Laboratory evolution of a *Saccharomyces cerevisiae* x *S. eubayanus* hybrid under simulated lager-brewing conditions: genetic diversity and phenotypic convergence

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Abstract

Saccharomyces pastorianus lager-brewing yeasts are domesticated hybrids of *S. cerevisiae* x *S. eubayanus* that display extensive inter-strain chromosome copy number variation and chromosomal recombinations. It is unclear to what extent such genome rearrangements are intrinsic to the domestication of hybrid brewing yeasts and whether they contribute to their industrial performance. Here, an allodiploid laboratory hybrid of *S. cerevisiae* and *S. eubayanus* was evolved for up to 418 generations on wort under simulated lager-brewing conditions in six independent sequential batch bioreactors. Characterization of 55 single-cell isolates from the evolved cultures showed large phenotypic diversity and whole-genome sequencing revealed a large array of mutations. Frequent loss of heterozygosity involved diverse, strain-specific chromosomal translocations, which differed from those observed in domesticated, aneuploid *S. pastorianus* brewing strains. In contrast to the extensive aneuploidy of domesticated *S. pastorianus* strains, the evolved isolates only showed limited (segmental) aneuploidy. Specific mutations could be linked to calcium-dependent flocculation, loss of maltotriose utilisation and loss of mitochondrial activity, three industrially relevant traits that also occur in domesticated *S. pastorianus* strains. This study indicates that fast acquisition of extensive aneuploidy is not required for genetic adaptation of *S. cerevisiae* x *S. eubayanus* hybrids to brewing environments. In addition, this work demonstrates that, consistent with the diversity of brewing strains for maltotriose utilization, domestication under brewing conditions can result in loss of this industrially relevant trait. These observations have important implications for the design of strategies to improve industrial performance of novel laboratory-made hybrids.

Introduction

Saccharomyces yeasts are popular eukaryotic models for studying genome hybridisation, chromosome (mis)segregation and aneuploidy (Botstein, et al. 1997; Sheltzer, et al. 2011). The genus *Saccharomyces* arose between 10 and 20 million years ago and currently comprises eight described species, as well as interspecies hybrids (Liti, et al. 2006; Hittinger 2013; Naseeb, et al. 2017). Absence of a prezygotic barrier between *Saccharomyces* species facilitates hybridization, although spore viabilities of the resulting hybrids is typically well below 10 % (Liti, et al. 2006; Hittinger 2013; Naseeb, et al. 2017). Several interspecies *Saccharomyces* hybrids are tightly associated with domestication in industrial processes. *S. pastorianus* lager-brewing yeasts are domesticated *S. cerevisiae* x *S. eubayanus* hybrids (Libkind, et al. 2011). Double and triple hybrids between *S. cerevisiae*, *S. kudriavzevii* and *S. uvarum* are closely associated with wine fermentation (González, et al. 2006; Querol and Bond 2009; Marsit and Dequin 2015). *S. bayanus* cider fermentation yeasts are domesticated *S. uvarum* X *S. eubayanus* hybrids (Naumov, et al. 2001). Reconstruction of the corresponding *Saccharomyces* hybrids in the laboratory showed improved performance, relative to the parental species. For example, laboratory-made *S. cerevisiae* x *S. eubayanus* hybrids combined sugar utilisation characteristics of *S. cerevisiae* and the superior performance at low temperatures of *S. eubayanus* (Hebly, et al. 2015; Krogerus, et al. 2017). Similarly, hybrids of *S. cerevisiae*, *S. kudriavzevii* and *S. uvarum* combined traits of their parental species relevant to industrial wine fermentation, such as flocculence, sugar utilisation kinetics, stress tolerance and aroma production (Coloretti, et al. 2006; Lopandic, et al. 2016).

The relevance of laboratory hybridization of *Saccharomyces* species extends beyond reconstruction of existing, domesticated hybrids. The ability of hybridisation to generate extensive phenotypic diversity has raised interest in the development of novel *Saccharomyces* hybrids for specific industrial processes (Krogerus, et al. 2017). For example, an *S. cerevisiae* x *S. paradoxus* hybrid produced high concentrations of aromatic compounds that are of interest for wine making

(Bellon, et al. 2011). Hybrids between *S. cerevisiae* and *S. arboricola* or *S. mikatae* were able to utilize the sugars in wort at low temperatures and produced particularly aromatic beer (Nikulin, et al. 2017). Laboratory hybrids of *S. cerevisiae* and *S. kudriavzevii* or *S. mikatae* yielded xylose-consuming strains with high inhibitor tolerance for 2nd generation biofuel production (Peris, et al. 2017).

The allo euploid genomes of laboratory hybrids of *Saccharomyces* species strongly differ from the extremely aneuploidy genomes of the domesticated strains used in traditional industrial processes. For example, the genomes of *S. pastorianus* lager-brewing yeasts contain between 45 and 79 chromosomes (Van den Broek, et al. 2015; Okuno, et al. 2016), a degree of aneuploidy that is not observed elsewhere in the *Saccharomyces* genus (Gorter de Vries, Pronk, et al. 2017). The current consensus is that all *S. pastorianus* strains have evolved from a single, common ancestor *S. cerevisiae* x *S. eubayanus* hybrid by centuries of domestication and strain improvement in brewing-related environments (Okuno, et al. 2016). However, it remains unclear when and how domestication resulted in the extensive chromosome copy number variations and phenotypic diversity of current *S. pastorianus* strains.

Hybrid genomes have a well-documented increased tendency to become aneuploid due to an increased rate of chromosome missegregation during mitosis and/or meiosis (Chambers, et al. 1996; Liti, et al. 2006). Aneuploidy reduces the efficiency of sporulation and can thereby complicate genetic modification, impeding breeding and targeted strain improvement (Santaguida and Amon 2015; Gorter de Vries, de Groot, et al. 2017). In evolutionary contexts, aneuploidy is generally seen as a transient adaptation mechanism, whose positive impacts are eventually taken over by more parsimonious mutations (Yona, et al. 2012). When grown mitotically, sporulated hybrid strains were prone to further chromosome missegregation resulting in more extensive chromosome copy number variations (Lopandic, et al. 2016). Even genomes of *Saccharomyces* hybrids that had not undergone meiosis displayed increased rates of mitotic chromosome missegregation during mitosis (Delneri, et al. 2003). Indeed, when evolved in lignocellulosic hydrolysates, cultures of *S. cerevisiae* X *S.*

kudriavzevii and *S. cerevisiae* x *S. mikatae* hybrids exhibited segmental and full-chromosome aneuploidy after only 50 generations (Peris, et al. 2017). Similarly, when evolved under wine fermentation conditions, *S. cerevisiae* x *S. kudriavzevii* hybrids displayed extensive genome reorganisations that led to a significant reduction of their genome content (Pérez Través, et al. 2014).

Genetic instability of hybrid genomes could be detrimental to stable, robust industrial performance. Therefore, to assess industrial applicability of new hybrids generated in the laboratory, it is important to determine their genome stability under industrially relevant conditions. Moreover, laboratory evolution under simulated industrial conditions can increase understanding of the selective pressures that shaped the genomes of domesticated microorganisms (Bachmann, et al. 2012; Gibbons, et al. 2012; Gibbons and Rinker 2015).

The goal of the present study was to investigate how a previously constructed allodiploid *S. cerevisiae* x *S. eubayanus* hybrid (Hebly, et al. 2015) evolves under simulated lager-brewing conditions, with a specific focus on genome dynamics and on acquisition or loss of brewing-related phenotypes. To mimic successive lager beer fermentation processes, the hybrid strain was subjected to sequential batch cultivation on industrial wort, in six independent bioreactor setups. After up to 418 generations, the genotypic and phenotype diversity generated in these laboratory evolution experiments was analysed by characterization of 55 single-cell isolates. After whole-genome resequencing of each isolate using 150 bp paired-end reads, sequence data were mapped to high-quality reference genomes of the parental strains to identify genomic changes. Phenotypic analysis of the isolates focused on the ability to utilise maltotriose, flocculation and the respiratory capacity. We interpreted these results in the context of the domestication history of *S. pastorianus* brewing strains as well as in relation to genome stability and industrial application of newly generated *Saccharomyces* hybrids.

Results

Simulating domestication under lager-brewing conditions in sequential batch bioreactors

Industrial lager brewing involves batch cultivation of *S. pastorianus* on wort, an extract from malted barley, at temperatures between 7 and 15 °C. After a brief initial aeration phase to enable oxygen-dependent biosynthesis of unsaturated fatty acids and sterols (Andreasen and Stier 1953; Gibson, et al. 2007), brewing fermentations are not aerated or stirred, leading to anaerobic conditions during the main fermentation (Briggs, et al. 2004). To simulate domestication under industrial lager-brewing conditions, a laboratory evolution regime was designed in which the laboratory-made *S. cerevisiae* x *S. eubayanus* hybrid IMS0408 was grown at 12 °C in sequential batch bioreactors on industrial wort. As in industrial brewing, each cultivation cycle was preceded by an aeration phase, after which cultures were incubated without sparging or stirring until a decline of the CO₂ production indicated a cessation of sugar consumption. The bioreactors were then partially emptied, leaving 7 % of the culture volume as inoculum for the next aeration and fermentation cycle, which was initiated by refilling the reactor with sterile wort (Figure 1A and 1B). To mimic the low sugar concentrations during early domestication of *S. pastorianus* (Meussdoerffer 2009), parallel duplicate experiments at 12 °C were not only performed with full-strength 17 °Plato wort ('High Gavity', experiments HG12.1 and HG12.2), but also with three-fold diluted wort ('Low Gavity', experiments LG12.1 and LG12.2). To enable a larger number of generations during 4 months of operation, additional duplicate experiments on three-fold diluted wort were performed at 30 °C (LG30.1 and LG30.2).

Figure 1: Laboratory evolution mimicking the domestication of lager-brewing yeast. The *S. cerevisiae* x *S. eubayanus* laboratory hybrid IMS0408 was grown in duplicate sequential batch bioreactors in 3-fold diluted wort at 30 °C (LG30.1 and LG30.2) and at 12 °C (LG12.1 and LG12.2), and in full-strength wort at 12 °C (HG12.1 and HG12.2). (A) Experimental design for simulated sequential lager-beer brewing cycles. Each cycle consisted of four phases: (i) (re)filling of the fermenter with fresh medium up to a total volume of 100 mL, (ii) aeration at 200 mL/min while stirring at 500 RPM, (iii) a batch fermentation phase without sparging or stirring, while flushing the bioreactor headspace with N₂ to enable accurate analysis of CO₂ production and (iv) removal of broth, leaving 7 mL to inoculate the next cycle. (B) Fermentation profiles of three consecutive cycles from experiment LG30.1, performed at 30 °C in 3-fold diluted wort. Percentage of CO₂ in the off gas, culture pH and dissolved oxygen (dO₂) concentration are indicated by red, blue and black symbols, respectively. Due to the lack of stirring and sparging, CO₂ was slowly released by the medium; emptying of the reactor was initiated when the offgas CO₂ concentration dropped to 70 % of its initial value as off-line analyses indicated that, at this point, all fermentable sugars had been consumed (C) Specific growth rates estimated from CO₂ production profiles during each cycle of the evolution lines. LG30.1 (blue circles) and LG30.2 (red circles) were grown on 3-fold diluted wort at 30 °C; LG12.1 (blue triangles) and LG12.2 (red triangles) were grown on 3-fold diluted wort at 12 °C. HG12.1 (blue squares) and HG12.2 (red squares) were evolved in full-strength wort at 12 °C. Since lack of sparging and stirring precluded exact estimates of specific growth rates, the calculated values should be taken as indicative.

Concentration of the wort and temperature strongly affected the length of the fermentation cycles, which was 17 h for LG30.1 and LG30.2, 93 h for LG12.1 and LG12.2 and 205 h for HG12.1 and HG12.2. Experiments LG30.1 and LG30.2 involved 117 and 118 batch cycles, respectively, LG12.1 and LG12.2 covered 58 and 57 cycles, respectively, and HG12.1 and HG12.2 covered 13 and 16 cycles, respectively. At the inoculum size of 7 % of the total culture volume, each cycle corresponded to approximately 4 generations. Specific growth rates, estimated from CO₂ production rates during the exponential growth phase of the batch cycles, were not significantly different during the first and the last five cycles of each experiment (Student's t-test, $p > 0.05$). Average specific growth rates were $0.35 \pm 0.02 \text{ h}^{-1}$ for LG30.1, $0.42 \pm 0.03 \text{ h}^{-1}$ for LG30.2, $0.070 \pm 0.013 \text{ h}^{-1}$ for LG12.1, $0.062 \pm 0.009 \text{ h}^{-1}$ for LG12.2, $0.068 \pm 0.007 \text{ h}^{-1}$ for HG12.1 and $0.098 \pm 0.018 \text{ h}^{-1}$ for HG12.2. While the initial specific growth rate was clearly higher at 30 °C than at 12 °C, initial growth rates on diluted and full-strength wort were not significantly different. However, CO₂ production from sugars continued much longer in full-strength wort. During brewing fermentation, depletion of nitrogen sources and oxygen limit biomass formation. Complete sugar conversion therefore depends on growth-independent alcoholic fermentation which, apparently, was much slower in cultures grown on full-strength wort. At the end of each evolution experiment, culture samples were streaked on YPD agar and 5 single colonies were isolated for each culture. For experiments LG12.1 and LG12.2, isolates were also made from intermediate samples after 29 cycles. Evolution lines LG30.2 and LG12.2 developed flocculence and isolates from these lines had two distinct colony morphologies: about half of the colonies were elevated and conically-shaped, while the other colonies shared the flat morphology of IMS0408. For each of these lines, five random colonies of each morphology were selected.

Prolonged growth under simulated brewing conditions did not cause large ploidy changes

Six independent sequential batch fermentation experiments under simulated brewing conditions, covering 52 to 468 generations, yielded 55 isolates. Staining with the DNA-binding fluorescent dye SYTOX Green and flow cytometry indicated genome sizes of the isolates between 17.6 and 23.5 Mbp (Supplementary table S1). These values did not differ significantly from the 21.3 ± 1.9 Mbp genome size measured for the parental laboratory hybrid IMS0408 and therefore indicated the absence of large changes in genome content such as whole-genome duplications. For a detailed genotypic analysis, the genomes of the 55 isolates were sequenced using 150 bp pair-end reads with 101- to 189-fold coverage. A high quality-reference genome was constructed by combining the chromosome-level contigs from assemblies of CEN.PK113-7D and CBS12357 generated with nanopore technology, including mitochondrial genome sequences (Salazar, et al. 2017; Brickwedde, et al. 2018).

Copy number analysis revealed whole-chromosome aneuploidies in only 5/55 isolates (Table 1). Relative to strain IMS0408, the total chromosome number of the isolates had not changed by more than one. Isolate IMS0556 (LG30.1) had gained a copy of *ScCHRVIII*, IMS0560 (LG30.1) had gained a copy of *SeCHRX*, IMS0565 (LG30.2) had lost *ScCHRXIV* and gained a copy of *SeCHRXIV*, IMS0595 (LG12.1) had gained a copy of *SeCHRVIII* and IMS0606 (LG12.2) had lost a copy of *SeCHRVIII*. Read alignments to mitochondrial genome sequences were absent from 14/55 isolates, while 1 isolate showed only a partial alignment, indicating complete (ρ^-) or partial loss (ρ^0) of the mitochondrial genome in 15/55 strains. Loss of respiratory competence was confirmed by the observation that these 15 isolates, in contrast to IMS0408 and isolates containing a full mitochondrial genome, were unable to grow on YP-ethanol (Supplementary Figure S1).

1 **Table 1: Overview of phenotypic and genotypic changes in isolates obtained after laboratory evolution of the allodiploid laboratory hybrid IMS0408**
2 **under simulated lager-brewing conditions.**

Experiment	GEN# ¹	Isolate	M ²	F ³	R ⁴	M _{DNA} ⁵	Aneuploidy ^{6,7}	Segmental aneuploidy and loss of heterozygosity ^{6,7}	SNPs ^{6,8}	INDELs ^{6,8,9}
LG12.1 (3x diluted wort, 12 °C)	Unevolved	IMS0408	+	-	+					
		IMS0538	+	-	+					
	116	IMS0539	+	-	+					
		IMS0540	+	-	+				<i>SeHDA2</i> ^{A1651T}	<i>SeMED2</i> ^{462+3N}
		IMS0541	+	-	-	p ⁰				
		IMS0542	+	-	+				<i>ScIQG1</i> ^{A2069C}	<i>SeKEX1</i> ^{1875-6N}
		IMS0543	+	+	+			Δ Sc(YKL032C-YKL054C)		
		IMS0544	+	+	+			Δ Sc(YKL032C-YKL054C), Δ Sc::Se(YLR154C-YLR _{end})	<i>SeSAC1</i> ^{G1093C}	
		IMS0545	+	+	+			Δ Sc(YKL032C-YKL054C)		<i>SeNAF1</i> ^{1404-30N,1436-30N}
		IMS0546	+	+	-	p ⁻		Δ Sc(YKL032C-YKL054C)		
		IMS0547	+	+	+			Δ Sc(YKL032C-YKL054C)		
	232	IMS0594	+	-	+					<i>SeELA1</i> ^{230+343N} , <i>SeIRA2</i> ^{1402-1N} , <i>ScFLO9</i> *
		IMS0595	+	-	+		2xSe(CHRVIII)	Δ Se::Sc(YOL _{end} -YOL072W)		
		IMS0596	+	-	+				<i>SeIRA2</i> ^{C1376A} , <i>ScYER188W</i> ^{T28A}	
		IMS0597	+	-	+			Δ Se::Sc(YOL _{end} -YOL057W)	<i>SeNUP1</i> ^{C1205T} , <i>ScPDC2</i> ^{G372A}	
		IMS0598	+	-	+				<i>SeSRT1</i> ^{C359T}	<i>SeASG1</i> ^{2488+3N}
		IMS0599	+	+	+			Δ Sc(YKL032C-YKL054C), Δ Se::Sc(YLR154C-YLR _{end})		
		IMS0600	+	-	+			Δ Se::Sc(YOL _{end} -YOL075W)	<i>SeMSR1</i> ^{A853C}	
		IMS0601	+	+	-	p ⁻		Δ Sc(YKL032C-YKL054C)		
		IMS0602	+	+	+			Δ Sc(YKL032C-YKL054C)		
		IMS0603	+	-	+			Δ Se::Sc(YNL _{end} -YNL123W), Δ Se::Sc(YOL _{end} -YOL013C), Δ Sc::Se(YOL013C-YOL006C)		

3 Footnotes: A complete list of mutations is provided in Supplementary Data File 1. ¹ Estimated number of generations prior to isolation. ² Ability to utilise the
4 sugar maltotriose. ³ Flocculation during growth on liquid medium. ⁴ Respiratory competence. ⁵ Presence of the mitochondrial DNA. An empty field indicates
5 presence of the *S. eubayanus* mitochondrial DNA, p⁻ indicates complete loss of the mitochondrial genome and p⁰ partial loss. ⁶ Sc and Se indicate sequences
6 on the *S. cerevisiae* and *S. eubayanus* subgenomes, respectively. ⁷ “ Δ ” indicates deletion, “2x” indicates duplication and “::” indicates substitution. ⁸ SNPs
7 and INDELs are only indicated when they affected an ORF and were non synonymous. ⁹ The number of nucleotides added (+N) or removed (-N) is indicated
8 after the coordinate of the last unchanged nucleotide.

Experiment	GEN# ¹	Isolate	M ²	F ³	R ⁴	M _{DNA} ⁵	Aneuploidy ^{6,7}	Segmental aneuploidy and loss of heterozygosity ^{6,7}	SNPs ^{6,8}	INDELS ^{6,8,9}
LG12.2 (3x diluted wort, 12 °C)	116	IMS0548	+	-	+			$\Delta Se::Sc(YKL_{end}-YKL057C), \Delta Sc::Se(YKL057C-YKR_{end})$		
		IMS0549	+	-	-	ρ^-				
		IMS0550	+	-	-	ρ^-				
		IMS0551	+	-	-	ρ^-		$\Delta Sc(YCL_{end}-YCL067C), \Delta Sc(YCR039C-YCR_{end})$		
		IMS0552	+	-	-	ρ^-		$\Delta Sc::Se(YHL_{end}-YHL023C)$		
	228	IMS0604	+	-	+			$\Delta Sc(YKL032C-YKL054C), \Delta Sc::Se(YLR305C-YLR_{end})$	<i>SeBET2</i> ^{G550A}	
		IMS0605	+	-	+			$\Delta Sc(YKL032C-YKL054C)$		
		IMS0606	+	-	-	ρ^-	$\Delta Se(CHRVIII)$	$\Delta Sc(YKL032C-YKL054C)$	<i>ScLRG1</i> ^{C2277G}	
		IMS0607	+	-	+			$\Delta Sc(YKL032C-YKL054C)$		<i>SeFLO11</i> *
		IMS0608	+	-	-	ρ^-		$\Delta Sc(YKL032C-YKL054C)$		
LG30.1 (3x diluted wort, 30 °C)	464	IMS0553	+	-	+					<i>SeNHX1</i> ^{1622+3N}
		IMS0554	-	-	+			$\Delta Sc::Se(YGR282C-YGR_{end})$		<i>ScCIS3</i> *
		IMS0555	+	-	+				<i>SeBAT1</i> ^{G1073A} , <i>ScMAL23</i> ^{G422A}	
		IMS0556	+	-	+		2xSc(CHRVIII)		<i>SeGMC1</i> ^{G1579A} , <i>ScSFL1</i> ^{T605A}	
		IMS0557	-	-	+				<i>ScMAL11</i> ^{G88T, A98G} , <i>ScMDL2</i> ^{C1451A}	<i>ScMAL11</i> ^{93-1N} , <i>SeYNL247W</i> ^{2062-1N}
		IMS0558	-	+	+			$\Delta Sc(YDR261C-YDR211W), \Delta Sc::Se(YGR218C-YGR_{end})$	<i>ScSFL1</i> ^{T605A}	<i>SeSFL1</i> ^{96+1N}
		IMS0559	+	+	+			$\Delta Se::Sc(YOR133W-YOR_{end})$	<i>ScSFL1</i> ^{T605A}	
		IMS0560	+	+	+		2xSe(CHRX)	$\Delta Sc(YDR261C-YDR211W), \Delta Se::Sc(YOR063W-YOR_{end})$	<i>ScSFL1</i> ^{T605A}	
		IMS0561	+	+	+			$\Delta Se::Sc(YBR275C-YBR_{end}), \Delta Se::Sc(YOR133W-YOR_{end})$	<i>ScSFL1</i> ^{T605A}	
		IMS0562	+	+	+			$\Delta Sc::Se(YNL061C-YNL055C), \Delta Se::Sc(YOR133W-YOR_{end})$	<i>ScSFL1</i> ^{T605A}	
LG30.2 (3x diluted wort, 30 °C)	468	IMS0563	-	-	+			$\Delta Sc(YGR279C-YGR_{end})::Se(YMR305C-YMR_{end})$	<i>SeROG3</i> ^{G191A} , <i>SeMSS51</i> ^{C448T}	<i>ScRTP1</i> ^{874-64N}
		IMS0564	+	-	+				<i>SeIZH3</i> ^{A526G} , <i>ScCST6</i> ^{C807A}	
		IMS0565	-	-	+		$\Delta Sc::Se(CHRXIV)$		<i>ScMAL11</i> ^{A1G}	
		IMS0566	+	-	-	ρ^-			<i>ScERG6</i> ^{C413T} , <i>ScBUL1</i> ^{G2110A}	<i>SeYBR238C</i> ^{315-36N}
		IMS0567	-	-	-	ρ^-		$\Delta Sc::Se(YDR051C-YDR_{end}), \Delta Sc::Se(YGR271W-YGR_{end})$		<i>ScULP2</i> ^{1469+1N}
HG12.1 (Full-strength wort, 12 °C)	52	IMS0609	+	-	+					
		IMS0610	+	-	-	ρ^-			<i>ScYBR259W</i> ^{C833A}	
		IMS0611	+	-	-	ρ^-			<i>ScFMP52</i> ^{G406C}	
		IMS0612	+	-	+					
		IMS0613	+	-	+				<i>SeGIC2</i> ^{C344G} , <i>SeSFL1</i> ^{C1390T}	<i>SeGCN2</i> ^{2274+18N, 2239+18N} , <i>SeTRA1</i> ^{4421-25N}
HG12.2 (Full-strength wort, 12 °C)	64	IMS0614	+	-	+			$\Delta Se(YAR050W-YAR_{end})::Se(YAL_{end}-YAL063C)$	<i>SeCAC2</i> ^{A994T}	
		IMS0615	+	-	+				<i>ScATG1</i> ^{A434C}	
		IMS0616	+	-	-	ρ^-			<i>SeSFL1</i> ^{C1390T}	
		IMS0617	+	-	+				<i>ScALR1</i> ^{G1645T}	
		IMS0618	+	-	-	ρ^-				

Footnotes: A complete list of mutations is provided in Supplementary Data File 1. ¹ Estimated number of generations prior to isolation. ² Ability to utilise the sugar maltotriose. ³ Flocculation during growth on liquid medium. ⁴ Respiratory competence. ⁵ Presence of the mitochondrial DNA. An empty field indicates presence of the *S. eubayanus* mitochondrial DNA, ρ^- indicates complete loss of the mitochondrial genome and ρ^0 partial loss. ⁶ Sc and Se indicate sequences on the *S. cerevisiae* and *S. eubayanus* subgenomes, respectively. ⁷ “ Δ ” indicates deletion, “2x” indicates duplication and “::” indicates substitution. ⁸ SNPs

16 and INDELs are only indicated when they affected an ORF and were non synonymous.⁹ The number of nucleotides added (+N) or removed (-N) is indicated
17 after the coordinate of the last unchanged nucleotide. *FLO9_{cer}** and *CIS3_{cer}** indicate coverage drops of these ORFs, for which the responsible mutation could
18 not be exactly reconstructed. *FLO11_{cer}** indicates an up to 10-fold increase in coverage in the middle of the open reading frame that indicates amplification
19 of internal additional repeats.

Chromosomal recombinations frequently caused loss of heterozygosity

Of the 55 evolved isolates, 29 displayed segmental copy number changes. In total, 20 of the 32 chromosomes in strain IMS0408 were affected in at least one isolate. Of the 55 evolved isolates, 24 showed chromosome segments with increased copy number and 41 showed chromosome segments with decreased copy number (Table 1, Figure 2). 17 internal recombinations resulting in deletions were observed: $\Delta Sc(YKL032C-YKL054C)$ occurred in 13 strains, $\Delta Sc(YDR261C-YDR211W)$ occurred in two strains, and $\Delta Sc(YCL_{end}-YCL067C)$ and $\Delta Sc(YCR039C-YCR_{end})$ occurred together in one strain. The internal recombinations $\Delta Sc(YKL032C-YKL054C)$ and $\Delta Sc(YDR261C-YDR211W)$ both resulted in loss of the sequence between the recombination sites. The recombination occurred between *IXR1* and *DEF1* in $\Delta Sc(YKL032C-YKL054C)$ and between Ty-transposons for $\Delta Sc(YDR261C-YDR211W)$. Finally, the concurrent loss of $Sc(YCL_{end}-YCL067C)$ and $Sc(YCR039C-YCR_{end})$ indicated loss of both ends of *ScCHRIII*, including telomeres, consistent with recombination into a circular chromosome III at the *HMLALPHA2* and *MATALPHA2* loci (Newlon, et al. 1991).

Figure 2: Total number of occurrences of whole-chromosome (A) and segmental (B) aneuploidy for each chromosome of IMS0408 among 55 isolates obtained after laboratory evolution under simulated lager fermentation conditions. For each chromosome, loss of genetic material is indicated in red and duplicated genetic material is indicated in blue. Loss or duplication of *S. cerevisiae* or *S. eubayanus* genetic material which was coupled with duplication or loss of the corresponding region of the other subgenome, is indicated by checked bars. *S. eubayanus* harbours two translocations relative to *S. cerevisiae*: between chromosomes II and IV, and between chromosomes VIII and XV. For simplicity, copy number affecting these regions were allocated based on the *S. cerevisiae* genome architecture.

The remaining 24 chromosome-segment duplications and losses reflected inter-chromosomal recombinations: one chromosomal region was replaced by an additional copy of another chromosomal region by a non-conservative recombination. The recombinations $\Delta Sc(YGR_{279C}-YGR_{end})::Se(YMR305C-YMR_{end})$ and $\Delta Se(YAR050W-YAR_{end})::Se(YAL_{end}-YAL063C)$ occurred between highly similar genes; the paralogs *SCW4* and *SCW10*, and *FLO1* and *FLO9*, respectively. In the remaining 22 cases, recombination occurred between homologous genes of each subgenome. No copy-number conservative chromosome translocations were identified.

Of the 26 observed recombinations, 23 occurred inside ORFs, and thus resulted in chimeric genes (Table 2). The homology between ORFs involved in recombinations varied from <70 % to 100 %, with a median homology of 82.41 %. Chimeric ORFs were reconstructed by extracting reads paired to the other chromosome from the sequencing data and using them for a local assembly. This approach allowed for identification of the recombination site at a resolution that, depending on sequence homology of the two ORFs, varied between 2 to 633 nucleotides. Due to length differences and relative INDELs between the original ORFs, recombined ORFs differed in length. However, all recombinations occurred in frame and no premature stop codons were introduced, suggesting that these chimeric ORFs might yield functional proteins.

59 **Table 2: Overview of all recombinations observed in 55 isolates obtained after laboratory evolution of strain IMS0408 under simulated lager**
60 **fermentation conditions.**

Recombination ^{1,2}	Affected isolates	Locus 1			Locus 2			Homology ⁵	Length chimeric ORF (bp)
		Name ¹	Length ³	Recombination ⁴	Name ¹	Length ³	Recombination ⁴		
ΔSe(YAR050W-YARend) ::Se(YALend-YAL063C)	IMS0614	SeFLO1	5517	711-734	SeFLO9	4752	712-735	79.23 %	4752
ΔSe::Sc(YBR275C-YBRend)	IMS0561	ScRIF1	5751	562-572	SeRIF1	5751	569-579	77.49 %	5745
ΔSc::Se(YDR051C-YDRend)	IMS0567	SeDET1	1005	426-434	ScDET1	1005	427-435	84.36 %	1005
ΔSc::Se(YGR218C-YGRend)	IMS0558	ScCRM1	3251	2626-2636	SeCRM1	3251	2627-2637	85.84 %	3251
ΔSc::Se(YGR271W-YGRend)	IMS0567	ScSLH1	5904	1335-1349	SeSLH1	5901	1336-1350	82.41 %	5901
ΔSc(YGR279C-YGRend) ::Se(YMR305C-YMRend)	IMS0563	SeSCW10	1146	892-899	ScSCW4	1161	908-915	<70 %*	1146
ΔSc::Se(YGR282C-YGRend)	IMS0554	SeBGL2	942	115-125	ScBGL2	942	116-126	87.75 %	942
ΔSc::Se(YHLend-YHL023C)	IMS0552	SeNPR3	3444	1551-1571	ScNPR3	3441	1561-1581	78.80 %	3432
ΔSe::Sc(YKLend-YKL057C), ΔSc::Se(YKL057C-YKrend)	IMS0548	SeNUP120	3114	1867-1868	ScNUP120	3114	1868-1869	80.14 %	3114
ΔSc::Se(YLR154C-YLRend)	IMS0544	Ribosomal DNA	-	-	Ribosomal DNA	-	-	-	-
ΔSe::Sc(YLR154C-YLRend)	IMS0599	Ribosomal DNA	-	-	Ribosomal DNA	-	-	-	-
ΔSc::Se(YLR305C-YLRend)	IMS0604	SeSTT4	5703	5013-5018	ScSTT4	5703	5014-5019	81.64 %	5703
ΔSc::Se(YNL061C-YNL055C)	IMS0562	ScNOP2	1857	1386-1391	SeNOP2	1860	1390-1395	87.11 %	1857
		ScPOR1	852	399-401	SePOR1	852	400-402	85.82 %	852
ΔSe::Sc(YNLend-YNL123W)	IMS0603	ScNMA111	2994	303-314	SeNMA111	2994	304-315	83.61 %	2994
ΔSc::Se(YOL013C-YOL006C)	IMS0603	SeTOP1	2304	2010-2021	ScTOP1	2311	2017-2028	82.64 %	2304
ΔSe::Sc(YOLend-YOL013C)	IMS0603	SeHRD1	1644	312-329	ScHRD1	1656	313-330	82.82 %	1656
ΔSe::Sc(YOLend-YOL057W)	IMS0597	ScYOL057W	3015	99-116	SeYOL057W	2133	100-117	<70 %*	2133
ΔSe::Sc(YOLend-YOL072W)	IMS0595	ScTHP1	1368	726-731	SeTHP1	1374	733-738	79.07 %	1368
ΔSe::Sc(YOLend-YOL075W)	IMS0600	SeYOL075C	3903	2409-2414	ScYOL075C	3885	2392-2397	81.85 %	3903
ΔSe::Sc(YOR063W-YORend)	IMS0560	ScRPL3	1164	1023-1063	SeRPL3	1164	1024-1064	94.59 %	1164
ΔSe::Sc(YOR133W-YORend)	IMS0559, IMS0561 and IMS0562	ScEFT1	2529	246-311	SeEFT1	2529	247-312	94.94 %	2529
ΔSc(YCLend-YCL067C), ΔSc(YCR039C-YCRend)	IMS0551	ScHMLALPHA2	633	1-633	ScMATALPHA2	633	1-633	100 %	633
ΔSc(YDR261C-YDR211W)	IMS0558 and IMS0560	TY-transposon	-	-	TY-transposon	-	-	-	-
ΔSc(YKL032C-YKL054C)	IMS0543-547, IMS0599, IMS0601 and IMS0602	ScIXR1	1794	944-955	ScDEF1	2217	1281-1292	<70 %*	1881
ΔSc(YKL032C-YKL054C)	IMS0604-608	ScIXR1	1794	332-341	ScDEF1	2217	1272-1281	<70 %*	1278

61 Footnotes: ¹ Sc and Se indicate sequences on the *S. cerevisiae* and *S. eubayanus* subgenomes, respectively. ² “Δ” indicates deletion and “::” indicates
62 substitution. ³ Length of the affected ORF in bp. ⁴ Nucleotide coordinates of the locus at which recombination could have occurred. Due to the high
63 homology of Locus 1 and Locus 2, they shared identical sequence stretches within which the recombination could have occurred. Recombination sites are
64 therefore indicated by a range of nucleotide coordinates within the ORFs. ⁵ Homology between Locus 1 and Locus 2 as determined by sequence alignment.

***IRA2*, *SFL1* and *MAL11* are mutated in multiple evolved isolates**

A total of 76 SNPs and 43 INDELs were identified in the genomes of the 55 isolates (Supplementary Data File 1A and 1B), of which 38 SNPs and 17 INDELs occurred in ORFs and were non-synonymous (Table 1). Gene ontology analysis of all genes affected by non-synonymous SNPs or INDELs did not yield a significant enrichment in specific biological processes, molecular functions or cellular components. However, the genes *IRA2*, *SFL1* and *MAL11* were affected in more than one strain. *IRA2* encodes a RAS GTPase-activating protein, which is disrupted in many *S. cerevisiae* genomes from the CEN.PK strain family (Tanaka, et al. 1990; Tanaka, et al. 1991; Nijkamp, et al. 2012). In strain IMS0408, the *ScIRA2* was indeed disrupted while the *SeIRA2* ORF was intact. However, *SeIRA2* was mutated in 6/10 isolates from LG12.1 after 232 generations. *SeIRA2* had a frameshift in IMS0594, a premature stop codon in IMS0596 and was completely lost in four isolates due to different loss of heterozygosities: $\Delta Se::Sc(YOL_{end}-YOL072W)$ in IMS0595, $\Delta Se::Sc(YOL_{end}-YOL057W)$ in IMS0597, $\Delta Se::Sc(YOL_{end}-YOL075W)$ in IMS0600 and $\Delta Se::Sc(YOL_{end}-YOL013C)$ in IMS0603.

SFL1 encodes a transcriptional repressor of flocculation genes, which was present both on *ScCHRXV* and *SeCHRVIII* (Atsushi, et al. 1989). *ScSFL1* was mutated in 6/10 isolates from LG30.1 after 464 generations, which harboured a non-conservative substitution at the 605th nucleotide, affecting its DNA binding domain (Atsushi, et al. 1989). *SeSFL1* had a frameshift in IMS0558 (LG30.1), a single nucleotide substitution in IMS0614 and IMS0617 (HG12.2) and was completely lost in four isolates of LG30.1 due to two losses of heterozygosity: $\Delta Se::Sc(YOR133W-YOR_{end})$ in IMS0559, IMS0561 and IMS0562, and $\Delta Se::Sc(YOR063W-YOR_{end})$ in IMS0560 (Table 1).

ScMAL11, also referred to as *AGT1*, encodes the only maltose transporter of the *MAL* gene family which enables efficient uptake of maltotriose in IMS0408 (Alves, et al. 2008). *ScMAL11* is located on the right arm of *ScCHRVII* and is absent in the *S. eubayanus* subgenome of IMS0408, which has no other maltotriose transporters (Hebly, et al. 2015; Brickwedde, et al. 2018). *ScMAL11* had a frameshift in IMS0557 (LG30.1) and lost its start codon in IMS0565 (LG30.2) (Table 1). In addition, *ScMAL11* was completely lost due to three different losses of heterozygosity: $\Delta Sc::Se(YGR282C-$

YGR_{end}) in IMS0554 (LG30.1), $\Delta Sc::Se(YGR218C-YGR_{end})$ in IMS0558 (LG30.1) and $\Delta Sc::Se(YGR271W-YGR_{end})$ in IMS0567 (LG30.2), and due to the non-conservative recombination $\Delta Sc(YGR279C-YGR_{end})::Se(YMR305C-YMR_{end})$ in IMS0563 (LG30.2).

Mutations in *SFL1* cause emergence of calcium-dependent flocculation

Eight of the 55 evolved isolates showed SNPs, INDELs or loss of heterozygosity in the flocculation inhibitor gene *SFL1*. In isolates IMS0558, IMS0559, IMS0560, IMS0561 and IMS0562 from LG30.1, both *SeSFL1* and *ScSFL1* were mutated, while in isolate IMS0556 from LG30.1 only *ScSFL1* was mutated and in isolates IMS0614 and IMS0617 from HG12.2, only *SeSFL1* was mutated. Evolved isolates that carried mutations in both *ScSFL1* and *SeSFL1* formed elevated conically-shaped colonies on YPD agar, while strain IMS0408 and evolved isolates with either an intact *SeSFL1* or *ScSFL1* did not (Figure 3A). Strains with mutations in both *ScSFL1* and *SeSFL1* also showed rapid sedimentation in micro-aerobic cultures on wort, which was not observed for the other evolved isolates or for strain IMS0408 (Figure 3F).

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Figure 3: Mutations in *ScSFL1* and *SeSFL1* correlate with flocculation in evolved isolates and reverse engineered strains. A) Colony morphology and phase-contrast microscopy images (100x) of YPD-grown cell suspensions of the non-evolved, non-flocculent strain (IMS0408) and of a typical flocculent evolved isolate (IMS0558). Resuspension in 50 mM EDTA (pH 7.0) eliminated flocculation. B) Biomass sedimentation of evolved isolates and engineered strains with mutations in *SeSFL1* and/or *ScSFL1*. Triplicate cultures of all strains were grown on YPD and sedimentation was measured as the decrease in OD₆₆₀ right underneath the meniscus of a stationary cell suspensions 60 s after the suspension had been vortexed.

SFL1 is a repressor of *FLO* effector genes, which mediate calcium-dependent flocculation in *S. pastorianus* strains. To test if the acquired mutations in *SFL1* affected calcium-dependent flocculation, cultures from each strain were inspected microscopically. In the absence of EDTA, all sedimenting strains formed large aggregates, while IMS0408 and strains with an intact *SeSFL1* or *ScSFL1* did not (Figure 3A). Upon resuspension in Ca^{2+} -chelating EDTA buffer, the aggregates reverted into a single-cell suspensions, thus confirming Ca^{2+} -dependent flocculation. To test if the mutations affecting *SeSFL1* and *ScSFL1* were responsible for the calcium-dependent flocculation, single and double knockout strains were made using CRISPR-Cas9 gene editing: IMX1824 ($\Delta\text{ScSFL1}::\text{mTurquoise2}$), IMX1825 ($\Delta\text{SeSFL1}::\text{Venus}$) and IMX1826 ($\Delta\text{ScSFL1}::\text{mTurquoise2}$, $\Delta\text{SeSFL1}::\text{Venus}$). Flocculation intensity was assessed by measuring the decrease in OD_{660} at the surface of stationary-phase cultures after 1 minute without shaking. Isolates with mutations affecting only *SeSFL1* or only *ScSFL1* did not sediment significantly faster than IMS0408, which showed a decrease in OD_{660} of 13.3 ± 5.8 % (Figure 3B). However, IMX1826 ($\text{mTurquoise2}::\Delta\text{ScSFL1}$, $\text{Venus}::\Delta\text{SeSFL1}$), IMS0558 (*ScSFL1*^{T605A}, *SeSFL1*^{96+1N}) and IMS0559 (*ScSFL1*^{T605A}, ΔSeSFL1) all showed decreases in OD_{660} above 53.6 %, indicating that disruption of both *ScSFL1* and *SeSFL1* causes a significant increase in flocculation (Student's T test, $p < 0.0011$). Since *SeSFL1* harboured three different mutations in isolates IMS0558, IMS0559, IMS0560, IMS0561 and IMS0562, calcium-dependent flocculation emerged by at least three independent mutations during the laboratory evolution experiment LG30.2.

Mutations in *ScMAL11* cause loss of maltotriose utilisation

ScMAL11, which encodes the sole maltotriose transporter in strain IMS0408, was mutated in six evolved isolates. To investigate if these mutations affected maltotriose fermentation, the isolates were grown micro-aerobically on diluted wort at 30 °C. The unevolved IMS0408 consumed 100 ± 0 % of the maltotriose in diluted wort, and evolved strains with an intact *ScMAL11* genes consumed $98 \pm$

3 %. In contrast, strains IMS0554, IMS0557, IMS0558, IMS0563, IMS0565 and IMS0567, which all harboured mutations in *ScMAL11*, did not show any maltotriose consumption; instead, the concentration increased by 14 ± 3 % on average, presumably due to water evaporation. To test if the mutations affecting *ScMAL11* were responsible for the loss of maltotriose utilisation, *ScMAL11* was deleted in strain IMS0408 using CRISPR-Cas9 gene editing, resulting in strain IMX1698 ($\Delta ScMAL11::mVenus$). Under the same conditions used to evaluate maltotriose utilization by the evolved strains, strain IMS0408 consumed 97 ± 5 % of the maltotriose while strain IMX1698 only consumed 1 ± 0 % of the maltotriose. These results confirmed that loss of *ScMAL11* function was responsible for loss of maltotriose utilization.

Discussion

Evolution of the laboratory *S. cerevisiae* x *S. eubayanus* IMS0408 under simulated lager-brewing conditions yielded a wide array of mutations, including SNPs, INDELs, chromosomal recombinations, aneuploidy and loss of mitochondrial DNA (Table 1, Additional file 2). SNPs were the most common type of mutation, with frequencies ranging between 0.004 and 0.039 per division (Figure 4). Based on a genome size of 24.2 Mbp, the rate at which single-nucleotide mutations occurred was between $1.7 \cdot 10^{-10}$ and $1.6 \cdot 10^{-9}$ per nucleotide per cell division, which is similar to a rate of $3.3 \cdot 10^{-10}$ per site per cell division reported for *S. cerevisiae* (Lynch et al. 2008). At a frequency of 0.003 and 0.010 per cell division, INDELs occurred up to 4.1-fold less frequently than SNPs, in accordance with their twofold lower occurrence in *S. cerevisiae* (Lang and Murray 2008). The higher incidence of both SNPs and INDELs in isolates evolved in full-strength wort (Figure 4) may be related to the higher concentrations of ethanol, a known mutagen, in these cultures (Voordeckers, et al. 2015). The rate of loss of mitochondrial DNA varied between 0.0001 and 0.007 per division (Figure 4), and was negatively correlated with the number of generations of selective growth, indicating loss of mitochondrial DNA is selected against. The percentage of respiratory deficient isolates, between 13 and 40 % for all evolutions, is consistent with observations during laboratory evolution under oxygen limitation, and has been associated with increased ethanol tolerance (Ibeas and Jimenez 1997; Taylor, et al. 2002). However, loss of respirative capacity is highly undesirable for lager brewing as it impedes biomass propagation (Gibson, et al. 2008).

Figure 4: Frequencies of different types of mutations observed in evolved isolates obtained after laboratory evolution of strain IMS0408 under simulated lager fermentation conditions. The mutations identified in all 55 isolates evolved under brewing conditions were classified by type, and the frequency of mutation per cell division was calculated for each isolate based on its estimated number of generations of growth under simulated brewing conditions. Average frequencies of mutation types and standard deviations are shown for isolates evolved on full-strength wort at 12 °C (black), on three-fold diluted wort at 12 °C (grey) and on threefold diluted wort at 30 °C (white). The frequencies are shown on a logarithmic scale, and p-values were determined using Student's t-test.

The frequency of chromosomal recombinations was estimated between 0.002 and 0.005 per division (Figure 4), which is similar to frequencies reported for *S. cerevisiae* (Dunham, et al. 2002). The observed recombinations were not reciprocal translocations. Instead, in all cases, genetic material was lost due to internal deletions, or genetic material from one chromosome was replaced by an additional copy of genetic material from another chromosome. This abundant loss of heterozygosity is consistent with the evolutionary history of *S. cerevisiae* (Magwene, et al. 2011) and with previously observed loss of genetic material in hybrids (Sipiczki 2008; Peris, et al. 2017; Smukowski Heil, et al. 2017; Smukowski Heil, et al. 2018). Under selective conditions, regions from one subgenome can be preferentially affected by loss of heterozygosity due to the fitness effects of genes they harbour. Due to its irreversibility, loss of heterozygosity is a determining mechanism for the evolution of hybrids during domestication (Pérez Través, et al. 2014). For example, in an *S. cerevisiae* x *S. uvarum* hybrid, the chromosomal region harbouring *ScPHO84* was preferentially retained at 30 ° C, while it was lost at the expense of its *S. uvarum* homolog at 15 ° C (Smukowski Heil, et al. 2017; Smukowski Heil, et al. 2018). Similarly, the superior growth of *S. cerevisiae* at 30 °C relative to *S. paradoxus* could be linked to *S. cerevisiae* alleles of 8 genes by investigating the effect of loss of heterozygosity in a laboratory hybrid (Weiss, et al. 2018). Moreover, the deletion of *S. cerevisiae* alleles in *S. uvarum* x *S. cerevisiae* had strong and varying impacts on fitness under glucose-, sulphate- and phosphate-limitation (Lancaster, et al. 2018). Overall, loss of heterozygosity is an irreversible process which enables rapid adaptation of hybrid genomes to the selective pressure of their growth environment. In the present study, loss of heterozygosity notably affected *ScMAL11* and *SeSFL1*, contributing to the acquired flocculation and loss of maltotriose utilization phenotypes. Other observed losses of heterozygosity may be due to genetic drift, or they may yield a selective advantage which we did not identify. Loss of heterozygosity may be advantageous by removing unfavourable dominant alleles, by enabling the expression of favourable recessive alleles, or by resolving redundancy, cross-talk and possible incompatibility between alleles and genes of both subgenomes (Piatkowska, et al. 2013; Gibson and Liti 2015).

Whole-chromosome aneuploidy was the rarest type of mutation, occurring only between 0 and 0.0003 times per division (Figure 4). The emergence of single chromosome aneuploidy in 9 % of the isolates after laboratory evolution is similar to observations during laboratory evolution of *S. cerevisiae* (Yona, et al. 2012; Voordeckers, et al. 2015; González-Ramos, et al. 2016; Gorter de Vries, Pronk, et al. 2017). However, the observed small extent of aneuploidy starkly contrasts with the massive aneuploidy of *S. pastorianus* brewing strains (Van den Broek, et al. 2015; Okuno, et al. 2016). These differences might of course be attributed to the difference in time scale between four months of laboratory evolution and several centuries of domestication. However, they might also be due to differences between the *S. cerevisiae* x *S. eubayanus* IMS0408 evolved in this study and the ancestral *S. pastorianus* hybrid. Firstly, IMS0408 was obtained by crossing a haploid *S. cerevisiae* strain with a haploid *S. eubayanus* spore (Hebly, et al. 2015). In contrast, the progenitor of brewing strains of *S. pastorianus* may have resulted from a cross of higher ploidy strains (Okuno, et al. 2016). Since higher ploidy leads to higher chromosome missegregation rates, they accumulate chromosome copy number changes faster (Storchova 2014). In addition, the higher initial ploidy leads to a smaller relative increase of genetic material when an additional copy is gained, which may ‘buffer’ deleterious effects of further changes in ploidy (Torres, et al. 2007). Moreover, the ancestral *S. pastorianus* hybrid may have had, or have acquired, mutations that stimulate extensive aneuploidy, such as mutations increasing the rate of chromosome missegregation or mutations increasing the tolerance against aneuploidy associated stresses (Torres, et al. 2010). Regardless of the origin of the extensive aneuploidy of *S. pastorianus*, our results show that euploid *S. cerevisiae* x *S. eubayanus* hybrids are not by definition prone to extensive aneuploidy under brewing-related experimental conditions. For industrial applications, the relative genetic stability of newly generated *Saccharomyces* hybrid strains reduces the chance of strain deterioration during the many generations involved in large-scale fermentation and/or biomass recycling (Krogerus, et al. 2017).

In addition to showing extensive aneuploidy, *S. pastorianus* strains harbour numerous chromosomal recombinations. During the laboratory evolution experiments, two types of

recombinations were observed: (i) intrachromosomal recombinations resulting in loss of chromosome segments, and (ii) interchromosomal recombinations resulting in loss of one chromosome segment and replacement by an additional copy of a segment from another chromosome, resulting in loss of heterozygosity. While in *S. cerevisiae*, chromosomal recombinations predominantly occur in repetitive regions of the genome (Dunham, et al. 2002; Fontdevila 2005), here 88 % of the observed recombinations occurred within ORFs. The average homology of the recombined ORFs did not exceed the average 85 % homology of ORFs in the *S. cerevisiae* and *S. eubayanus* subgenomes. Instead, the high rate of recombinations at ORFs could reflect a correlation between transcriptional activity and recombination (Thomas and Rothstein 1989). In all cases, the reading frames were conserved, resulting in chimeric ORFs which could encode functional chimeric proteins, with altered length and sequences compared to the parental genomes. The potential selective advantage of such chimeric proteins is illustrated by recurring recombinations between ammonium permease *MEP2* alleles in a *S. cerevisiae* and *S. uvarum* hybrid during laboratory evolution under nitrogen limitation conditions (Dunn, et al. 2013). The formation of chimeric ORFs has even led to the emergence of novel gene functions, as illustrated by the formation of a maltotriose transporter by recombination of three non-maltotriose transporter genes in *S. eubayanus* during laboratory evolution (Brouwers, et al. 2018).

The predominant occurrence of recombinations within ORFs in the evolved isolates has also been observed in the genomes of brewing strains of *S. pastorianus*, which all share identical recombinations at the *ZUO1*, *MAT*, *HSP82* and *XRN1/KEM1* loci (Hewitt, et al. 2014; Walther, et al. 2014; Okuno, et al. 2016). These common recombinations suggest that all *S. pastorianus* isolates descend from a common ancestor (Monerawela and Bond 2018). However, since identical recombinations have been observed in independent evolutions, identical recombinations might reflect parallel evolution due to a strong selective advantage under brewing-related conditions and/or a predisposition of specific loci for recombination (Dunham, et al. 2002; Dunn, et al. 2013). Of the recombination loci found in the present study, only *EFT1* and *MAT* loci were associated with

recombinations in *S. pastorianus*. Moreover, the recombinations at these loci in the evolved isolates were different from those in *S. pastorianus* (Hewitt, et al. 2014; Walther, et al. 2014; Okuno, et al. 2016; Monerawela and Bond 2017). All interchromosomal recombinations observed in this study were unique. These results, obtained under brewing-related conditions, are consistent with the notion that recombination sites are largely aleatory and that all modern *S. pastorianus* strains descend from a single hybrid ancestor.

Different recombination events resulted in the loss of heterozygosity, in four isolates each, of the right arm of *SeCHR XV*, including *SeSFL1*, and of the right arm of *ScCHR XI*, including *ScMAL11*. These events directly affected two phenotypes relevant for brewing fermentation: calcium-dependent flocculation, which led to fast biomass sedimentation, and loss of maltotriose utilisation. Biomass sedimentation can be strongly selected for in sequential batch bioreactors, as it increases the chance that cells are retained in the bioreactor during the emptying phase (Oud, et al. 2013; Hope, et al. 2017). A similar selective advantage is likely to have played a role in the early domestication of *S. pastorianus*, as sedimenting yeast remaining in fermentation vessels was more likely to be used in a next fermentation. Flocculation is a key characteristic of current lager-brewing yeasts (also referred to as bottom-fermenting yeasts), as it simplifies biomass separation at the end of the fermentation (Ferreira, et al. 2010). The present study illustrates how this aspect of brewing yeast domestication can be rapidly reproduced under simulated laboratory conditions.

At first glance, loss of the ability to utilize maltotriose, an abundant fermentable sugar in wort, appears to be undesirable from an evolutionary perspective. However, as demonstrated in studies on laboratory evolution of *S. cerevisiae* in sequential batch cultures on sugar mixtures, the selective advantage of consuming a specific sugar from a sugar mixture correlates with the number of generations of growth on that sugar during each cultivation cycle (Wisselink, et al. 2009; Verhoeven, et al. 2018). *Saccharomyces* yeasts, including strain IMS0408 generally prefer glucose and maltose over maltotriose (Alves, et al. 2008; Hebly, et al. 2015; Brickwedde, et al. 2017). As a

consequence, maltotriose consumption from wort typically only occurs when growth has already ceased due to oxygen limitation and/or nitrogen source depletion, which results in few or no generations of growth on this trisaccharide. However, loss of maltotriose utilization in six isolates in two independent evolution experiments strongly suggests that loss of *ScMAL11* expression was not merely neutral but even conferred a selective advantage. These results are consistent with the existence of many *S. pastorianus* strains with poor maltotriose utilisation and with the truncation of *ScMAL11* in all *S. pastorianus* strains, including good maltotriose utilizers (Vidgren, et al. 2009; Gibson, et al. 2013). In the latter strains, maltotriose utilisation depends on alternative transporters such as Mty1 (Dietvorst, et al. 2005; Salema-Oom, et al. 2005). It is therefore unclear if a selective advantage of the loss of *ScMAL11* reflects specific properties of this gene or its encoded transporter or, alternatively, a general negative impact of maltotriose utilisation under brewing-related conditions. In analogy with observations on maltose utilization by *S. cerevisiae*, unrestricted Mal11-mediated maltotriose-proton symport might cause maltotriose-accelerated death (Postma, et al. 1990; Jansen, et al. 2004). Alternatively, expression of the Mal11 transporter might compete with superior maltose transporters for intracellular trafficking, membrane integration and/or membrane space (Vidgren 2010; Libkind, et al. 2011). Indeed, laboratory evolution to obtain improved maltotriose utilisation resulted in reduced maltose uptake in *S. pastorianus* (Brickwedde, et al. 2017).

Saccharomyces hybrids are commonly applied for industrial applications such as lager beer brewing and wine fermentation (Naumov, et al. 2001; González, et al. 2006; Libkind, et al. 2011). Recently, novel hybrids have been generated, that performed well under a broad range of industrially-relevant conditions (Bellon, et al. 2011; Hebly, et al. 2015; Lopandic, et al. 2016; Krogerus, et al. 2017; Nikulin, et al. 2017; Peris, et al. 2017). While the performance and phenotypic diversity of laboratory hybrids support their application in industrial processes, further strain development of such hybrids could improve their performance (Steensels, et al. 2014; Krogerus, et al. 2017). Especially in food and beverage fermentation processes, consumer acceptance issues largely preclude use of targeted genetic modification techniques. Laboratory evolution offers an interesting

307 alternative strategy for ‘non-GMO’ strain improvement (Bachmann, et al. 2015). However, as
 308 exemplified by the loss, in independent laboratory evolution experiments, of *MAL11* and of the
 309 mitochondrial genome, mutations that yield increased fitness under simulated industrial
 310 fermentation conditions are not necessarily advantageous for industrial performance. Therefore,
 311 instead of faithfully reconstructing industrial conditions in the laboratory, laboratory evolution
 312 experiments should be designed to specifically select for desired phenotypes. For example, a recent
 313 study illustrated how maltotriose fermentation kinetics of an *S. pastorianus* hybrid could be
 314 improved by laboratory evolution in carbon-limited chemostats grown on a maltotriose-enriched
 315 sugar mixture (Brickwedde, et al. 2017).

Materials and Methods

Yeast strains and media

Saccharomyces strains used in this study are listed in Supplementary Table 3. Yeast strains and *E. coli* strains containing plasmids were stocked in 1 mL aliquots after addition of 30 % v/v glycerol to the cultures and stored at -80 °C. For preparation of stock cultures and inocula of bioreactors, yeast strains were routinely propagated in shake flasks containing 100 mL YPD (10 g.L⁻¹ yeast extract, 20 g.L⁻¹ yeast peptone and 20 g.L⁻¹ glucose) at 30 °C and 200 RPM in an Brunswick Innova43/43R shaker (Eppendorf Nederland B.V., Nijmegen, The Netherlands). For cultivation on solid media, YPD medium was supplemented with 20 g.L⁻¹ Bacto agar (Becton Dickinson, Breda, The Netherlands) and incubation was done at 30 °C. Synthetic medium (SM), containing 3 g.L⁻¹ KH₂PO₄, 0.5 g.L⁻¹ MgSO₄·7H₂O, 5 g.L⁻¹ (NH₄)₂SO₄, 1 mL.L⁻¹ of a trace element solution and 1 mL.L⁻¹ of a vitamin solution, was prepared as previously described (Verduyn, et al. 1992). SM maltotriose was supplemented with 20 g.L⁻¹ of maltotriose and SM ethanol with 20 mL.L⁻¹ of ethanol. Selection for the amdS marker was performed on SM-AC: SM with 0.6 g.L⁻¹ acetamide and 6.6 g.L⁻¹ K₂SO₄ instead of (NH₄)₂SO₄ as nitrogen source (Solis-Escalante, et al. 2013). For counter selection of the amdS marker, strains were first grown on YPD and then on SM-FAC: SM supplemented with 2.3 g.L⁻¹ fluoroacetamide (Solis-Escalante, et al. 2013). Industrial wort was provided by HEINEKEN Supply Chain B.V., Zoeterwoude, the Netherlands, and contained 14.4 g/L glucose, 2.3 g/L fructose, 85.9 g/L maltose, 26.8 g/L maltotriose and 269 mg/L free amino nitrogen. The wort was supplemented with 1.5 mg.L⁻¹ Zn²⁺ by addition of ZnSO₄·7H₂O, then autoclaved for 30 min at 121 °C and, prior to use, filtered through Nalgene 0.2 µm SFCA bottle-top filters (ThermoFisher Scientific, Waltham, MA). For experiments performed with diluted wort, two volumes of sterile demineralized water were added per volume of wort. To prevent excessive foaming during the aeration phase of the bioreactor experiments, (un)diluted wort was supplemented with 0.2 mL.L⁻¹ of sterile Pluronic PE 6100 antifoam (Sigma-Aldrich, Zwijndrecht, the Netherlands).

Table 3: Plasmids used throughout this study.

Name	Relevant genotype	Origin
pUD711	ori <i>bla</i> gRNA- <i>ScSFL1</i>	GeneArt™
pUD712	ori <i>bla</i> gRNA- <i>SeSFL1</i>	GeneArt™
pUDE481	ori <i>bla</i> ARS4/CEN6 <i>hyg^R</i> <i>ScTDH3p</i> -mTurquoise2- <i>ScADH1t</i>	(Gorter de Vries, et al. 2018)
pUDE482	ori <i>bla</i> ARS4/CEN6 <i>hyg^R</i> <i>ScTEF1p</i> -Venus- <i>ScENO2t</i>	(Gorter de Vries, et al. 2018)
pUDP004	ori <i>bla</i> panARSopt amdSYM <i>Spcas9</i>	(Gorter de Vries, de Groot, et al. 2017)
pUDP045	ori <i>bla</i> panARSopt amdSYM <i>Spcas9</i> gRNA- <i>ScMAL11</i>	(Gorter de Vries, et al. 2018)
pUDP104	ori <i>bla</i> panARSopt amdSYM <i>Spcas9</i> gRNA- <i>ScSFL1</i>	This study
pUDP105	ori <i>bla</i> panARSopt amdSYM <i>Spcas9</i> gRNA- <i>SeSFL1</i>	This study

Analytical methods and statistics

Optical density at 660 nm was measured with a Libra S11 spectrophotometer (Biochrom, Cambridge, UK). HPLC analysis of sugar and metabolite concentrations was performed with an Agilent Infinity 1260 chromatography system (Agilent Technologies, Santa Clara, CA) with an Aminex HPX-87 column (Bio-Rad, Lunteren, The Netherlands) at 65 °C, eluted with 5 mM H₂SO₄. Significance of data was assessed by an unpaired two-tailed Student's t-test with a 95 % confidence interval.

Laboratory evolution and single colony isolation

The hybrid yeast strain IMS0408 was evolved under three different conditions in duplicate in Minifors 2 bioreactors (INFORS HT, Velp, the Netherlands) with a working volume of 100 mL: on diluted wort at 30 °C (LG30.1 and LG30.2), on diluted wort at 12 °C (LG12.1 and LG12.2) and on full-strength wort at 12 °C (HG12.1 and HG12.2). Sequential batch cultivation was performed with 10 and 30 mL.min⁻¹ of headspace N₂ flushing at 12 and 30 °C, respectively. The percentage of CO₂ in the outlet gas stream, the culture pH and the dissolved oxygen concentration in the broth were continuously monitored. The end of a batch cultivation cycle was automatically triggered when the percentage of CO₂ in the offgas decreased below 75 % and 10 % of the maximum value reached during that cycle for growth on diluted wort and full-strength wort, respectively. These CO₂ percentages correspond to the moment at which sugar utilisation was complete in the first batch cycle for each condition, as determined by HPLC measurements. When the CO₂ threshold was reached, the reactor was emptied while stirring at 1200 RPM leaving about 7 mL to inoculate the next batch. Upon addition of fresh medium, the broth was stirred at 500 RPM and sparged with 500 mL.min⁻¹ of pressurized air during 5

min for diluted wort or 12 h for wort. During the remainder of each batch cultivation cycle, the medium was not sparged or stirred and the pH was not adjusted. LG30.1 and LG30.2 were carried out for 116 and 117 cycles respectively, LG12.1 and LG12.2 were carried out for 29 cycles and HG12.1 and HG12.2 were carried out for 13 and 16 cycles, respectively. Culture samples from all six reactors were then streaked on YPD plates and after three subsequent restreaks, frozen stock cultures of single colony isolates were prepared. By default, five isolates were obtained for each culture. For LG12.1 and LG30.1, two different colony morphologies were observed, therefore five elevated and conically-shaped colonies and five regular flat colonies were stocked. The experiments at 12 °C on diluted wort were continued for four months until a total of 58 and 57 cycles was reached for LG12.1 and LG12.2, respectively, and five single-cell isolates were obtained for each reactor as described above.

Genomic DNA extraction and whole genome sequencing

Yeast cultures were incubated in 500-mL shake-flasks containing 100 mL YPD at 30°C on an orbital shaker set at 200 RPM until the strains reached stationary phase at an OD₆₆₀ between 12 and 20. Genomic DNA was isolated using the Qiagen 100/G kit (Qiagen, Hilden, Germany) according to the manufacturer's instructions and quantified using a Qubit® Fluorometer 2.0 (ThermoFisher Scientific). For IMS0408 and the evolved isolates, genomic DNA was sequenced at Novogene Bioinformatics Technology Co., Ltd (Yuen Long, Hong Kong) on a HiSeq2500 sequencer (Illumina, San Diego, CA) with 150 bp paired-end reads using PCR-free library preparation.

Genome analysis

A high quality reference genome was constructed by combining near-complete assemblies of *S. cerevisiae* CEN.PK113-7D (Salazar, et al. 2017) and *S. eubayanus* CBS12357^T (Brickwedde, et al. 2018). The kanMX marker present in IMS0408 was inserted as an additional contig (Wach, et al. 1994). For each evolved strain, raw Illumina reads were aligned against the reference genome using the Burrows–Wheeler Alignment tool (BWA, version 0.7.15-r1142) and further processed using SAMtools (version 1.3.1) and Pilon (version 1.18) for variant calling (Li, et al. 2009; Li and Durbin 2010; Walker,

et al. 2014). SNPs and INDELs that were also called or which were ambiguous in IMS0408, were disregarded. Copy number was determined based on read coverage analysis. Chromosomal translocations were detected using Breakdancer (version 1.3.6) (Chen, et al. 2009). Only translocations which were supported by at least 10 % of the reads aligned at that locus and which were absent in strain IMS0408 were considered. All SNPs, INDELs, recombinations and copy number changes were manually confirmed by visualising the generated .bam files in the Integrative Genomics Viewer (IGV) software (Robinson, et al. 2011). A complete list of identified mutations is provided in Supplementary Data File 1. For chimeric open-reading-frame reconstruction, reads aligning within 3 kbp of an identified recombination site and their paired reads were extracted using Python and were assembled using SPAdes (Bankevich, et al. 2012). The resulting contigs were aligned against ORFs of genes the genes affected by the recombination to identify the recombination point, and the complete recombined ORF was reconstructed. Original and recombined ORFs were then aligned and translated using CloneManager (version 9.51, Sci-Ed Software, Denver, CO) to determine whether the translocation had introduced frameshifts or premature stop codons.

DNA content determination by flow cytometric analysis.

Exponential-phase shake flask cultures on YPD were diluted to an OD₆₆₀ of 1. A 1 mL sample (approximately 10⁷ cells) was then washed in cold demineralized water and resuspended in 800 µL 70 % ethanol while vortexing. After addition of another 800 µL 70 % ethanol, fixed cells were stored at 4 °C until further staining and analysis. DNA was then stained with SYTOX Green as described previously (Haase and Reed 2002). Samples were analysed on a Accuri C6 flow cytometer (BD Biosciences, Franklin Lakes, NJ) equipped with a 488-nm laser and the median fluorescence of cells in the 1n and 2n phases of the cell cycle was determined using FlowJo (BD Biosciences). The 1n and 2n medians of strains CEN.PK113-7D (n), CEN.PK122 (2n) and FRY153 (3n) were used to create a standard curve of fluorescence versus genome size with a linear curve fit, as performed previously (Van den Broek, et al. 2015). The genome size of each tested strain was estimated by averaging predicted genome sizes of the 1n and 2n population in assays on three independent cultures.

Identification of strains with respiratory deficiency

Respiratory competence was assessed through their ability to grow on ethanol. Samples from 24 h shake-flask cultures on YPD (30 °C, 200 RPM) were washed twice with demineralized water and used to inoculate duplicate aerobic shake flasks containing 100 mL of SM with 2 % ethanol to an OD₆₆₀ of 0.2. After 72 h incubation at 30 °C and 200 RPM, OD₆₆₀ was measured.

Assay for calcium-dependence of flocculation

Two 100 µL aliquots from overnight cultures on YPD were washed with sterile demineralized water. One aliquot was resuspended in demineralized water and the other in 50 mM EDTA (pH 7.0). Both samples were imaged at 100 x magnification under a Z1 microscope (Carl Zeiss BV, Breda, the Netherlands) to assess flocculence and its reversal by EDTA chelation of calcium ions.

Plasmid construction

All plasmids were propagated in *E. coli* DH5α (Table 3). Plasmid pUD711 and pUD712 were designed as previously described (Gorter de Vries, de Groot, et al. 2017) and de novo synthesised at GeneArt (ThermoFisher Scientific) containing the sequence 5' GGTCTCGCAAATAACAAGTATGAGTCCGTGAGGACGAAACGAGTAAGCTCGTCTTGTTATAGTCACGGATCG AGTTTTAGAGCTAGAAATAGCAAGTTAAAATAAGGCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTCG GTGCTTTTGGCCGGCATGGTCCCAGCCTCCTCGCTGGCGCCGGCTGGGCAACATGCTTCGGCATGGCGAATGG GACACAGGCAAATTCATCTGATGAGTCCGTGAGGACGAAACGAGTAAGCTCGTCATGAATATCGCATTTTGT GGGTTTTAGAGCTAGAAATAGCAAGTTAAAATAAGGCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTC GGTGCTTTTGGCCGGCATGGTCCCAGCCTCCTCGCTGGCGCCGGCTGGGCAACATGCTTCGGCATGGCGAATG GGACACAGCGAGACC 3' for pUD711 and 5' GGTCTCGCAAATAACAAGTATGAGTCCGTGAGGACGAAACGAGTAAGCTCGTCTTGTTATAGTCACGGATCG AGTTTTAGAGCTAGAAATAGCAAGTTAAAATAAGGCTAGTCCGTTATCAACTTGAAAAAGTGGCACCGAGTCG GTGCTTTTGGCCGGCATGGTCCCAGCCTCCTCGCTGGCGCCGGCTGGGCAACATGCTTCGGCATGGCGAATGG GACACAGCGAGACC 3' for pUD712. Plasmid pUDP104, expressing gRNA_{ScsFL1} and *cas9*, was constructed by Golden Gate cloning by digesting pUDP004 and pUD711 using BsaI and ligating with T4 ligase (Engler, et al. 2008). Similarly, plasmid pUDP105, expressing gRNA_{SeSFL1} and *cas9*, was

constructed from pUDP004 and pUD712. Correct assembly was verified by restriction analysis using P_{ddl}.

Strain construction

The *ScTEF1p*-Venus-*ScENO2t* repair fragment with flanks for homologous recombination in the *ScMAL11* locus was PCR amplified from plasmid pUDE481 using primers 12989 and 12990 (Table 4). The *ScTDH3p*-mTurquoise2-*ScADH1t* repair fragment with flanks for homologous recombination in the *ScSFL1* locus was PCR amplified from plasmid pUDE482 using primers 13564 and 13565. The *ScTEF1p*-Venus-*ScENO2t* repair fragment with flanks for homologous recombination in the *SeSFL1* locus was PCR amplified from plasmid pUDE481 using primers 13566 and 13567.

Table 4: Primers used in this study.

name	Sequence (3' to 5')	Purpose
12989	TTGGTGTTCCTTCTGATGCTACATAGAAGAACATCAACAACCTAAAAAATAGTATAATACACACT GGCTTAAGATGAC	Insertion Venus in <i>MAL11</i>
12990	TTGGGAGCAGTCAAGGGATTCTTATTTCTTCCAAAAAACAACCTTTTACCGTCTCA TGCTCAGC	Insertion Venus in <i>MAL11</i>
13564	ACTTTTCAGCTAGCAAGAAGGATCTCTTTTAAACACTCTATACAGGTGCACACAAAGGACCAGAT GTCAACACAGCTAC	Insertion mTurquoise2 in <i>ScSFL1</i>
13565	CGGAGTTGGTAAAAATATAGTTATAATCACAAGGATCAGGAGGAAAAAGAAAAAAGTGATTTAT CGTCTCACATCCAGC	Insertion mTurquoise2 in <i>ScSFL1</i>
13566	GTAAGAAAGCAGAAAAAAGAGAAAAAGAAACGATCTCGACTAGAGTGACGGGTTGACACA CTGGCTTAAGATGAC	Insertion Venus in <i>SeSFL1</i>
13567	AAAATGGAAGGGTAAAAATGGAAGGAAGAAGTGAAAGTGAAAAATACAATGTCGTC TCATGCTCAGC	Insertion Venus in <i>SeSFL1</i>

All strains were transformed by electroporation as described previously, with 300 ng of gRNA/Cas9 expression plasmid and 1 µg of repair fragment (Gorter de Vries, de Groot, et al. 2017). Strains IMX1698 (mVenus::Δ*ScMAL11*), IMX1824 (mTurquoise2::Δ*ScSFL1*) and IMX1825 (Venus::Δ*SeSFL1*) were constructed by transforming IMS0408 with the appropriate repair fragments and plasmids pUDP045, pUDP104 and pUDP105, respectively. Strain IMX1826 (mTurquoise2::Δ*ScSFL1* Venus::Δ*SeSFL1*) was constructed by transforming IMX1824 using the appropriate repair fragment and plasmid pUDP105. After electroporation, cells were transferred to 20 mL SM-Ac medium to select successful transformants and incubated at 30 °C for 3 to 5 days. After growth was observed, 200 µL of culture was transferred to 20 mL fresh SM-Ac and incubated similarly during 24h. Finally, 200 µL from the second culture was transferred to 20 mL fresh YPD medium to maximize expression

of fluorescent proteins. Successfully gene-edited cells were sorted using the BD FACSaria™ II SORP Cell Sorter (BD Biosciences) as described previously (Gorter de Vries, et al. 2018). The plasmids were cured from strains IMX1698, IMX1824, IMX1825 and IMX1826 by subsequent growth on YPD and plating on SM-FAC. After confirmation of the correct genotype by colony PCR, randomly picked colonies were used to prepare frozen stocks.

Biomass sedimentation assay

IMS0408, IMS0556, IMS0558, IMS0559, IMS0617, IMX1824, IMX1825 and IMX1826 were grown in triplicate during 72h in vented 50 mL Bio-One Cellstar Cellreactor tubes (Sigma-Aldrich) on 20 mL YPD at 30 °C and 200 RPM until stationary phase. For each sample, the biomass was resuspended by vigorous vortexing and 1 mL was sampled immediately after vortexing from right underneath the meniscus. After 60 s of stationary incubation, another sample was taken by the same procedure. Biomass sedimentation was quantified as the ratio of the OD₆₆₀ values of the two samples.

Evaluation of maltotriose fermentation

Each strain was grown microaerobically in 100 mL serum bottles containing 100 mL medium and shaken at 200 RPM. Medium (full-strength or diluted wort) and incubation temperature (12 °C or 30 °C) were the same as in the evolution experiment from which a strain had been isolated. Strain IMS0408 was included as a control for each condition. Bottles were inoculated to an OD₆₆₀ of 0.2 from aerobic shake-flask precultures grown on same medium and under the same conditions. During cultivation, 8 to 12 samples were taken at regular intervals for OD₆₆₀ measurements and metabolite analysis by HPLC. When no further sugar consumption was recorded over an interval of at least 48h, the fermentation was considered finished.

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Author contributions

ARGdV planned and executed the laboratory evolution, made single colony isolates, prepared genomic DNA and characterised ploidy and respiratory deficiency. ARGdV, ACAvA and LHK characterised flocculation and maltotriose utilisation of the isolates. ARGdV and MAV performed the reverse engineering of deletions of *SFL1* and *MAL11* and characterized the reverse engineered phenotypes. ARGdV and MvdB performed the bioinformatics analysis. ARGdV, JTP and JMGD wrote the manuscript. AS, NB and TA provided critical feedback throughout the study. JMGD and JTP supervised the study. All authors read and approved the final manuscript.

Disclosure declaration

The authors declare no competing financial interests.

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