplasmic hydrolysis by invertase and active transport of

22 sucrose occur in parallel (**Figure 1**).

23 This assumption is supported by a previous work from Barford et al. (Barford et al. 1992), in

24 which a combination of direct sucrose uptake and extracellular hydrolysis with subsequent

25 transport of its monomers was proposed to explain the higher glycolytic rate of the evolved *S.*

26 *cerevisiae* 248 UNSW 703100 strain when grown on sucrose, as compared to an equimolar

27 mixture of the hexoses. Moreover, recently, Prado and co-workers (Prado et al. 2020) argued that

28 a mixed mode of sucrose utilization by the *S. cerevisiae* strain LBGA-01 was responsible for its

29 improved performance at high temperature.

16

1   Since the reference *S. cerevisiae* S288c strain does not have a functional *AGT1* allele, which

2   is incorrectly annotated as a synonym of *MAL11* in the yeastgenome.org database (Trichez et al.

3   2019), we used as a reference the *AGT1* sequence of the industrial *S. cerevisiae* CAT-1 strain

4   (Babrzadeh et al. 2012). The JP1 and the UFMG-CM-Y259 strains presente several (> 20) amino

5   acid changes, with respect to the sequence of the CAT-1 strain (**Supplementary Table S6 and**

6   **Figure S7**). Again, these differences remain to be explored experimentally to see whether *AGT1*

7   in the two strains investigated here encodes an improved sucrose transporter.

8   **Sucrose-grown *S. cerevisiae* UFMG-CM-Y259 cells display higher maximum specific**

9   **growth rate than glucose-grown cells**

10   A previous study from our group (Beato et al. 2016) revealed the capacity of some *S.*

11   *cerevisiae* strains, including UFMG-CM-Y259, to grow faster on sucrose than on glucose

12   through experiments carried out using microtiter plates as a cultivation system and optical

13   density measurements for assessing cell concentration. As the calculation of the maximum

14   specific growth rate can lead to different values depending on e.g. the cultivation system and the

15   cell concentration measurements used (Stevenson et al. 2016), we sought to investigate whether

16   such observations would hold in a well-controlled cultivation system, i.e. a bioreactor combined

17   with direct cell concentration measurements via dry mass determinations. We performed

18   cultivations of the three *S. cerevisiae* strains on glucose as the sole carbon and energy source

19   (**Figure 5; Table 2; Supplementary Figures S1-S5, Supplementary Tables S1-S3**), otherwise

20   under identical conditions as the cultivations hitherto discussed. The UFMG-CM-Y259 strain

21   displayed higher maximum specific growth rate on sucrose (μ_{MAX} = 0.37 ± 0.01 h^{-1}; **Table 1;**

22   **Table S2**) than on glucose (μ_{MAX} = 0.29 ± 0.00 h^{-1}; **Table 2; Table S2**), which corroborates the

23   aforementioned study. Results for the JP1 strain from both studies are also in agreement, in the

24   sense that the strain presented a slightly higher μ_{MAX} on sucrose than on glucose, although this

25   difference was not statistically significant in the previous study (at the 95% confidence level,

26   triplicate experiments in (Beato et al. 2016)). In the present study, we did not perform statistical

27   analysis, since we only performed duplicate experiments in bioreactors, but μ_{MAX} on sucrose and

28   on glucose for JP1 was 0.32 and 0.28 h^{-1}, respectively. For the CEN.PK113-7D strain, while the

29   results from the previous study indicated no statistical difference between μ_{MAX} on sucrose and

30   on glucose, here the value on sucrose (0.21 h^{-1}) was much lower than on glucose (0.31 h^{-1}). We

31   speculate that this could be due to the fact that in the previous study (Beato et al. 2016),

17

1   absorbance measurements, instead of the dry cell mass measurements employed here, were used

2   to calculate μ_{MAX}. Since the CEN.PK113-7D strain is haploid, and haploid cells are typically

3   smaller than diploid ones, the wavelength used in spectrophotometric analysis (typically around

4   600 nm) is closer to the size of the cells, causing bigger differences between direct cell

5   concentration and turbidity measurements (Stevenson et al. 2016).

6   In principle, the maximum specific growth rate on sucrose is not expected to exceed that on

7   glucose because the yeast *S. cerevisiae* is subjected to glucose repression (Gancedo 1998),

8   meaning that any carbon and energy source different from glucose eventually present in the

9   medium will have their consumption delayed, as long as glucose is available. This preference

10   mechanism suggests that the microbe will display a higher specific growth rate on glucose than

11   on any other carbon and energy source. As previously discussed in this work, direct sucrose

12   uptake could explain enhanced growth rates on this disaccharide over related carbon sources.

13   Alongside the superior growth rate, we observed a slightly higher substrate uptake rate during

14   sucrose cultivation ($q_{SMAX\_sucrose}$ = -12.84 ± 0.21 $mmol_{GLCeq}$ g_{DM}^{-1} h^{-1}; **Table 1; Table S2**), as

15   compared to growth on glucose alone ($q_{SMAX\_glucose}$ = -11.96 ± 0.43 $mmol_{GLCeq}$ g_{DM}^{-1} h^{-1}; **Table

16   2; Table S2**) for the yeast UFMG-CM-Y259. The remaining physiological parameters were

17   equivalent in both growth conditions.

18   Through another perspective, glucose and sucrose differently impact signaling cascades in the

19   cells, such as the one leading to the regulation of the protein kinase A (PKA) activity. A theory

20   related to the PKA signaling cascade has also been proposed to explain high specific growth

21   rates of *S. cerevisiae* on sucrose (Badotti et al. 2008; Marques et al. 2016). This cascade is

22   activated when the Gpr1-Gpa2 coupled receptor senses either glucose or sucrose, with the

23   affinity of Gpr1p being higher for the latter sugar (Lemaire et al. 2004). Once activated, the PKA

24   protein can exert all its function, including regulating the synthesis and degradation of storage

25   carbohydrates (Wingender-Drissen and Becker 1983; WERA et al. 1999; François et al. 2012),

26   the metabolic flux through the glycolytic pathway (Dihazi et al. 2003; Portela et al. 2006), and

27   gluconeogenesis (Mazón et al. 1982).

28   To this point, the mechanisms behind faster growth on sucrose over glucose remain an open

29   question. Further investigation, for instance by means of systems biology approaches, is needed

30   to elucidate sucrose regulation in the yeast *S. cerevisiae*.

18

S. cerevisiae **UFMG-CM-Y259 and JP1 display similar physiology during growth on sucrose or on fructose**

At last, we performed aerobic batch cultivations with fructose as the sole carbon and energy source (**Figure 6; Table 2; Supplementary Figures S1-S5, Supplementary Tables S1-S3**), again otherwise under identical conditions, when compared to all other cultivations described in this work. The physiological parameters were compared with those obtained from sucrose cultivations. The maximum specific growth rate between cultivations on the monosaccharide or on the disaccharide differed only for the CEN.PK113-7D strain, that grew faster when cultivated on the hexose ($\mu_{max\_Fructose}$ = 0.32 ± 0.01 h^{-1} and $\mu_{max\_Sucrose}$ = 0.21 ± 0.01 h^{-1}). The industrial (JP1) and the indigenous (UFMG-CM-Y259) strains' physiologies were equivalent when the carbon and energy sources sucrose or fructose are compared. Interestingly, the UFMG-CM-Y259 strain's physiology on fructose was more similar to that on sucrose than to the performance on glucose, contrasting the observations for the CEN.PK113-7D strain. This suggests that even the regulatory mechanisms triggered by hexoses in the yeast *S. cerevisiae* are strain-dependent.

Considering only glucose and fructose, transport has been pinpointed as a critical step for the different behaviors observed in the metabolism of such hexoses in *S. cerevisiae* (Luyten et al. 2002; Berthels et al. 2004). Among all the 20 proteins that constitute the group of hexose transporters and glucose sensors present in this microbe (Kruckeberg 1996; Özcan and Johnston 1999), Hxt1p to Hxt7p are the main ones in the context of glucose and/or fructose catabolism (Kruckeberg 1996; Luyten et al. 2002; Verwaal et al. 2002; Guillaume et al. 2007; Karpel et al. 2008). Expression of these proteins is strain dependent (Verwaal et al. 2002; PEREZ et al. 2005) and their substrate affinities vary from one transporter to the other (Ozcan and Johnston 1995). Enhanced fructose fermentation capacity has been demonstrated to be associated to the Hxt3p transporter (Guillaume et al. 2007). In the latter study (Guillaume et al. 2007), the molecular basis behind a higher fructose utilization capacity displayed by the commercial wine *S. cerevisiae* Fermichamp was revealed, in comparison to that of standard *S. cerevisiae* wine strains, namely the mutations T200A and G415N in the *HXT3* DNA sequence were responsible for such phenotype. Our sequence analysis of the *HXT3* open reading frame (ORF) did not show these same amino acid changes for any of the strains studied here (**Supplementary Figure S8**, **Supplementary Tables S7**). However, the T50A and V428C non-conservative amino acid substitutions were found in the *HXT3* ORF of the UFMG-CM-Y259 strain, and the K320Q

19

1 substitution in this ORF in JP1. These could be leads to be followed to verify whether these

2 *HXT3* alleles are capable of transporting fructose with different affinities and/or capacities.

3 Besides the transport step, hexose phosphorylation could also contribute to the different

4 growth physiologies exhibited by the UFMG-CM-Y259 strain on the monosaccharides, since it

5 is the first step in yeast glycolysis. *In vitro* measurements have shown that the proteins

6 hexokinase 1 and 2 target both monosaccharides (glucose or fructose), with lower affinity and

7 higher relative maximum velocity of reaction (V_{max}) for fructose (Lobo and Maitra 1977).

8 Glucokinase, on the other hand, is insensitive to fructose (Lobo and Maitra 1977). Among these

9 isozymes, hexokinase 2 is the major kinase in this glycolytic step, as its absence dramatically

10 changes *S. cerevisiae*'s physiology (Diderich et al. 2001). It is noteworthy that while the

11 conversion of fructose into fructose-6-phosphate takes only one step, to convert glucose into the

12 same metabolite two steps are required (phosphorylation and isomerization).

13

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Table 1. Relevant physiological parameters of *S. cerevisiae* CEN.PK113-7D, UFMG-CM-Y259, and JP1 during aerobic batch cultivations with either sucrose or a glucose/fructose mixture as sole carbon and energy source. All parameters were calculated for the exponential growth phase. Residual substrate refers to the remaining concentration of substrate at the end of the experiment. The data represent the mean of two experiments and the average deviation.

	CEN.PK113-7D		UFMG-CM-Y259		JP1	
	Sucrose	Glucose & Fructose	Sucrose	Glucose & Fructose	Sucrose	Glucose & Fructose
μ_{MAX} (h^{-1})	0.21 ± 0.01	0.26 ± 0.01	0.37 ± 0.01	0.35 ± 0.03	0.32 ± 0.00	0.30 ± 0.02
$Y_{X/S}$ (g$_{DM}$ gGLC$_{eq}$$^{-1}$)	0.14 ± 0.01	0.12 ± 0.00	0.16 ± 0.01	0.16 ± 0.01	0.14 ± 0.00	0.17 ± 0.01
$Y_{ethanol/S}$ (g gGLC$_{eq}$$^{-1}$)	0.36 ± 0.01	0.37 ± 0.00	0.41 ± 0.01	0.38 ± 0.00	0.41 ± 0.00	0.44 ± 0.02
qS_{MAX} (mmolGLC$_{eq}$ g$_{DM}$$^{-1}$ h^{-1})	-8.23 ± 0.37	-11.96 ± 0.02	-12.84 ± 0.21	-12.19 ± 0.63	-12.29 ± 0.12	-10.08 ± 0.39
qCO_2 (mmol g$_{DM}$$^{-1}$ h^{-1})	14.48 ± 0.22	19.48 ± 0.08	22.50 ± 2.00	20.96 ± 1.55	21.74 ± 0.19	17.53 ± 1.80
qO_2 (mmol g$_{DM}$$^{-1}$ h^{-1})	-2.96 ± 0.16	-2.56 ± 0.38	-3.34 ± 0.21	-2.88 ± 0.18	-2.19 ± 0.32	-2.15 ± 0.31
RQ (mmolCO$_2$ mmolO$_2$$^{-1}$)	4.91 ± 0.34	7.78 ± 1.19	6.81 ± 1.03	7.34 ± 1.00	10.13 ± 1.40	8.19 ± 0.33
Residual substrate (gGLC$_{eq}$ l^{-1})	0.10 ± 0.74	0.37 ± 0.94	1.03 ± 0.84	0.91 ± 0.40	1.79 ± 0.15	0.90 ± 0.11
Carbon recovery (%)	94.91 ± 2.86	93.75 ± 0.95	104.55 ± 2.95	98.80 ± 1.74	101.59 ± 0.11	107.25 ± 5.47
Electron balance (%)	95.35 ± 3.05	93.61 ± 0.44	107.25 ± 0.76	99.09 ± 0.96	102.13 ± 0.49	110.76 ± 5.34

Table 2. Relevant physiological parameters of *S. cerevisiae* CEN.PK113-7D, UFMG-CM-Y259, and JP1 during aerobic batch cultivations with either glucose or fructose as sole carbon and energy source. All parameters were calculated for the exponential growth phase. Residual substrate refers to the remaining substrate concentration at the end of the experiment. The data represent the mean of two experiments and the average deviation.

	CEN.PK113-7D		UFMG-CM-Y259		JP1	
	Glucose	Fructose	Glucose	Fructose	Glucose	Fructose
μ_{MAX} (h^{-1})	0.31 ± 0.01	0.32 ± 0.01	0.29 ± 0.00	0.36 ± 0.02	0.28 ± 0.02	0.32 ± 0.02
$Y_{X/S}$ (g$_{DM}$ gGLC$_{eq}$$^{-1}$)	0.13 ± 0.01	0.13 ± 0.00	0.14 ± 0.00	0.16 ± 0.00	0.13 ± 0.01	0.15 ± 0.01
$Y_{Ethanol/S}$ (g gGLC$_{eq}$$^{-1}$)	0.41 ± 0.01	0.38 ± 0.00	0.40 ± 0.03	0.37 ± 0.00	0.42 ± 0.00	0.38 ± 0.00
q_{SMAX} (mmolGLC$_{eq}$ g$_{DM}$$^{-1}$ h^{-1})	-13.22 ± 0.23	-13.46 ± 0.51	-11.96 ± 0.43	-12.12 ± 0.84	-11.67 ± 0.45	-12.24 ± 0.07
q_{CO_2} (mmol g$_{DM}$$^{-1}$ h^{-1})	23.57 ± 0.22	23.48 ± 0.34	20.77 ± 0.50	20.44 ± 1.39	19.75 ± 0.31	20.89 ± 0.05
q_{O_2} (mmol g$_{DM}$$^{-1}$ h^{-1})	-2.45 ± 0.14	-2.84 ± 0.11	-2.63 ± 0.31	-3.03 ± 0.12	-2.12 ± 0.09	-2.43 ± 0.02
RQ (mmolCO$_2$ mmolO$_2$$^{-1}$)	9.66 ± 0.66	8.28 ± 0.19	8.01 ± 1.12	6.73 ± 0.20	9.31 ± 0.25	8.60 ± 0.04
Residual substrate (gGLC$_{eq}$.l^{-1})	0.15 ± 0.12	2.72 ± 1.35	0.03 ± 0.03	2.28 ± 0.19	0.73 ± 0.08	1.34 ± 0.10
Carbon recovery (%)	100.73 ± 0.07	97.57 ± 0.66	103.25 ± 6.11	99.04 ± 0.94	101.44 ± 1.35	97.12 ± 0.51
Electron balance (%)	101.14 ± 0.70	97.41 ± 0.01	99.05 ± 2.05	100.00 ± 0.96	104.06 ± 0.53	97.25 ± 0.66

Figure 1. Schematic illustration of the two mechanisms of sucrose utilization in *S. cerevisiae*: A) periplasmic hydrolysis by invertase (Suc) and passive transport of glucose and fructose into the cells; B) active sucrose transport via the **A**lpha-**G**lucoside **T**ransporter (Agt1) and ATP-dependent export of H$^+$ via the **P**lasma **M**embrane **AT**Pase (Pma1).

Figure 2. Substrate and metabolites concentrations/amounts during aerobic batch cultivation of *S. cerevisiae* CEN.PK113-7D (▲), JP1 (♦), and UFMG-CM-Y259 (■) cells <u>with sucrose as sole carbon and energy source</u>. Substrate represents the sum of sucrose, glucose and fructose concentrations. Dashed lines represent trend lines. One representative dataset of duplicate independent experiments is shown.

Figure 3. Sugar concentrations and periplasmic invertase activity during aerobic batch cultivation of *S. cerevisiae* CEN.PK113-7D (A), UFMG-CM-Y259 (B) and JP1 (C) cells on synthetic medium with <u>sucrose as sole carbon and energy source</u>. Sucrose (●); glucose (■); fructose (*closed* ∇); invertase (▲). Experiments were performed in duplicate. One representative dataset of duplicate independent experiments is shown.

Figure 4. Substrate and metabolites concentrations/amounts during aerobic batch cultivation of *S. cerevisiae* CEN.PK113-7D (▲), JP1 (♦), and UFMG-CM-Y259 (■) cells with an <u>equimolar mixture of glucose and fructose as sole carbon and energy source</u>. Substrate represent the sum of glucose and fructose concentrations. Dashed lines represent trend lines. One representative dataset of duplicate independent experiments is shown.

Figure 5. Substrate and metabolites concentrations/amounts during aerobic batch cultivation of *S. cerevisiae* CEN.PK113-7D (▲), JP1 (♦), and UFMG-CM-Y259 (■) cells with <u>glucose as sole carbon and energy source</u>. Dashed lines represent trend lines. One representative dataset of duplicate independent experiments is shown.

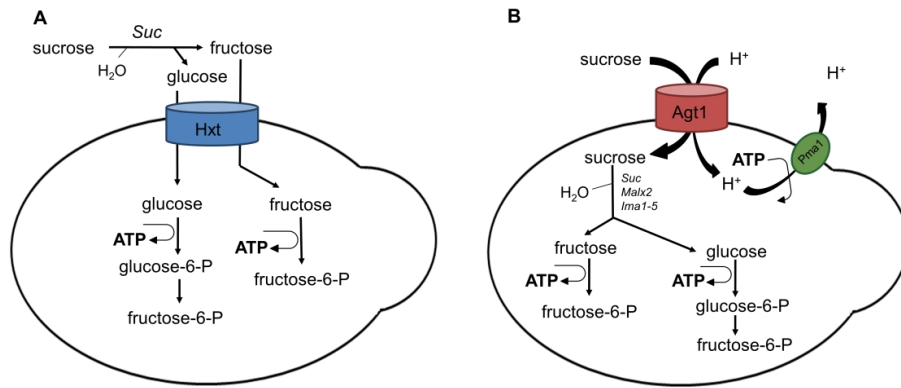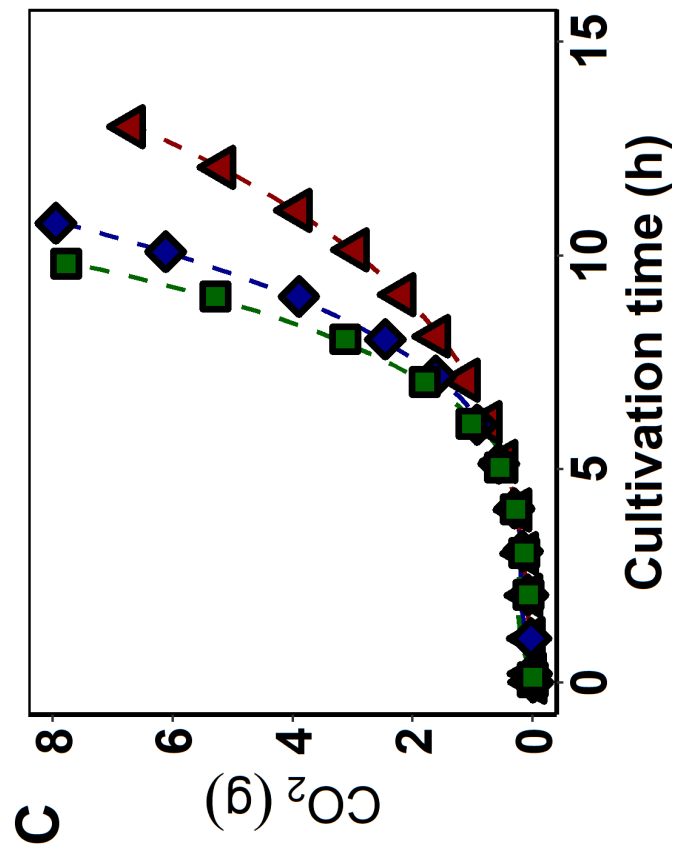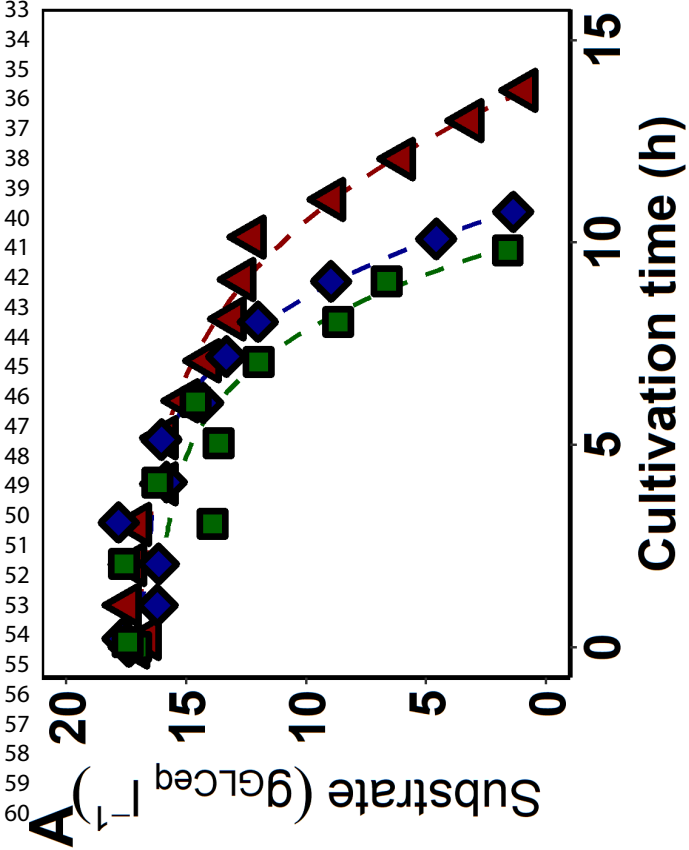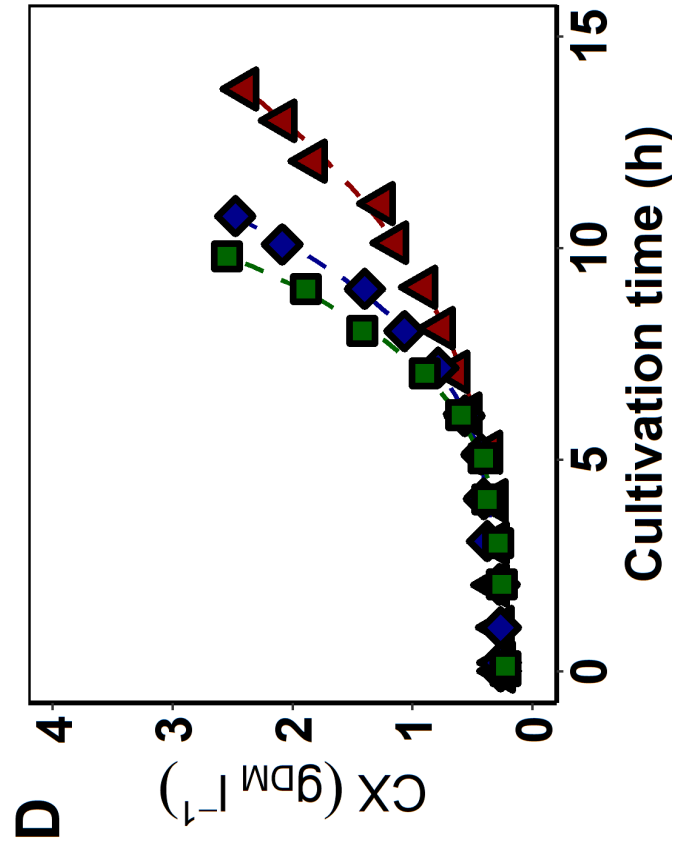**Figure 6.** Substrate and metabolites concentrations/amounts during aerobic batch cultivation of *S. cerevisiae* CEN.PK113-7D (▲), JP1 (♦), and UFMG-CM-Y259 (■) cells with <u>fructose as sole carbon and energy source</u>. Dashed lines represent trend lines. One representative dataset of duplicate independent experiments is shown.
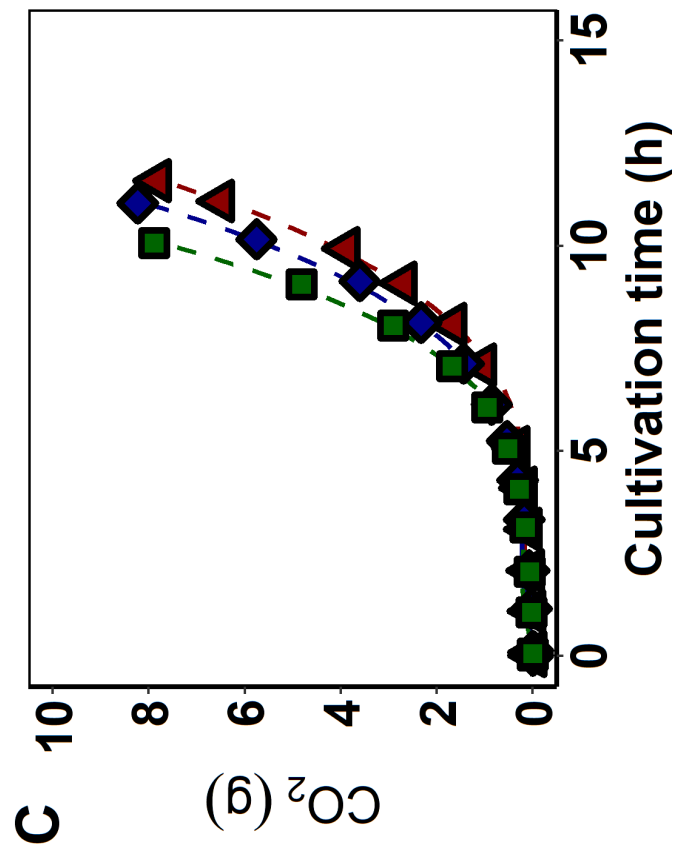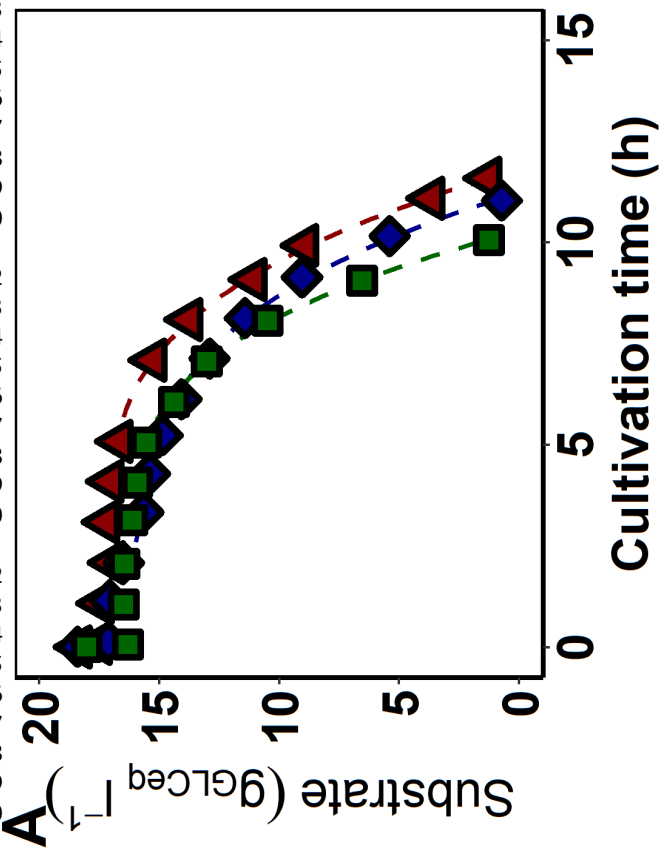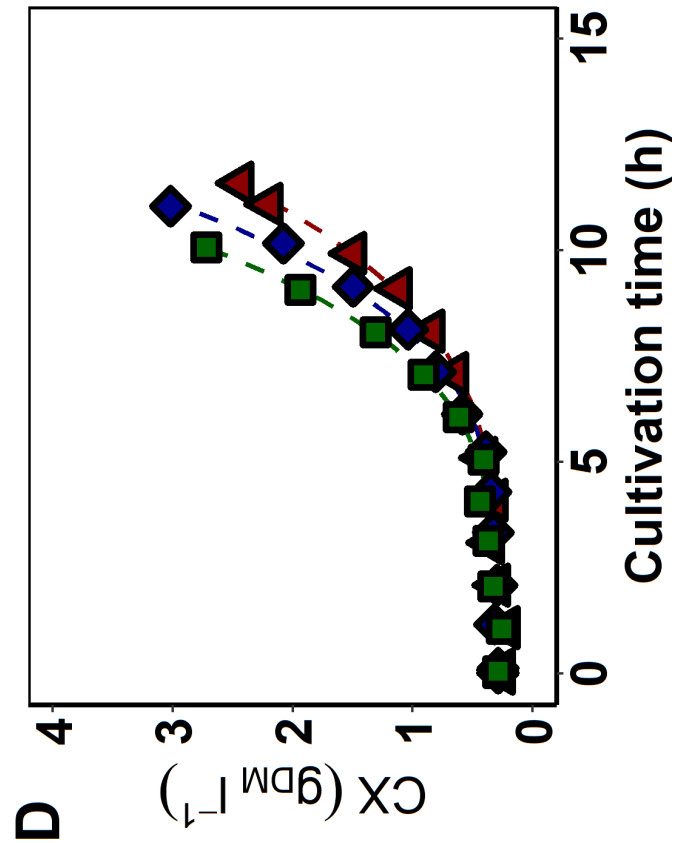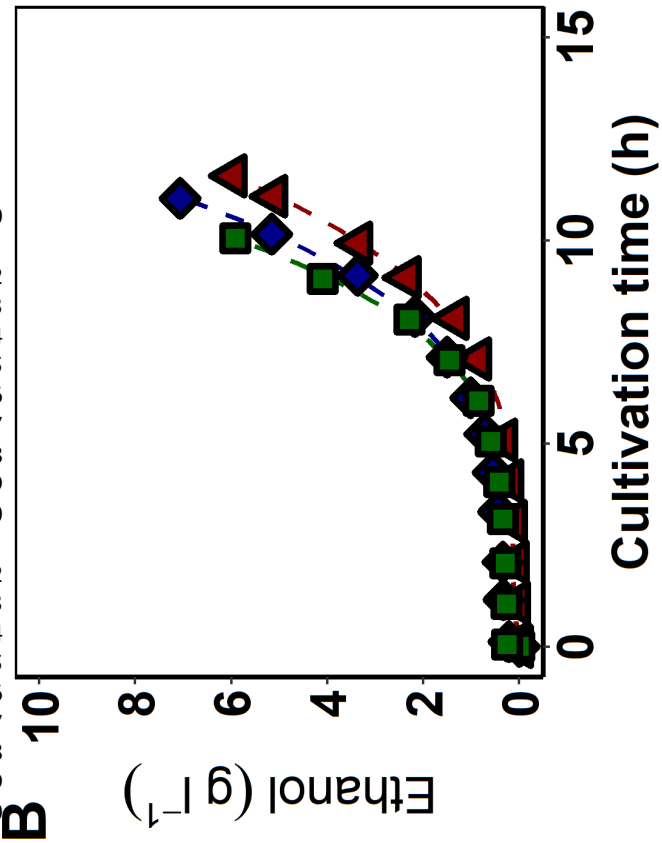
Figure 1. Schematic illustration of the two mechanisms of sucrose utilization in *S. cerevisiae*: A) periplasmic hydrolysis by invertase (Suc) and passive transport of glucose and fructose into the cells; B) active sucrose transport via the Alpha-Glucoside Transporter (Agt1) and ATP-dependent export of H$^+$ via the Plasma Membrane ATPase (Pma1).
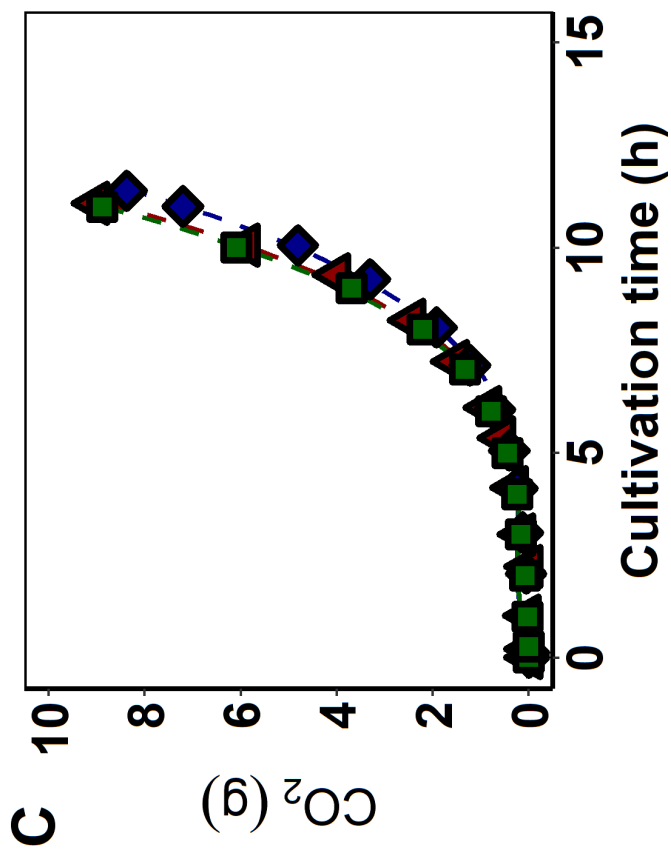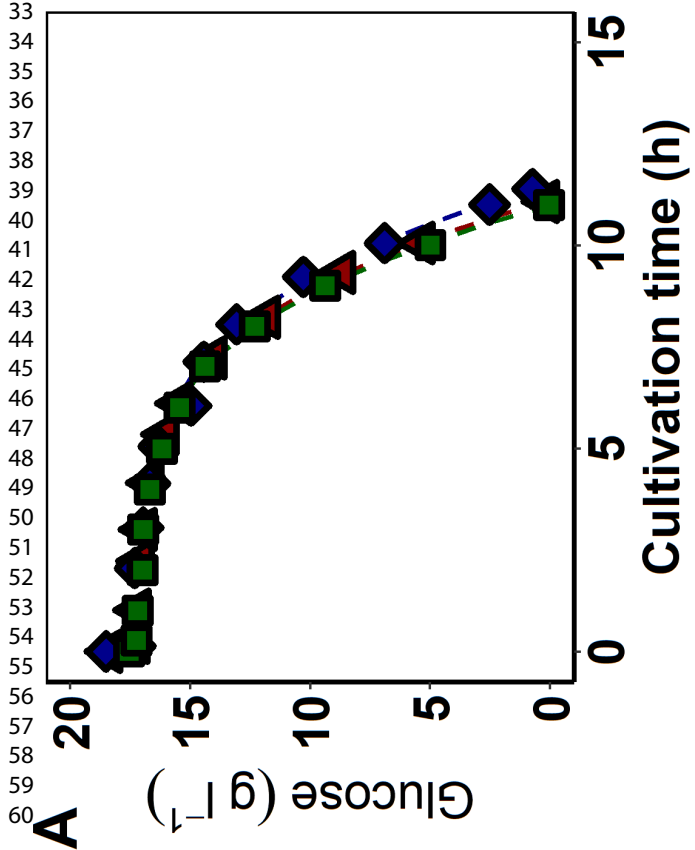
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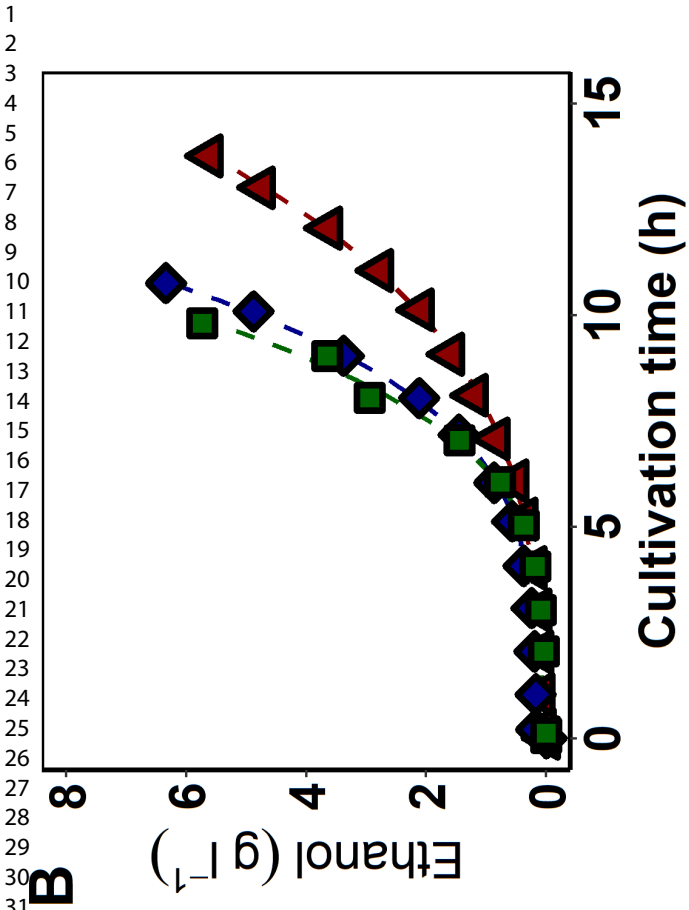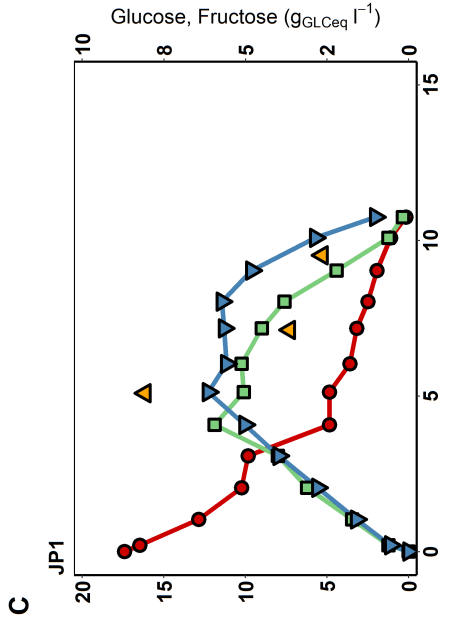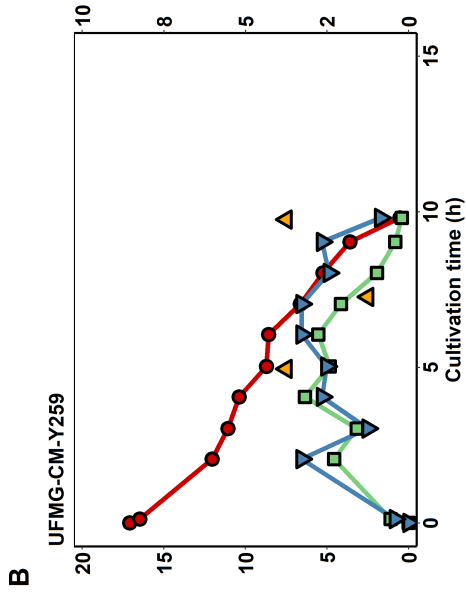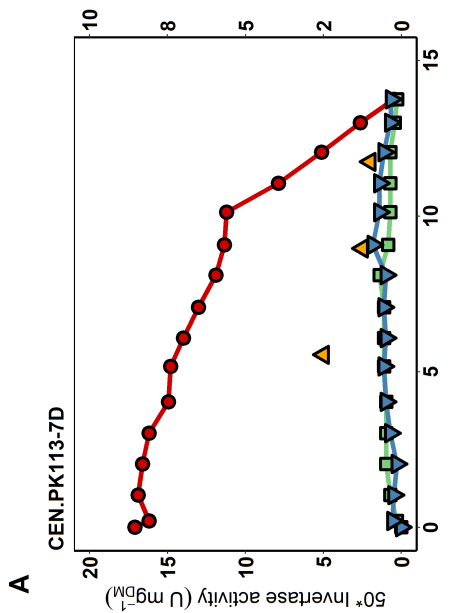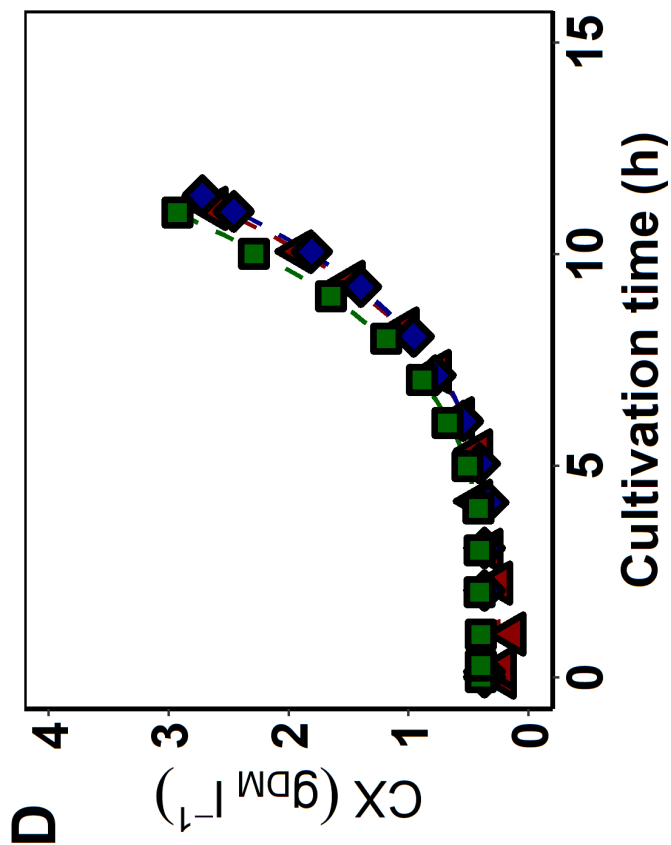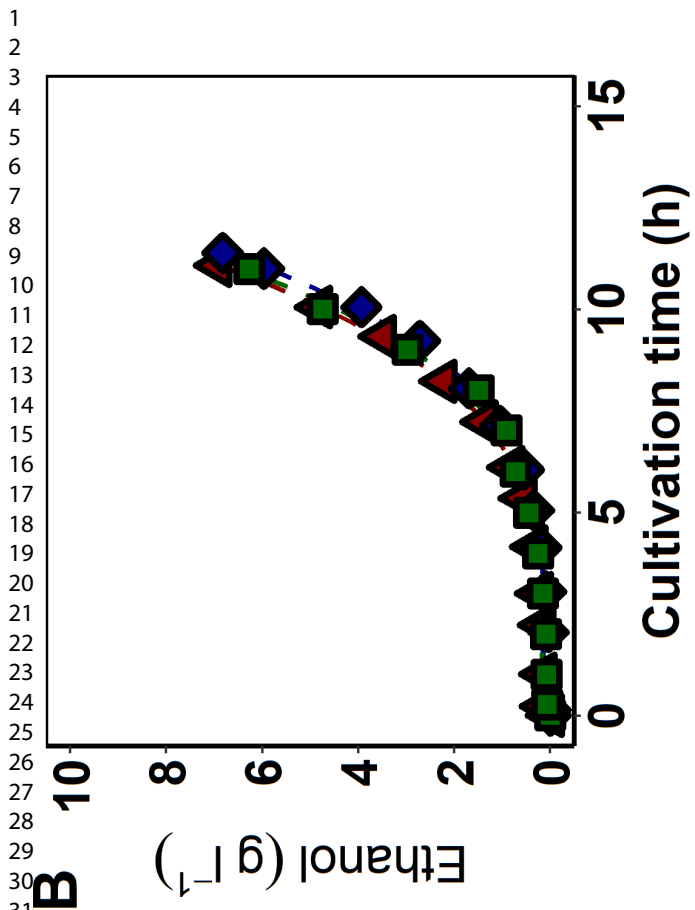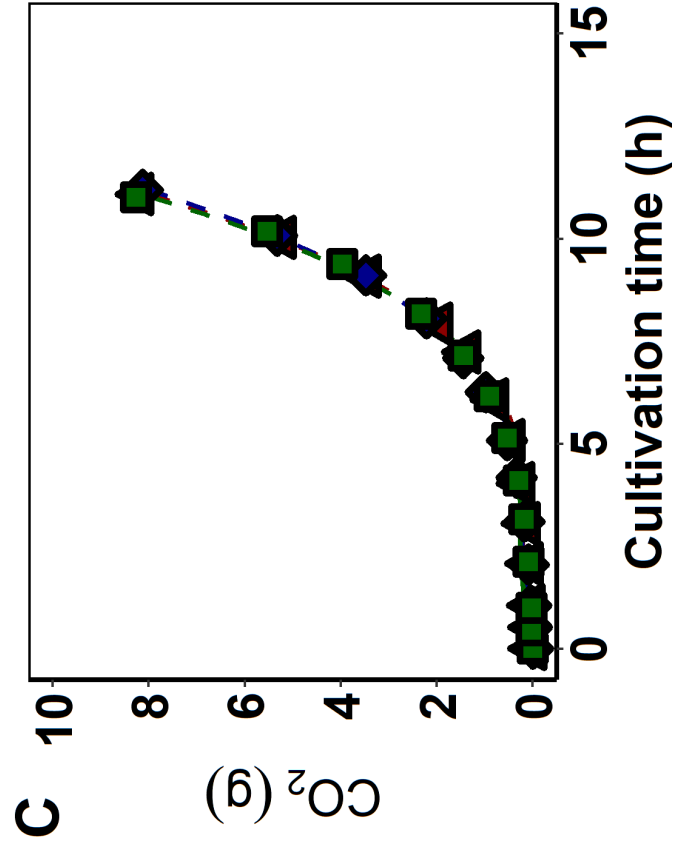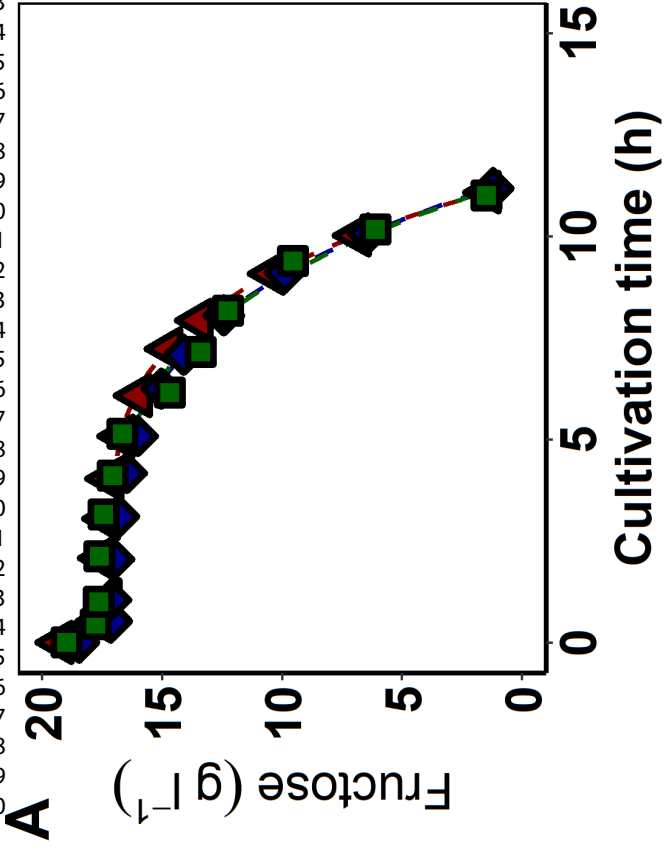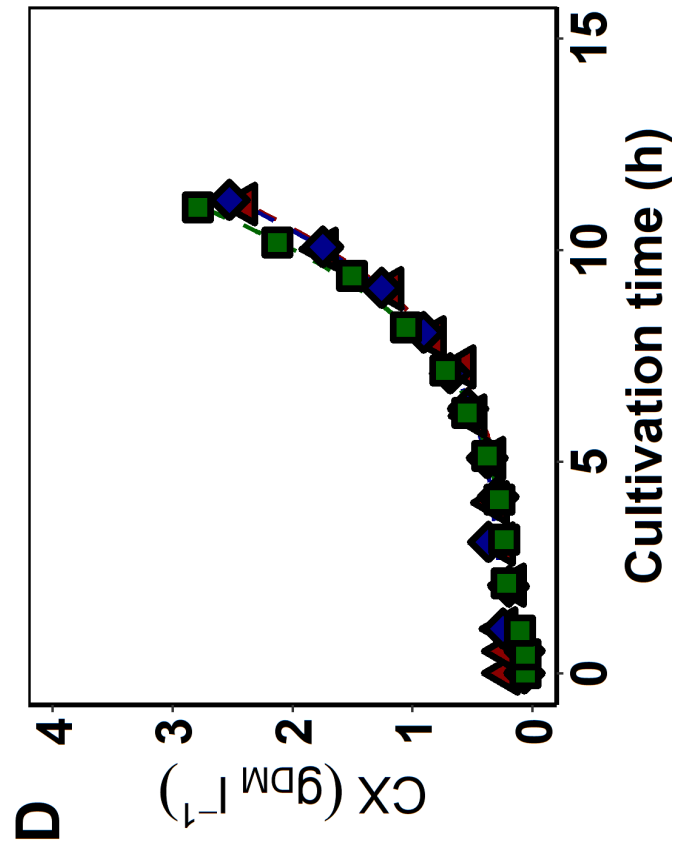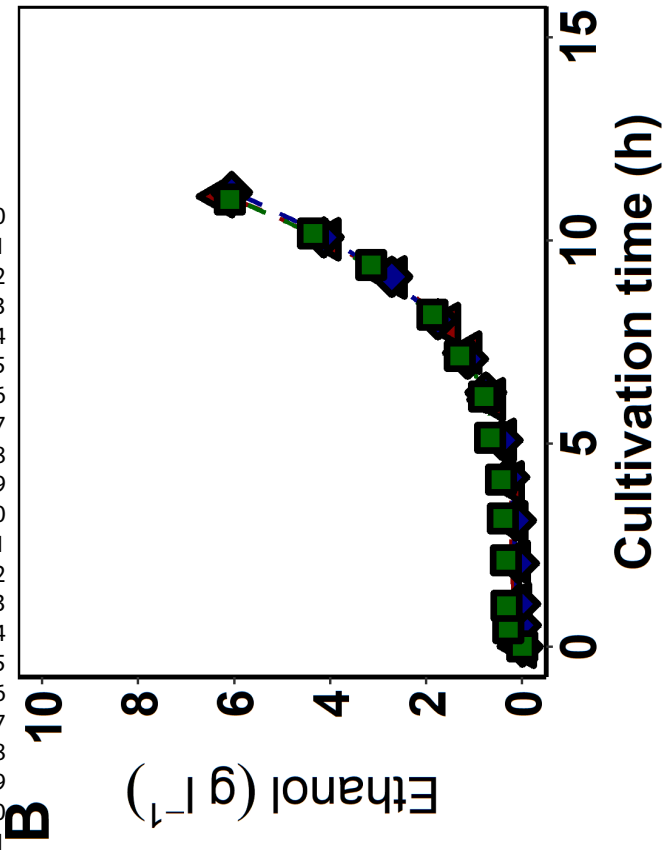
Supplementary Material (Figures and Tables)

Figure S1. Percentage of substrate metabolized via the fermentative pathway by *S. cerevisiae* CEN.PK113-7D, UFMG-CM-Y259, and JP1 during aerobic growth on sucrose as sole carbon and energy source. Error bars represent the average deviation of the values obtained from duplicate experiments.
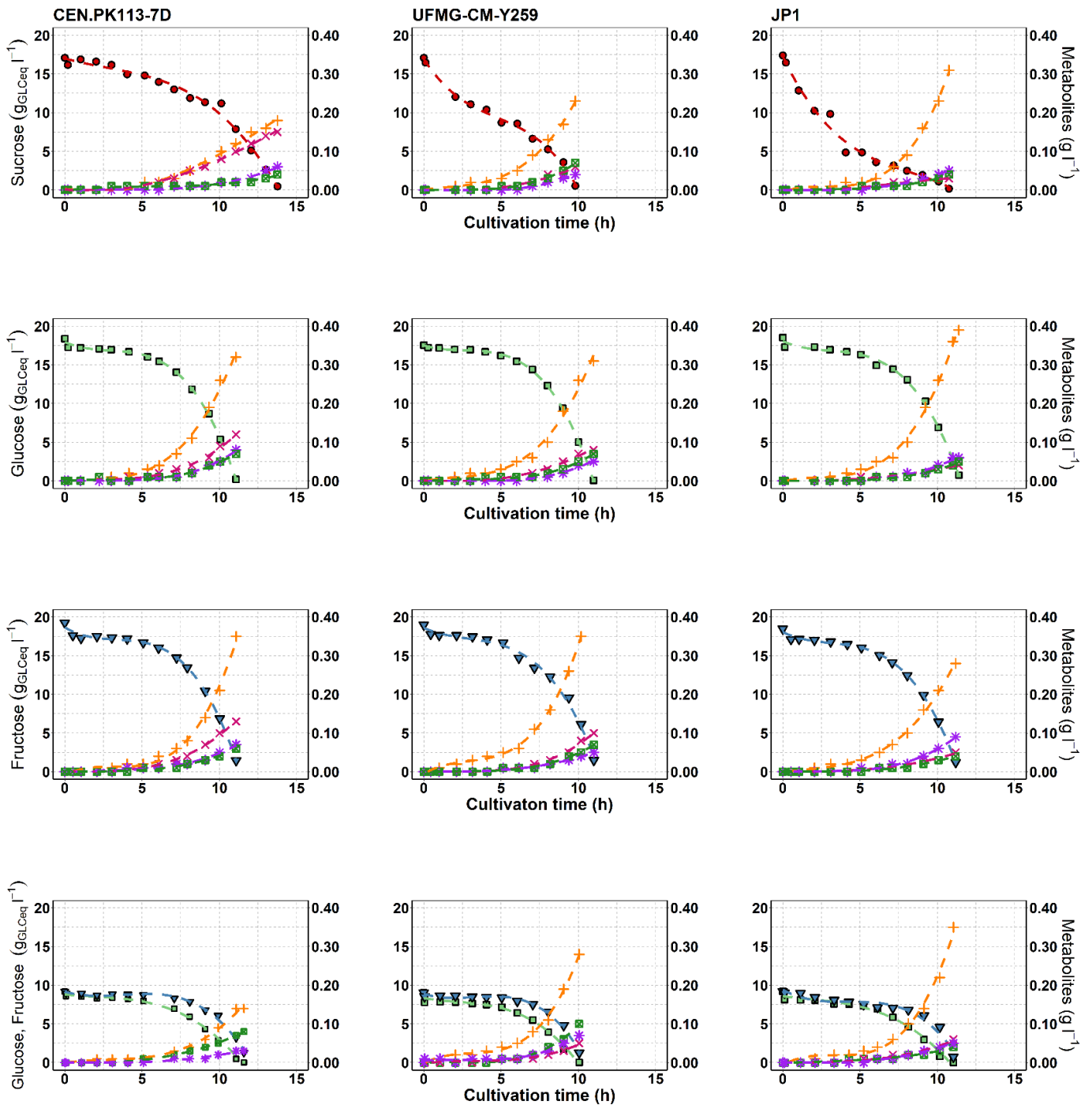
Figure S2. Substrate and metabolites concentrations during aerobic batch cultivation of S. cerevisiae CEN.PK113-7D, UFMG-CM-Y259 and JP1 with either sucrose, glucose, fructose or an equimolar mixture of glucose and fructose as sole carbon and energy source. Sucrose (●); Glucose

(■), Fructose (*closed* ∇); Glycerol (+); Acetate (X); Succinate (*); Lactate (crossed □). Dashed lines represent trend lines. Experiments were performed in duplicate. Data shown in the plots are from a single growth experiment.
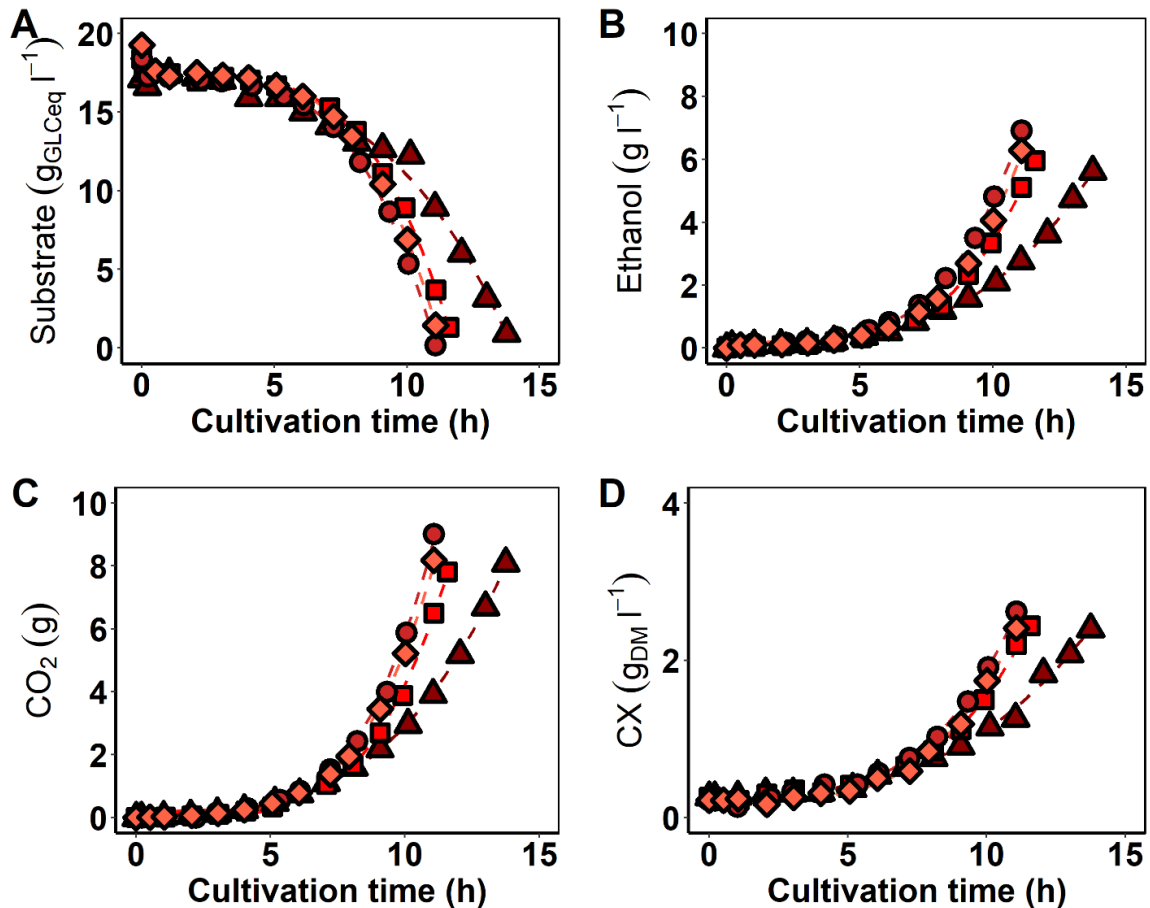
Figure S3. Substrate and metabolites concentrations/amounts during aerobic batch cultivation of <u>*S. cerevisiae*</u> CEN.PK113-7D with sucrose (▲), an equimolar mixture of glucose and fructose (■), glucose (●), or fructose (♦) as sole carbon and energy source. For the glucose + fructose experiment, substrate represents the sum of glucose and fructose concentrations. Dashed lines represent trend lines. Experiments were performed in duplicate. Data shown in the plots are from a single growth experiment.
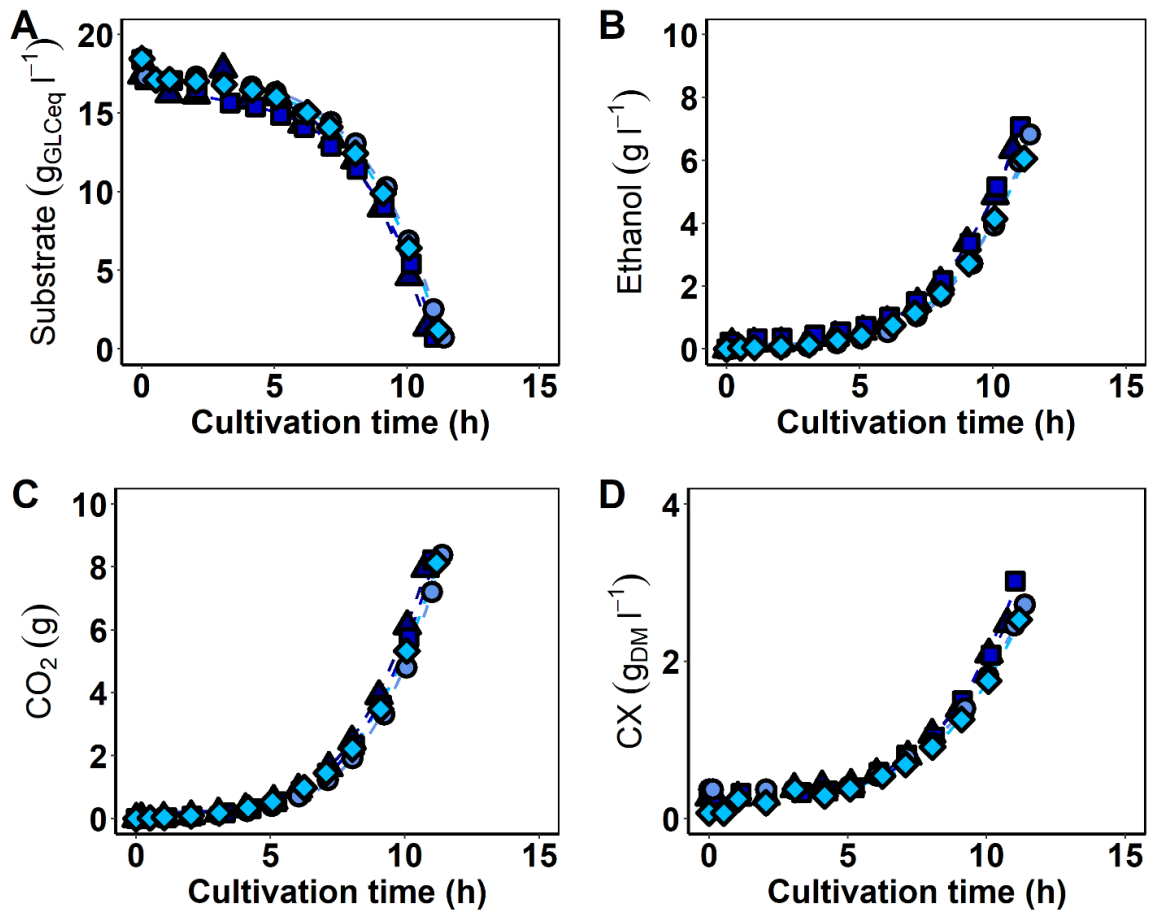
Figure S4. Substrate and metabolites concentrations/amounts during aerobic batch cultivation of *S. cerevisiae* JP1 with sucrose (▲), an equimolar mixture of glucose and fructose (■), glucose (●), or fructose (♦) as sole carbon and energy source. For the glucose + fructose experiment, substrate represents the sum of glucose and fructose concentrations. Dashed lines represent trend lines. Experiments were performed in duplicate. Data shown in the plots are from a single growth experiment.
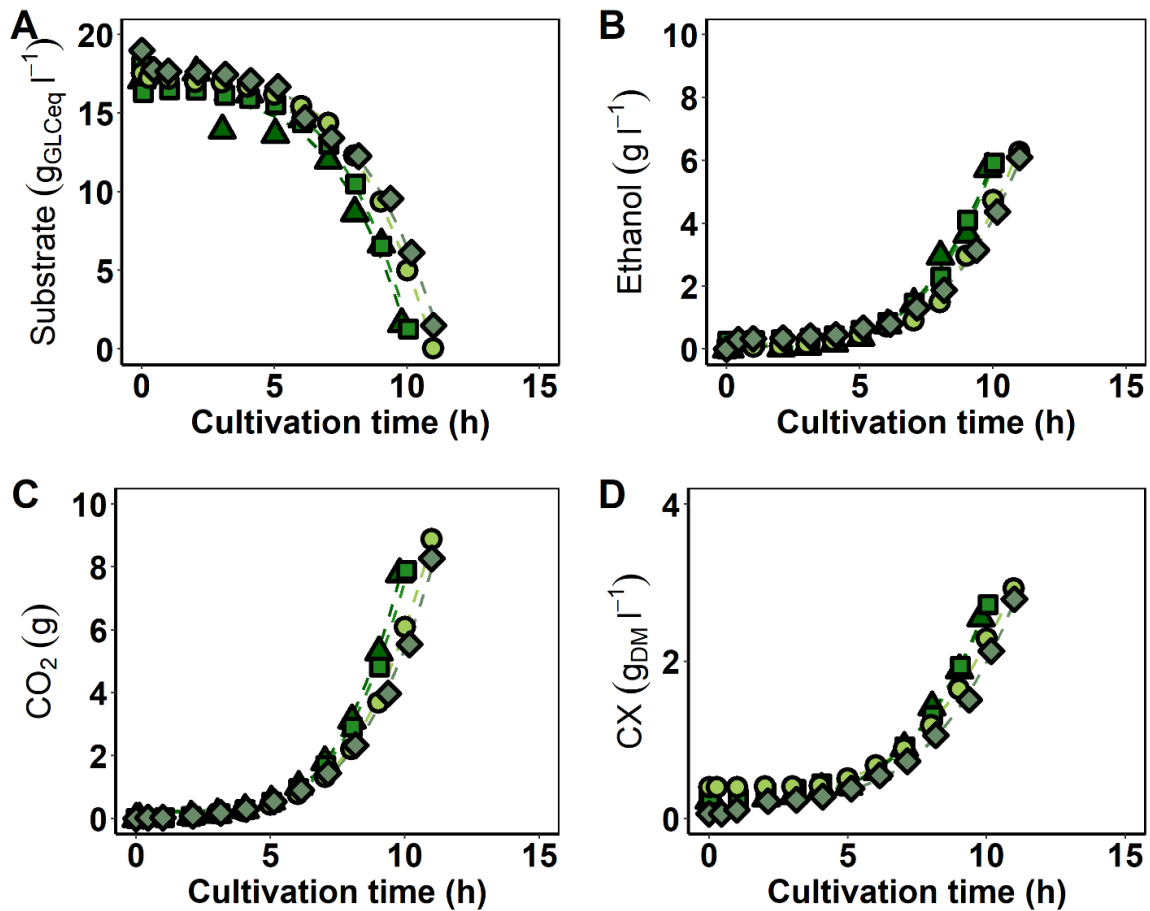
Figure S5. Substrate and metabolites concentrations/amounts during aerobic batch cultivation of _S. cerevisiae_ UFMG-CM-Y259 with sucrose (▲), an equimolar mixture of glucose and fructose (■), glucose (●), or fructose (♦) as sole carbon and energy source. For the glucose + fructose experiment, substrate represents the sum of glucose and fructose concentrations. Dashed lines represent trend lines. Experiments were performed in duplicate. Data shown in the plots are from a single growth experiment.

Table S1. Physiological parameters of *S. cerevisiae* CEN.PK113-7D during aerobic batch cultivations with either glucose, sucrose, an equimolar mixture of glucose and fructose, or fructose as sole carbon and energy source. All parameters were calculated for the exponential growth phase. The data represent the mean of two experiments and the average deviation.

	Strain: CEN.PK113-7D										
	Glucose			Sucrose			Glucose & Fructose			Fructose		
μ_{MAX} (h^{-1})	0.31	±	0.01	0.21	±	0.01	0.26	±	0.01	0.32	±	0.01
$Y_{X/S}$ (g_{DM} $gGLC_{eq}^{-1}$)	0.13	±	0.01	0.14	±	0.01	0.12	±	0.00	0.13	±	0.00
$Y_{Ethanol/S}$ (g $gGLC_{eq}^{-1}$)	0.41	±	0.01	0.36	±	0.01	0.37	±	0.00	0.38	±	0.00
$Y_{CO2/S}$ (g $gGLC_{eq}^{-1}$)	0.44	±	0.00	0.43	±	0.01	0.40	±	0.00	0.43	±	0.01
$Y_{glycerol/S}$ (g $gGLC_{eq}^{-1}$)	0.02	±	0.00	0.01	±	0.00	0.02	±	0.01	0.03	±	0.00
$Y_{acetate/S}$ (g $gGLC_{eq}^{-1}$)	0.01	±	0.00	0.01	±	0.00	0.01	±	0.00	0.01	±	0.00
$Y_{succinate/S}$ (g $gGLC_{eq}^{-1}$)	0.00	±	0.00	0.00	±	0.00	0.00	±	0.00	0.00	±	0.00
$Y_{lactate/S}$ (g $gGLC_{eq}^{-1}$)	0.00	±	0.00	0.00	±	0.00	0.00	±	0.00	0.00	±	0.00
qS_{MAX} ($mmolGLC_{eq}$ g_{DM}^{-1} h^{-1})	-13.22	±	0.23	-8.23	±	0.37	-11.96	±	0.02	-13.46	±	0.51
qCO_2 (mmol g_{DM}^{-1} h^{-1})	23.57	±	0.22	14.48	±	0.22	19.48	±	0.08	23.48	±	0.34
qO_2 (mmol g_{DM}^{-1} h^{-1})	-2.45	±	0.14	-2.96	±	0.16	-2.56	±	0.38	-2.84	±	0.11
RQ ($mmolCO_2$ $mmolO_2^{-1}$)	9.66	±	0.66	4.91	±	0.34	7.78	±	1.19	8.28	±	0.19
$qEthanol$ (mmol g_{DM}^{-1} h^{-1})	21.03	±	0.74	11.43	±	0.34	17.46	±	0.09	20.08	±	0.81
$qglycerol$ (mmol g_{DM}^{-1} h^{-1})	0.53	±	0.02	0.19	±	0.01	0.37	±	0.12	0.72	±	0.01
$qacetate$ (mmol g_{DM}^{-1} h^{-1})	0.21	±	0.05	0.32	±	0.06	0.28	±	0.12	0.26	±	0.03
$qsuccinate$ (mmol g_{DM}^{-1} h^{-1})	0.08	±	0.00	0.04	±	0.01	0.05	±	0.02	0.07	±	0.02
$qlactate$ (mmol g_{DM}^{-1} h^{-1})	0.13	±	0.06	0.02	±	0.02	0.05	±	0.00	0.08	±	0.01
Residual substrate ($gGLC_{eq}$ l^{-1})	0.15	±	0.12	0.10	±	0.74	0.37	±	0.94	2.72	±	1.35
Carbon recovery (%)	100.73	±	0.07	94.91	±	2.86	93.75	±	0.95	97.57	±	0.66
Electron balance (%)	101.14	±	0.70	95.35	±	3.05	93.61	±	0.44	97.41	±	0.01

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Table S2. Physiological parameters of *S. cerevisiae* JP1 during aerobic batch cultivations with either glucose, sucrose, an equimolar mixture of glucose and fructose, or fructose as sole carbon and energy source. All parameters were calculated for the exponential growth phase. The data represent the mean of two experiments and the average deviation.

	Strain: JP1			
	Glucose	Sucrose	Glucose & Fructose	Fructose
μ_{MAX} (h^{-1})	0.28 ± 0.02	0.32 ± 0.00	0.30 ± 0.02	0.32 ± 0.02
$Y_{X/S}$ (g$_{DM}$ gGLC$_{eq}^{-1}$)	0.13 ± 0.01	0.14 ± 0.00	0.17 ± 0.01	0.15 ± 0.01
$Y_{Ethanol/S}$ (g gGLC$_{eq}^{-1}$)	0.42 ± 0.00	0.41 ± 0.00	0.44 ± 0.02	0.38 ± 0.00
$Y_{CO2/S}$ (g gGLC$_{eq}^{-1}$)	0.41 ± 0.01	0.43 ± 0.00	0.42 ± 0.03	0.42 ± 0.00
$Y_{glycerol/S}$ (g gGLC$_{eq}^{-1}$)	0.02 ± 0.00	0.02 ± 0.00	0.02 ± 0.00	0.02 ± 0.00
$Y_{acetate/S}$ (g gGLC$_{eq}^{-1}$)	0.00 ± 0.00	0.00 ± 0.00	0.00 ± 0.00	0.00 ± 0.00
$Y_{succinate/S}$ (g gGLC$_{eq}^{-1}$)	0.00 ± 0.00	0.00 ± 0.00	0.00 ± 0.00	0.00 ± 0.00
$Y_{lactate/S}$ (g gGLC$_{eq}^{-1}$)	0.00 ± 0.00	0.00 ± 0.00	0.00 ± 0.00	0.00 ± 0.00
qS_{MAX} (mmolGLC$_{eq}$ g$_{DM}^{-1}$ h^{-1})	-11.67 ± 0.45	-12.29 ± 0.12	-10.08 ± 0.39	-12.24 ± 0.07
qCO_2 (mmol g$_{DM}^{-1}$ h^{-1})	19.75 ± 0.31	21.74 ± 0.19	17.53 ± 1.80	20.89 ± 0.05
qO_2 (mmol g$_{DM}^{-1}$ h^{-1})	-2.12 ± 0.09	-2.19 ± 0.32	-2.15 ± 0.31	-2.43 ± 0.02
RQ (mmolCO$_2$ mmolO$_2^{-1}$)	9.31 ± 0.25	10.13 ± 1.40	8.19 ± 0.33	8.60 ± 0.04
$q_{Ethanol}$ (mmol g$_{DM}^{-1}$ h^{-1})	19.23 ± 0.83	19.64 ± 0.26	17.18 ± 1.54	18.23 ± 0.14
$q_{glycerol}$ (mmol g$_{DM}^{-1}$ h^{-1})	0.52 ± 0.08	0.48 ± 0.03	0.42 ± 0.00	0.39 ± 0.07
$q_{acetate}$ (mmol g$_{DM}^{-1}$ h^{-1})	0.17 ± 0.09	0.08 ± 0.00	0.08 ± 0.00	0.10 ± 0.01
$q_{succinate}$ (mmol g$_{DM}^{-1}$ h^{-1})	0.07 ± 0.00	0.06 ± 0.01	0.04 ± 0.02	0.05 ± 0.01
$q_{lactate}$ (mmol g$_{DM}^{-1}$ h^{-1})	0.07 ± 0.01	0.05 ± 0.01	0.06 ± 0.00	0.07 ± 0.02
Residual substrate (gGLC$_{eq}$ l^{-1})	0.73 ± 0.08	1.79 ± 0.15	0.90 ± 0.11	1.34 ± 0.10
Carbon recovery (%)	101.44 ± 1.35	101.59 ± 0.11	107.25 ± 5.47	97.12 ± 0.51
Electron balance (%)	104.06 ± 0.53	102.13 ± 0.49	110.76 ± 5.34	97.25 ± 0.66

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Table S3. . Physiological parameters of _S. cerevisiae_ UFMG-CM-Y259 during aerobic batch cultivations with either glucose, sucrose,
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an equimolar mixture of glucose and fructose, or fructose as sole carbon and energy source. All parameters were calculated for the
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exponential growth phase. The data represent the mean of two experiments and the average deviation.
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	Strain: UFMG-CM-Y259										
	Glucose			Sucrose			Glucose & Fructose			Fructose		
μ_{MAX} (h^{-1})	0.29	±	0.00	0.37	±	0.01	0.35	±	0.03	0.36	±	0.02
$Y_{X/S}$ (g$_{DM}$ gGLC$_{eq}^{-1}$)	0.14	±	0.00	0.16	±	0.01	0.16	±	0.01	0.16	±	0.00
$Y_{Ethanol/S}$ (g gGLC$_{eq}^{-1}$)	0.40	±	0.03	0.41	±	0.01	0.38	±	0.00	0.37	±	0.00
$Y_{CO2/S}$ (g gGLC$_{eq}^{-1}$)	0.45	±	0.02	0.43	±	0.05	0.42	±	0.01	0.41	±	0.00
$Y_{glycerol/S}$ (g gGLC$_{eq}^{-1}$)	0.02	±	0.00	0.02	±	0.00	0.02	±	0.00	0.03	±	0.01
$Y_{acetate/S}$ (g gGLC$_{eq}^{-1}$)	0.01	±	0.00	0.00	±	0.00	0.01	±	0.00	0.01	±	0.00
$Y_{succinate/S}$ (g gGLC$_{eq}^{-1}$)	0.00	±	0.00	0.00	±	0.00	0.00	±	0.00	0.00	±	0.00
$Y_{lactate/S}$ (g gGLC$_{eq}^{-1}$)	0.00	±	0.00	0.00	±	0.00	0.00	±	0.00	0.00	±	0.00
qS_{MAX} (mmolGLC$_{eq}$ g$_{DM}^{-1}$ h^{-1})	-11.96	±	0.43	-12.84	±	0.21	-12.19	±	0.63	-12.12	±	0.84
qCO_2 (mmol g$_{DM}^{-1}$ h^{-1})	20.77	±	0.50	22.50	±	2.00	20.96	±	1.55	20.44	±	1.39
qO_2 (mmol g$_{DM}^{-1}$ h^{-1})	-2.63	±	0.31	-3.34	±	0.21	-2.88	±	0.18	-3.03	±	0.12
RQ (mmolCO$_2$ mmolO$_2^{-1}$)	8.01	±	1.12	6.81	±	1.03	7.34	±	1.00	6.73	±	0.20
$q_{Ethanol}$ (mmol g$_{DM}^{-1}$ h^{-1})	17.63	±	0.76	20.82	±	0.65	18.00	±	1.06	17.57	±	1.26
$q_{glycerol}$ (mmol g$_{DM}^{-1}$ h^{-1})	0.47	±	0.07	0.48	±	0.05	0.40	±	0.07	0.60	±	0.13
$q_{acetate}$ (mmol g$_{DM}^{-1}$ h^{-1})	0.21	±	0.03	0.15	±	0.01	0.20	±	0.03	0.30	±	0.07
$q_{succinate}$ (mmol g$_{DM}^{-1}$ h^{-1})	0.05	±	0.00	0.05	±	0.00	0.08	±	0.03	0.07	±	0.00
$q_{lactate}$ (mmol g$_{DM}^{-1}$ h^{-1})	0.10	±	0.02	0.12	±	0.01	0.12	±	0.03	0.09	±	0.00
Residual substrate (gGLC$_{eq}$ l^{-1})	0.03	±	0.03	1.03	±	0.84	0.91	±	0.40	2.28	±	0.19
Carbon recovery (%)	103.25	±	6.11	104.55	±	2.95	98.80	±	1.74	99.04	±	0.94
Electron balance (%)	99.05	±	2.05	107.25	±	0.76	99.09	±	0.96	100.00	±	0.96

Table S4. Point mutations in the *SUC2* sequence of *S. cerevisiae* strains JP1 and UFMG-CM-Y259 with respect to the reference strain S288C. Nucleotide and aminoacid (aa) changes are indicated in red and blue, respectively. Colored in grey are mutations shared by the two strains.

Gene: *SUC2*		Accession number: V01311			
Mutation					
Position (bp)	**Original codon**	**aa**	**Mutated codon**	**aa**	**Type**
JP1					
228	GCT	A	ACT	T	Missense
249	AAT	N	CAT	H	Missense
261	CAA	Q	GAA	E	Missense
338	ACG	T	ACC	T	Silent
386	CGC	A	GCA	A	Silent
437	TCT	S	TCC	S	Silent
599	TCT	S	TCG	S	Silent
621	CTA	L	TTA	L	Silent
641	AAC	N	AAT	N	Silent
665	TAC	Y	TAT	Y	Silent
716	TCT	S	TCC	S	Silent
866	TTG	L	TTA	L	Silent
1082	GGT	G	GGC	G	Silent
1085	CCC	P	CCA	P	Silent
1103	ACT	T	ACC	T	Silent
1190	GTT	V	GTC	V	Silent
1223	TTT	F	TTC	F	Silent
1224	GCC	A	CCC	P	Missense
1544	ACT	T	ACC	T	Silent
UFMG-CM-Y259					
135	TTG	L	CTG	L	Silent
249	AAT	N	CAT	H	Missense
261	CAA	Q	GAA	E	Missense
338	ACG	T	ACC	T	Silent
386	GCG	A	GCA	A	Silent
437	TCT	S	TCC	S	Silent
473	AAG	K	AAA	K	Silent
641	AAC	N	AAT	N	Silent
1082	GGT	G	GGC	G	Silent
1103	ACT	T	ACC	T	Silent
1334	GTC	V	GTT	V	Silent

Strain		*SUC2* sequence	
JP1	1	MLLQAFLFLLAGFAAKISASMTNETSDRPLVHFTPNKGWMNDPNGLWYDE	50
UFMG-CM-Y259		MLLQAFLFLLAGFAAKISASMTNETSDRPLVHFTPNKGWMNDPNGLWYDE	
CEN.PK113-7D		MLLQAFLFLLAGFAAKISASMTNETSDRPLVHFTPNKGWMNDPNGLWYDE	
S288C		MLLQAFLFLLAGFAAKISASMTNETSDRPLVHFTPNKGWMNDPNGLWYDE	
	51	KDAKWHLYFQYNPNDTVWGTPLFWGHTTSDDLTHWEDEPIAIAPKRNDSG	100
		KDAKWHLYFQYNPNDTVWGTPLFWGHATSDDLTHWEDEPIAIAPKRNDSG	
		KDAKWHLYFQYNPNDTVWGTPLFWGHATSDDLTNWEDQPIAIAPKRNDSG	
		KDAKWHLYFQYNPNDTVWGTPLFWGHATSDDLTNWEDQPIAIAPKRNDSG	
	101	AFSGSMVVDYNNTSGFFNDTIDPRQRCVAIWTYNTPESEEQYISYSLDGG	150
		AFSGSMVVDYNNTSGFFNDTIDPRQRCVAIWTYNTPESEEQYISYSLDGG	
		AFSGSMVVDYNNTSGFFNDTIDPRQRCVAIWTYNTPESEEQYISYSLDGG	
		AFSGSMVVDYNNTSGFFNDTIDPRQRCVAIWTYNTPESEEQYISYSLDGG	
	151	YTFTEYQKNPVLAANSTQFRDPKVFWYEPSQKWIMTAAKSQDYKIEIYSS	200
		YTFTEYQKNPVLAANSTQFRDPKVFWYEPSQKWIMTAAKSQDYKIEIYSS	
		YTFTEYQKNPVLAANSTQFRDPKVFWYEPSQKWIMTAAKSQDYKIEIYSS	
		YTFTEYQKNPVLAANSTQFRDPKVFWYEPSQKWIMTAAKSQDYKIEIYSS	
	201	DDLKSWKLESAFANEGFLGYQYECPGLIEVPTEQDPSKSYWVMFISINPG	250
		DDLKSWKLESAFANEGFLGYQYECPGLIEVPTEQDPSKSYWVMFISINPG	
		DDLKSWKLESAFANEGFLGYQYECPGLIEVPTEQDPSKSYWVMFISINPG	
		DDLKSWKLESAFANEGFLGYQYECPGLIEVPTEQDPSKSYWVMFISINPG	
	251	APAGGSFNQYFVGSFNGTHFEAFDNQSRVVDFGKDYYALQTFFNTDPTYG	300
		APAGGSFNQYFVGSFNGTHFEAFDNQSRVVDFGKDYYALQTFFNTDPTYG	
		APAGGSFNQYFVGSFNGTHFEAFDNQSRVVDFGKDYYALQTFFNTDPTYG	
		APAGGSFNQYFVGSFNGTHFEAFDNQSRVVDFGKDYYALQTFFNTDPTYG	
	301	SALGIAWASNWEYSAFVPTNPWRSSMSLVRKFSLNTEYQANPETELINLK	350
		SALGIAWASNWEYSAFVPTNPWRSSMSLVRKFSLNTEYQANPETELINLK	
		SALGIAWASNWEYSAFVPTNPWRSSMSLVRKFSLNTEYQANPETELINLK	
		SALGIAWASNWEYSAFVPTNPWRSSMSLVRKFSLNTEYQANPETELINLK	
	351	AEPILNISNAGPWSRFATNTTLTKANSYNVDLSNSTGTLEFELVYAVNTT	400
		AEPILNISNAGPWSRFATNTTLTKANSYNVDLSNSTGTLEFELVYAVNTT	
		AEPILNISNAGPWSRFATNTTLTKANSYNVDLSNSTGTLEFELVYAVNTT	
		AEPILNISNAGPWSRFATNTTLTKANSYNVDLSNSTGTLEFELVYAVNTT	
	401	QTISKSVFPDLSLWFKGLEDPEEYLRMGFEVSASSFFLDRGNSKVKFVKE	450
		QTISKSVFADLSLWFKGLEDPEEYLRMGFEVSASSFFLDRGNSKVKFVKE	
		QTISKSVFADLSLWFKGLEDPEEYLRMGFEVSASSFFLDRGNSKVKFVKE	
		QTISKSVFADLSLWFKGLEDPEEYLRMGFEVSASSFFLDRGNSKVKFVKE	
	451	NPYFTNRMSVNNQPFKSENDLSYYKVYGLLDQNILELYFNDGDVVSTNTY	500
		NPYFTNRMSVNNQPFKSENDLSYYKVYGLLDQNILELYFNDGDVVSTNTY	
		NPYFTNRMSVNNQPFKSENDLSYYKVYGLLDQNILELYFNDGDVVSTNTY	
		NPYFTNRMSVNNQPFKSENDLSYYKVYGLLDQNILELYFNDGDVVSTNTY	
	501	FMTTGNALGSVNMTTGVDNLFYIDKFQVREVK	532
		FMTTGNALGSVNMTTGVDNLFYIDKFQVREVK	
		FMTTGNALGSVNMTTGVDNLFYIDKFQVREVK	
		FMTTGNALGSVNMTTGVDNLFYIDKFQVREVK	

Figure S6. Alignment of the translated *SUC2* sequence of *S. cerevisiae* CEN.PK113-7D, JP1, UFMG-CM-Y259, and S288C (reference). Missense mutations in at least one strain are highlighted in yellow. The alignment was performed using the Basic Local Alignment Search Tool with a translated nucleotide query (BLASTX) (Altschul et al. 1997).

Table S5. Point mutations within 1000 bp upstream of
the *SUC2* ORF of strains JP1 and UFMG-CM-Y259
with respect to the reference strain S288C. Colored in
grey are mutations shared by the two strains.

Position (bp)	nucleotide mutation		
JP1			
-929	C	→	A
-805	A	→	C
-776	C	→	T
-704	C	→	T
-698	A	→	G
-184	C	→	CTTTT, CTTTTTTTTT
-86	CT	→	C
UFMG-CM-Y259			
-971	GT	→	G
-805	A	→	C
-776	C	→	T
-737	G	→	A
-317	T	→	C
-236	A	→	T
-184	C	→	CTTTT, CTTTTTTTTT
-170	A	→	T

Table S6. Point mutations in the *AGT1* sequence of *S. cerevisiae* strains JP1 and UFMG-CM-Y259 with respect to the reference strain CAT-1 (S288C does not have a functional *AGT1* allele). Nucleotide and aminoacid (aa) changes are indicated in red and blue, respectively. Colored in grey are mutations shared by the two strains.

Gene: *AGT1*			Accession number: MF374788		
		Mutation			
Position (bp)	**Original codon**	**aa**	**Mutated codon**	**aa**	**Type**
JP1					
118	ATT	N	GAT	D	Missense
233	ACG	T	ATG	M	Missense
305	ATA	I	AAA	K	Missense
383	AAC	N	AGC	S	Missense
489	GTC	V	GTT	V	Silent
491 492	CAA	Q	CTT	L	Missense
523	CCT	P	ACT	T	Missense
634	GTG	V	ATG (80%)	M	Missense
675	CAG	Q	CAA	Q	Silent
676	GGT	G	AGT (68%)	S	Missense
			TGT (32%)	C	Missense
682 684	ACT	T	GCT (69%)*	A	Missense
			ACC (27%)*	T	Silent
688	ACT	T	TCT (25%)	S	Missense
819	TCT	S	TCC	S	Silent
997	ATT	I	GTT	V	Missense
1024	TTG	L	ATG	M	Missense
1075	GAT	D	AAT	N	Missense
1123	GCT	A	ACT	T	Missense
1142	ACT	T	AGT	S	Missense
1153 1154	TGT	C	GTT	V	Missense
1225	GTA	V	CTA	L	Missense
1336	TTA	L	CTA	L	Silent
1342	GTT	V	ATT	I	Missense
1375	GGC	G	AGC	S	Missense
1461	GTA	V	GTT	V	Silent
1462	ACT	T	GCT	A	Missense
1525	ATC	I	CTC	L	Missense
1539	ATC	I	ATT	I	Silent
1667	AGT	S	ACT	T	Missense
UFMG-CM-Y259					

Gene: *AGT1*		Accession number: MF374788			
Mutation					
Position (bp)	**Original codon**	**aa**	**Mutated codon**	**aa**	**Type**
118	ATT	N	GAT	D	Missense
233	ACG	T	ATG	M	Missense
288	TTA	L	TTG	L	Silent
305	ATA	I	AAA	K	Missense
383	AAC	N	AGC	S	Missense
491 492	CAA	Q	CTT	L	Missense
523	CCT	P	ACT	T	Missense
675	CAG	Q	CAA	Q	Silent
676	GGT	G	AGT (70%)	S	Missense
			TGT (30%)	C	Missense
682	ACT	T	GCT (71%)	A	Missense
684			ACC (29%)	T	Silent
688	ACT	T	TCT (26%)	S	Missense
819	TCT	S	TCC	S	Silent
997	ATT	I	GTT	V	Missense
1075	GAT	D	AAT	N	Missense
1123	GCT	A	ACT	T	Missense
1142	ACT	T	AGT	S	Missense
1153	TGT	C	GTT	V	Missense
1225	GTA	V	CTA	L	Missense
1342 1344	GTT	V	ATC	I	Missense
1375	GGC	G	AGC	S	Missense
1462	ACT	T	GCT	A	Missense
1525	ATC	I	CTC	L	Missense
1667	AGT	S	ACT	T	Missense

Figure S7. Alignment of the translated *AGT1* sequence of *S. cerevisiae* JP1 and UFMG-CM-Y259, and CAT-1 (reference). Missense mutations in at least one strain are highlighted in yellow. The alignment was performed using the Basic Local Alignment Search Tool with a translated nucleotide query (BLASTX) (Altschul et al. 1997).

Table S7. Point mutations in the *HXT3* sequence of *S. cerevisiae* strains JP1 and UFMG-CM-Y259 with respect to the reference strain S288C. Nucleotide and aminoacid (aa) changes are indicated in red and blue, respectively. Colored in grey are mutations shared by the two strains.

Gene: *HXT3*	Accession number:				
Mutation					
Position (bp)	**Original codon**	**aa**	**Mutate codon**	**aa**	**Type**
JP1					
501	TAT	Y	TAC	Y	Silent
906	GCT	A	GCC	A	Silent
913 915	TCA	S	ACT	T	Missense
921	TCA	S	TCT	S	Silent
927	GGT	G	GGC	G	Silent
930	GAG	E	GAA	E	Silent
933	TTG	L	TTA	L	Silent
945	AAG	K	AAA	K	Silent
948	CCG	P	CCA	P	Silent
958 960	AAG	K	CAA	Q	Missense
1443	CCA	P	CCG	P	Silent
UFMG-CM-Y259					
36	AAG	K	AAA	K	Silent
148	ACC	T	GCC	A	Missense
159	AAT	N	AAC	N	Silent
174	GCA	A	GCC	A	Silent
393	GCT	A	GCC	A	Silent
414	GGT	G	GGA	G	Silent
480	TCC	S	TCT	S	Silent
501	TAT	Y	TAC	Y	Silent
867	GAG	E	GAA	E	Silent
948	CCG	P	CCA	P	Silent
1159	TTG	L	CTG	L	Silent
1174	ATT	I	GTT	V	Missense
1225	CTA	L	TTA	L	Silent
1282	GTC → TC				Deletion
1284	GTC	V	TGT	C	Missense
1288	GC → CGC				Insertion
1291	GCC	A	GCA	A	Silent
1452	ACT	T	ACC	T	Silent

Strain		*HXT3* sequence		
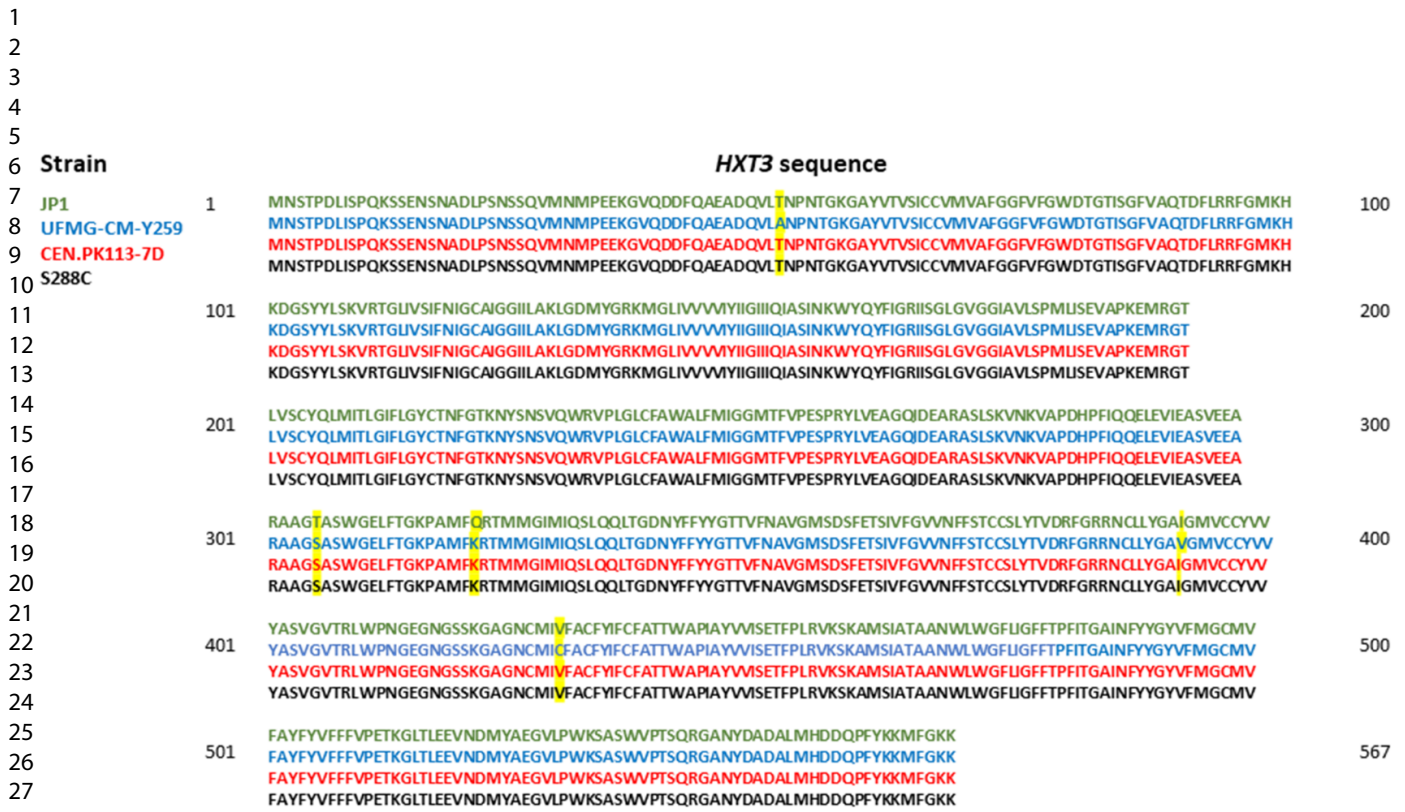
Figure S8. Alignment of the translated *HXT3* sequence of *S. cerevisiae* CEN.PK113-7D, JP1, UFMG-CM-Y259, and S288C (reference). Missense mutations in at least one strain are highlighted in yellow. The alignment was performed using the Basic Local Alignment Search Tool with a translated nucleotide query (BLASTX) (Altschul et al. 1997).

Table S9. Point mutations within 1000 bp upstream of the *HXT3* ORF of *S. cerevisiae* strains JP1 and UFMG-CM-Y259 with respect to the reference strain S288C. Only one mutation is shared by the two strains (in gray color).

Position (bp)	nucleotide mutation		
JP1			
-118	G	→	A
-601	A	→	T
UFMG-CM-Y259			
-65	T	→	C
-124	G	→	A
-125	CT	→	C
-140	GA	→	G

Position (bp)	nucleotide mutation		
-519	T	→	C
-586	C	→	G
-601	A	→	T
-689	T	→	TAA
-820	A	→	AGC
-858	C	→	T
-904	C	→	T
-925	T	→	C
-926	C	→	G

References

Altschul SF, Madden TL, Schäffer AA, et al (1997) Gapped BLAST and PSI-BLAST: a new generation of protein database search programs. Nucleic Acids Res 25:3389–3402. https://doi.org/10.1093/nar/25.17.3389

Reviewers' Comments to Authors:

Reviewer: 1

Comments to the Author
The authors tried to address the comments raised by the reviewers, however some key issues hasn't been well elucidated.
For example, the authors proposed that the sugar transporter would affect the cell growth on sucrose, glucose and fructose, however there was lack of direct evidence. Suggest the authors conduct transcriptional analysis of different sugar transporters with various affinities.

We agree with the reviewer that such a transcriptional analysis would improve the quality of the work and that we would be able to ascertain how and which sugar transporters influence the physiology of the different *S. cerevisiae* strains during growth on different sugars. However, we would have to rerun all 24 bioreactor cultivations (=4 sugars X 3 strains X 2 replicates), which at this point is unfeasible. Thus, we cannot perform these analyses, unfortunately, and I will leave it to the editor to decide whether they are crucial or not for acceptance of our manuscript.

Reviewer: 2

Comments to the Author
The author improved the manuscript according to the comments of the reviewers. The methods and experiment design are now detailed enough for readers to repeat. Through sequencing the genomes of JP1 and Y259 strains, they also explained some conclusions like mutation of SUC2. Although it still needs further study for quantitative physiology of yeast growth on sucrose, just for this manuscript it can be published.

We would like to thank the reviewer for her/his comments. We understand that no action is required from this assessment.

Reviewer: 3

Comments to the Author
In the revised manuscript, the authors have adequately addressed my comments. As a results, the quality of the manuscript has been significantly improved.

We would like to thank the reviewer for her/his comments.

I recommend acceptance of the revise manuscript after they address my minor comments as follows.

1. Page 13, line 15-18, please remove semicolons between each questions.
Semicolons removed.

2. Page 17 line 3-4, and page 19 line 10-11, please change "x" to the symbol for multiple "×", e.g., "2×".
Changed.

3. Page 34, line 11-13, please keep the consistent style of the reference.
Reference style corrected for consistence.

4. Some references are not shown intact, e.g. "De Deken RH (1966) The Crabtree Effect: A Regulatory System in Yeast" (Page 33, line 1), "E4tech, Re-Cord, WUR (2015) From the Sugar Platform to biofuels and biochemicals" (Page 33, line 14).
Both references were completed.

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5. Page 40-42, please remove the underlines in the figure legends and table description. Since the tables are very similar to each other, same situation with the Figures, we thought that underlining/highlighting the differences among them would facilitate interpretation by the reader. We will leave it like this for the moment, but if the editor considers that the underlining should indeed be removed, we would not oppose to this. We believe this is something simple that could be fixed during the editing steps, in case our manuscript is accepted for publication.