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Experimental determination of codon usage-dependent selective pressure on high copy-number genes in *Saccharomyces cerevisiae*

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Keywords: codon usage, yeast, evolution, selective pressure, recombinant gene expression, plasmid copy number

Abstract

One of the central hypotheses in the theory of codon usage evolution is that in highly expressed genes particular codon usage patterns arise because they facilitate efficient gene expression and are thus selected for in evolution. Here we use plasmid copy number assays and growth rate measurements to explore details of the relationship between codon usage, gene expression level, and selective pressure in *Saccharomyces cerevisiae*. We find that when high expression levels are required optimal codon usage is beneficial and provides a fitness advantage, consistent with evolutionary theory. However, when high expression levels are not required, optimal codon usage is surprisingly and strongly selected against. We show that this selection acts at the level of protein synthesis, and we exclude a number of molecular mechanisms as the source for this negative selective pressure including nutrient and ribosome limitations and proteotoxicity effects. These findings inform our understanding of the evolution of codon usage bias, as well as the design of recombinant protein expression systems.

Keywords: codon usage, yeast, evolution, selective pressure, recombinant gene expression, plasmid

Introduction

Because the genetic code uses 64 codons to encode 20 amino acids (Crick, Barnett, Brenner, & Watts-Tobin, 1961), most amino acids can be encoded by multiple synonymous codons. In all organisms where codon usage has been investigated, synonymous codons are not used randomly but some codons are preferred over others (see eg Behura & Severson, 2013) for a recent review). Correlations between biased codon usage and a number of other parameters have been detected including cellular tRNA content (Ikemura, 1982), translational efficiency (Sharp & Li, 1986), translational accuracy (Zhou, Weems, & Wilke, 2009), RNA structure (Hartl, Moriyama, & Sawyer, 1994), protein structure (Oresic, Dehn, Korenblum, & Shalloway, 2003), genomic GC content (Comeron, Kreitman, & Aguade, 1999), recombination (Comeron et al., 1999), splicing (Chamary & Hurst, 2005), and gene conversion rates (Galtier, 2003).

Translational efficiency has gained experimental support as one causative parameter that can lead to codon usage bias (Carlini & Stephan, 2003; Chu et al., 2014; Hense et al., 2010; Zhou et al., 2013). Since synonymous mutations can have substantial effects on expressed protein levels, selection for high gene expression levels will favour codon usage patterns compatible with such high levels, but will avoid patterns that restrict attainable expression levels. However, the exact relationship between codon usage, protein expression, other associated parameters and selective pressure are still unclear.

One issue is that selective pressure is difficult to measure directly as an experimental parameter. Moriya *et al.* used copy number variations in high copy number vectors in *Saccharomyces cerevisiae* to estimate the direction and extent of selective pressure when these vectors contained different genes (Makanae, Kintaka, Makino, Kitano, & Moriya, 2013; Moriya, Shimizu-Yoshida, & Kitano, 2006). While natural 2 micron circles employ a sophisticated system to stabilise copy numbers (McQuaid et al., 2017), yeast cloning vectors containing solely the 2 micron origin of replication exhibit strong copy number variations and appear to be inherited in a stochastic fashion (Moriya et al., 2006). In cases where such vectors contain genes that affect the growth rate of the cell in a copy-number dependent fashion, cells with reduced or increased copy number (depending on whether the gene in question has a negative or positive effect on growth rate) have a growth advantage. Since such cells are also more likely to produce daughter cells with copy numbers that are lower/higher than the population average, the average copy number per cell changes until a new balance is achieved between the effect of the gene on growth rates, the metabolic cost of maintaining high numbers of vectors in the cell, and the ability to successfully inherit the vector and its selectable marker to the majority of daughter cells.

Here, we introduce reporter genes with differing codon usage into 2 micron cloning vectors and study the effect of varying codon usage on the high-copy number vector content and growth rate of transformed yeast cells. Surprisingly, our results suggest that codon optimisation is selected against unless high level expression of the gene provides a competitive growth advantage.

Results

Selection against high copy numbers of a codon optimised gene in yeast

We previously described a dual luciferase reporter system allowing us to assess the effect of codon usage variation on Firefly luciferase expression levels (Chu et al., 2014; Chu, Barnes, & von der Haar, 2011). The original reporter system was implemented using centromeric (low copy number) vectors, which contained three different Firefly luciferase variants encoded only by unfavourable codons (minFLuc), by the wild-type gene from *Photinus pyralis* which has random codon usage from the point of view of yeast (staFLuc), or by the fully codonadapted gene (maxFLuc). All vectors also contained an invariant *Renilla* luciferase gene for normalisation. For this vector system we had demonstrated that the expression ratio measured using dual luciferase assays corresponds well to the expression ratio measured by western blotting (Chu et al., 2011). We were therefore surprised to observe that, when the same expression system was implemented in multi-copy vectors based on the 2 micron origin of replication, the codon-dependent variation in expression levels was indistinguishable from the centromeric system when measured in dual luciferase assays, but differed markedly when assessed using western blots (figure 1). Note that all experiments shown in the current study were conducted with cytoplasmic firefly luciferase variants which contain deletions of the last three amino acids to remove a peroxisomal targeting signal (Chu et al., 2014).

We reasoned that our results could be explained if plasmid copy numbers changed substantially between expression plasmids containing the different codon usage variants. The dual luciferase assay measures expression normalised per plasmid, whereas the western blotting approach measures total numbers per cell, so that changes in plasmid copy number would only become visible in the latter assay. To test this assumption directly we measured the plasmid copy number for the different luciferase variants using a qPCR approach, and found that introduction of poorly adapted, wild type, and strongly codon adapted sequences sequentially reduced the steady-state plasmid copy number over a four-fold range compared to the vector control (a plasmid which only expresses *Renilla* luciferase but no firefly luciferase, figure 2b). This implies that expression of a strongly codon-adapted gene is less favourable and more strongly selected against than expression of weakly codon adapted genes. This effect is also visible at the levels of the growth rates of the respective transformants, which become successively reduced with increasing codon adaptation of the recombinant genes (figure 2c). This reduced growth rate appears due to lower cell division rates, as we do not detect any changes in the proportion of viable cells (data not shown).

Copy number selection is dependent on translational activity

To test whether selection against codon optimised genes in this system arose at the level of protein synthesis, we introduced a point mutation into the codon adapted maxFLuc construct that resulted in a premature stop signal at the fifth codon of the open reading frame (maxFLuc K5X). This prevents almost all of the translational activity on the mRNA, without significantly affecting the overall sequence or nucleotide composition of the construct nor the transcriptional activity of the promoter. The K5X mutation substantially rescued both the reduction in plasmid copy number and the reduction in growth rate observed with the

translated maxFLuc construct (figure 3). Thus, the major effect on fitness requires ongoing translation of the full-length ORF.

We explored a number of potential mechanisms which by our reasoning could have caused the selection against codon optimisation. We determined that the three constructs produced approximately 2.1×10^{-14} , 4.5×10^{-14} and 5.9×10^{-14} g firefly luciferase protein per cell (figure 4A). In other words, the highest expressing construct produced just over 1% of the 5×10^{-12} g of protein contained in a haploid yeast cell (von der Haar, 2007). Most of the resources required for protein synthesis, including amino acids and energy, should scale linearly with the amount of protein made, and a 1-2% increase in resource usage would be unlikely to cause a 10-20% reduction in growth rate. Consistently, we do not observe any change in the effect of maxFLuc expression on the growth rate if we double the concentration of carbon, nitrogen, or amino acids in the medium (figure 4B).

We and others previously predicted that, under fast growth conditions, providing ribosomes for the expression of additional proteins could only be accommodated at the cost of reduced growth (Chu et al., 2011; Chu & von der Haar, 2012; Shah, Ding, Niemczyk, Kudla, & Plotkin, 2013). We therefore estimated the number of ribosomes required to produce the observed protein levels from the three FLuc variants. This number essentially scales with the number of proteins produced per unit time, and inversely scales with the time spent to complete translation of the ORF (ie more ribosomes are needed to make more protein at the same speed, but less ribosomes are needed to make the same amount of protein at higher speed). In the case of our FLuc constructs, the increased speed of translation outweighs the more frequent initiation events for the maxFLuc construct, resulting in a predicted net decrease of ribosome usage with increasing codon optimisation (figure 4C). This decrease in ribosome usage for codon optimised genes is expected, and is thought to be one of the genome-wide drivers for codon optimisation as this maximises cell-wide ribosome availability (Shah & Gilchrist, 2011). In the context of our experiments, it rules out that ribosome shortages explain the selective pressure against codon optimised constructs. We also analysed whether tRNA shortages might explain the observed reduction in growth rate, but again find that tRNAs should actually be most limiting for the least optimised construct which is decoded by the highest proportion of low-abundance tRNAs (the average tRNA gene copy number per codon is 7.6, 10.2 and 12.6 for min, sta and maxFLuc, respectively).

Lastly, we reasoned that simple proteotoxic effects might explain our results. However, the responsiveness of a maxFLuc expressing construct to the proteotoxic drug AZC (Trotter, Berenfeld, Krause, Petsko, & Gray, 2001) is unaltered (figure 4D), even though we would expect that any pre-existing proteotoxicity should increase the sensitivity to this drug. We also did not detect any significant increases in the expression of the disaggregase Hsp104, which is usually induced under conditions of proteotoxic stress (figure 4E, Bösl, Grimminger, & Walter, 2006).

In summary, although we clearly observe negative selective pressure against codon optimised firefly luciferase, our experiments do not offer a simple explanation for this effect.

The effect of high copy number codon adapted genes on fitness is context dependent

While we have no immediate mechanistic explanation for the strong negative selective pressure arising from codon optimisation in our construct, selection against the optimisation of a gene that is of no benefit to the cell makes some sense from an evolutionary perspective. To test how the dynamics of negative and positive selection interplay in the case of proteins that are of benefit to the cell, we repeated the FLuc expression experiments with the *HIS3* gene, a natural yeast gene which catalyses the 6th step of the histidine biosynthesis pathway (Struhl & Davis, 1977). The yeast strain used in our experiments carries a chromosomal *his3* deletion and can only grow in the absence of histidine if the gene is provided extrachromosomally.

We determined the growth rates of yeast strains containing poorly adapted, wild-type, and well adapted *HIS3* genes under three different growth conditions. In the presence of histidine in the growth medium, the gene is not beneficial to the cell, similar to the FLuc genes in the experiments described above. In contrast, in the absence of histidine *HIS3* is essential, although it is not required at high expression levels (Chu et al., 2014). Lastly, in the absence of histidine and the presence of a competitive inhibitor of His3 enzymatic activity, 3-aminotriazole (3-AT), high levels of His3 protein are required to overcome the inhibitor (Durfee et al., 1993).

When His3 is not required, we observe a similar pattern of consecutively reduced fitness as for the FLuc genes (figure 5, upper panel). This confirms that the fitness effects observed upon high level FLuc expression are not specific to heterologous proteins and hold also upon high level expression of a homologous gene. When low level His3 activity is required, the effect of the different constructs on growth rates is initially relatively unaffected, although codon usage effects become statistically insignificant. This lack of significance is possibly attributable to the lower number of repeats used for this condition (figure 5, middle panel). However when very high His3 levels are required in the presence of 3-AT, the codon-optimised gene confers a clear fitness advantage over the less optimised genes (figure 5, bottom panel). Links between preferred codon usage and the requirement for high expression levels have previously been established largely by way of correlation, our observations provide experimental confirmation of this notion.

To ask whether the effect of codon usage on selective pressure is quantitatively determined by basic yeast physiology, or is dependent on the genetic background of the strain in question, we compared the growth rates of a number of genetically tractable strains from the *Saccharomyces* genome resequencing project (SGRP, Cubillos, Louis, & Liti, 2009) in the presence and absence of the maxFLuc gene (figure 6). We found that the proportional reduction in growth rate upon expression of the gene ranged from 10% in the mildest affected strains, to 50% in the worst affected strains. Although many of the SGRP strains flocculate strongly and determination of growth rates for these strains is less accurate than for BY4741, this wide range indicates that the genetic make-up of strains strongly modulates the correspondence between codon usage and selective pressure.

Discussion

Codon usage bias is thought to arise from an interplay between evolutionary drift, ie stochastic changes of one codon for another, and evolutionary selection, ie the effect of growth advantages arising through the use of particular subsets of codons in particular subsets of genes (Shah and Gilchrist, 2011). Our work highlights an unexpected element of such selection which appears to act against the use of normally favoured codons when these are used out of their normal context.

In interpreting our results, it is useful to define in how far plasmid copy number changes reflect selection mechanisms applying during the actual evolution of organisms. Codon usage is thought to have effects at both the individual gene level and at genome-wide levels. Effects at individual gene level include permitting high translational activity or conversely restricting translational activity to low levels (Tarrant & von der Haar (2014) and references therein), as well as effects on RNA stability (Presnyak et al., 2015) and transcriptional activity (Zhou et al., 2016). Genome-wide effects include compliance with particular GC bias (Palidwor, Perkins, & Xia, 2010), optimisation of ribosome usage by reducing the average dwell time of ribosomes on mRNAs (Chu & von der Haar, 2012; Shah et al., 2013) and other effects (Plotkin & Kudla, 2011). Genes contained in high copy number plasmids are in principle subject to effects at both levels, but because changing the sequence of such a plasmid is equivalent to changing the sequence of multiple genes simultaneously in the genome, we expect the selective forces acting on this system to be more similar to genome-wide selective forces than would be the case in a single copy plasmid.

The observation of particularly strong codon usage bias in highly expressed genes (Sharp, Tuohy, & Mosurski, 1986) implies that highly expressed genes show such bias, but also that low-expressed genes do not. It is thought that the absence of bias in genes with low expression levels arises because codon usage is at equilibrium, rather than being selected against (Sharp, Emery, & Zeng, 2010). Interestingly, our data suggest that codon bias can be selected against in genes with low expression levels. Moreover, the successive reduction in plasmid copy numbers with increasing codon optimisation (figure 2) suggests that negative selection is not restricted to extreme codon usage but applies to genes with close to equilibrium codon usage. Consistent with this notion, statistical enrichment of non-preferred codons in low expressed genes has been shown in some studies (Neafsey & Galagan, 2007).

The principal underlying assumption for explaining codon usage bias in highly expressed genes, that codon usage is selected for when high expression levels are required, has widespread support and is generally accepted. However, to our knowledge this has never been directly experimentally tested. By linking codon usage variants of the endogenous yeast *HIS3* gene to growth conditions where this gene is required at different expression levels, we provide an experimental test for this hypothesis. Our findings are entirely consistent with the prevailing hypothesis.

While our results are informative for understanding basic evolutionary mechanisms, they also provide useful information for the design of recombinant protein expression systems. 2 micron vector systems are used for the production of various yeast-derived biopharmaceuticals (Finnis et al., 2010; Gerngross, 2004; Thim et al., 1986). If plasmid copy

numbers in these cases are subject to the same effects we describe here, productivity could be boosted by introducing measures that stabilise plasmid copy number, although this could also produce adverse effects by increasing the selection for non-expressing mutants. The observation of substantial strain variability in the effect of recombinant gene expression on growth rates (figure 6) also indicates that the genetic variability of yeast could be harnessed for balancing these effects in order to stabilise high level expression systems.

Materials and Methods

Strains, plasmids and media. The main yeast strain used was BY4741 (Mata *ura3Δ0 his3Δ0 leu2Δ0 met15Δ0*, Brachmann et al., 1998). Genetically tractable strains from the SGRP collection (Cubillos et al., 2009) were obtained from the National Collection of Yeast Cultures (NCYC, UK, haploid Mata strains from SGRP strain set 2). Luciferase expression vectors were generated by transferring *XmaI/EcoRI* *RLuc* and *BamHI/SalI* *CFLuc* fragments from the centromeric vectors described in (Chu et al., 2014) into pBEVY-U (Miller, Martinat, & Hyman, 1998). *HIS3* expression vectors were generated by cloning the C-terminally HA-tagged *HIS3* variants described in (Chu et al., 2014) as *BamHI/PstI* fragments into the same vector. Plasmid sequences, maps, and the plasmids themselves are available through Addgene (table 1). Yeast strains were transformed as described (Gietz & Schiestl, 2007) and grown in SC -Ura medium (2% glucose, 0.67% Yeast Nitrogen Base without amino acids [BD, UK], and Kaiser Synthetic Complete Drop-Out Mixture lacking uracil [Formedium, UK] as indicated in the manufacturer's instructions).

Growth rate measurements. Growth rates were measured in 24- or 48-well cell culture plates using automated plate readers. 1 ml medium per well (0.5 ml for 48-well plates) were inoculated with material from transformed yeast colonies grown on selective agar plates and grown overnight in a standard shaking incubator. Following overnight growth, 1 ml or 0.5 ml of fresh medium per well contained in a new plate was inoculated with 10 μ l or 5 μ l of culture from the overnight plate. Plates were then incubated in Spectrostar Nano plate readers (BMG Lab Tech, UK) and incubated at 30°C under constant shaking, with automated *oD* measurements every 30 minutes until the culture had reached stationary phase. To analyse the resulting data, logarithmic curves were fitted to seven-time-point windows along the entire incubation time, and the highest growth rate returned from this fitting exercise was reported as the maximum logarithmic growth rate for each well.

qPCR assays. These were adapted from the procedure described in (Moriya et al., 2006). Briefly, 2 *oD*₆₀₀ units of cells were resuspended in 850 μ l of 1.2 M sorbitol, 100 μ l of 200 mM sodium phosphate buffer (pH 7.2), and 50 μ l lyticase (Sigma Aldrich, UK, L2524, resuspended in sodium phosphate buffer at 5 units/ μ l). The suspension was incubated at 37°C for 30 minutes, and then subjected to 3 cycles of incubation for ten minutes at 95°C followed by incubation for fifteen minutes at -80°. The extract was clarified by centrifugation in a microcentrifuge at 13,000 rpm for 5 minutes, and 0.1 μ l of the supernatant were used as input for the qPCR reactions.

For the latter, we used primers against the *URA3* marker gene on the pBEVY-U vectors (qURA3f, AGCAGAATTGTCATGCAAGG, qURA3r, TTCCACCCATGTCTCTTTGA) and against the *LEU3* gene for standardisation (Chu et al., 2014). qPCR reactions were assembled using QuantiFast SYBR Green PCR Kits (QIAGEN, UK) according to the manufacturer's instructions,

and reactions were run using a two-step amplification protocol. Ct values were determined in the logarithmic amplification range and converted to fold change values as described (Pfaffl, 2001). All reported data were determined using triplicate biological replicates, each assayed in technical duplicate.

Protein extraction and western blotting were conducted as described (von der Haar, 2007), using antibodies from Sigma-Aldrich, UK (anti-firefly luciferase, L0159, anti-HA tag, H6908, and peroxidase-labelled anti-rabbit IgG, A9169). Anti-Hsp104 antibodies were as described (Adam, Jossé, & Tuite, 2017).

Data analysis and statistics. Data were analysed using Python in the Jupyter Notebook environment. All raw data and data analysis scripts used to prepare individual figures are available for download from Github¹. Statistical significance between samples was tested using ANOVA followed by Tukey's Honestly Significant Difference test.

To calculate the numbers of ribosomes required for synthesis of the different luciferase variants, absolute numbers of luciferase molecules per cell were first determined using quantitative western blotting approach as described (von der Haar, 2007) and purified firefly luciferase (Sigma-Aldrich, UK) as standard. The resulting values were 208000, 441000, and 589000 molecules per cell for the min, sta and max constructs respectively (rounded to the nearest thousand). staFLuc protein turnover rate was determined by monitoring loss of the firefly luciferase signal in western blots following treatment of growing cells with cycloheximide, and was found to be 0.008 min⁻¹. This rate was assumed to be equal for the min and max constructs. Transit times for ribosomes over the different FLuc variants were used as published (Chu et al., 2014) and are 308, 141, and 68.6 seconds for min, sta and max, respectively, and the cellular ribosome pool was assumed to be 185000 ribosomes per haploid cell (von der Haar, 2008).

The proportion of ribosomes required in the steady state for synthesis of the different FLuc variants was then calculated as

$$R = N_{FLuc} * (\mu + \tau_{FLuc}) * t_{FLuc} N_{Ribo}$$

where N_{FLuc} is the number of luciferase molecules per cell, N_{Ribo} the number of ribosomes per cell, μ the growth rate, τ_{FLuc} the luciferase turnover rate, and t_{FLuc} the time it takes for a ribosome to transit the luciferase open reading frame.

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¹ <https://github.com/tobiasvonderhaar/codonselection>

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Figure Legends

Figure 1. Expression levels of three codon usage variants of firefly luciferase. Expression levels were assessed either by dual luciferase assay and are shown as the ratio of expression between the variant firefly luciferases and an invariant *Renilla* luciferase gene also contained in the expression vectors (black and grey dots, n = 8), or were assessed by western blotting in which case they are shown as absolute firefly expression levels (not normalised to *Renilla* levels, n = 3).

Figure 2. The expression of different firefly luciferase variants generates distinct fitness defects. A, the steady state copy number of 2 micron plasmids is modulated by the genes they contain. **B,** increasingly optimised codon usage leads to a decrease in steady state plasmid copy numbers as assessed by real time PCR (n=3). **C,** increasingly optimised codon usage in the recombinant luciferase gene leads to a decrease in growth rate (n = 6). In both panels B and C, “vector” refers to a vector expressing only *Renilla* luciferase without a firefly luciferase gene. Statistical significance of results was assessed by ANOVA and Tukeys’ HSD test. Significance to the vector control is indicated as: ns, p>0.05; **, p<0.01; ***, p<0.001.

Figure 3. Fitness defects associated with the expression of codon optimised firefly luciferase are dependent on translation of the gene. A, plasmid copy numbers of a vector containing the codon optimised gene, the same gene with a nonsense mutation in the fifth codon, or the vector control are indicated (n = 3). **B,** growth rate defects of the same strains as in panel A (n = 6). Statistical significance of results was assessed by ANOVA and Tukeys’ HSD test. Significance to the vector control is indicated as: ns, p>0.05; *, p<0.05; **, p<0.01.

Figure 4. Exploring the mechanistic origin of fitness defects associated with the expression of a codon-optimised gene. A, the number of protein molecules per cell expressed from the staFLuc gene was determined by comparing western blot signals of recombinant luciferase with signals derived from controlled numbers of cells. **B,** estimated numbers of ribosomes required at steady state for translation of the different codon variants (see text for explanation). **C,** growth rate defects associated with expression of the codon optimised firefly luciferase gene are not altered in media supplemented with different nutrients (n=4). **D,** sensitivity of yeast strains containing the codon optimised firefly luciferase gene or a vector control to the protein unfolding drug azetidine-2-carboxylic acid (AZC). **E,** expression levels of the molecular chaperone Hsp104 (relative to the loading control Pgk1).

Figure 5. The interplay between required gene expression levels and fitness effects arising from codon optimisation. Growth rates of strains with a chromosomal deletion of the *HIS3* gene but containing codon variants of the same gene on 2 micron plasmids are shown. The plasmid borne-gene is expected to be superfluous in -Ura medium as this contains histidine, essential but required at low expression levels in -Ura/-His medium; and essential and required at high expression levels in the same medium containing 3-aminotriazole (3-AT), a competitive inhibitor of His3 enzymatic activity. Statistical significance of results was assessed by ANOVA and Tukeys’ HSD test. n>=3 for all conditions. Significance to the vector control (top panel) or to the minHIS3 strains (middle and bottom panels) is indicated as: ns, p>0.05; ***, p<0.001.

Figure 6. Fitness defects caused by expression of codon optimised firefly luciferase are strain dependent. Top, growth rates of strains containing the maxFLuc vector (grey dots) or a control vector (black dots). Strains are shown in descending order of average growth rate when containing the vector control construct (n = 3). Bottom, average growth rate loss in % for strains containing the firefly luciferase gene compared to the vector control.

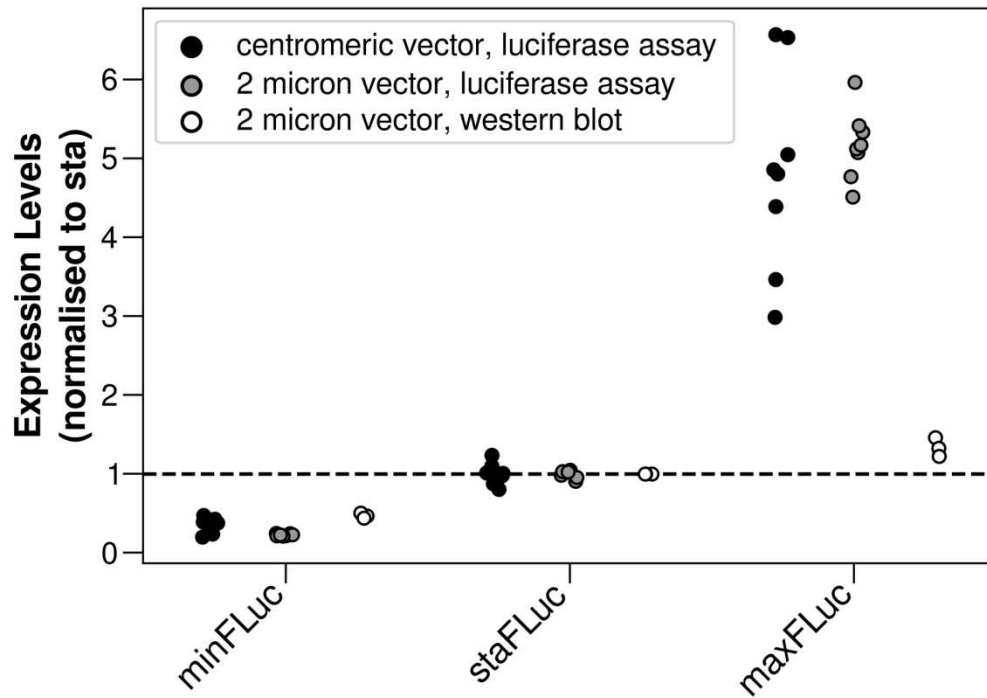


Figure 1. Expression levels of three codon usage variants of firefly luciferase. Expression levels were assessed either by dual luciferase assay and are shown as the ratio of expression between the variant firefly luciferases and an invariant Renilla luciferase gene also contained in the expression vectors (black and grey dots, n = 8), or were assessed by western blotting in which case they are shown as absolute firefly expression levels (not normalised to Renilla levels, n = 3).

79x56mm (600 x 600 DPI)

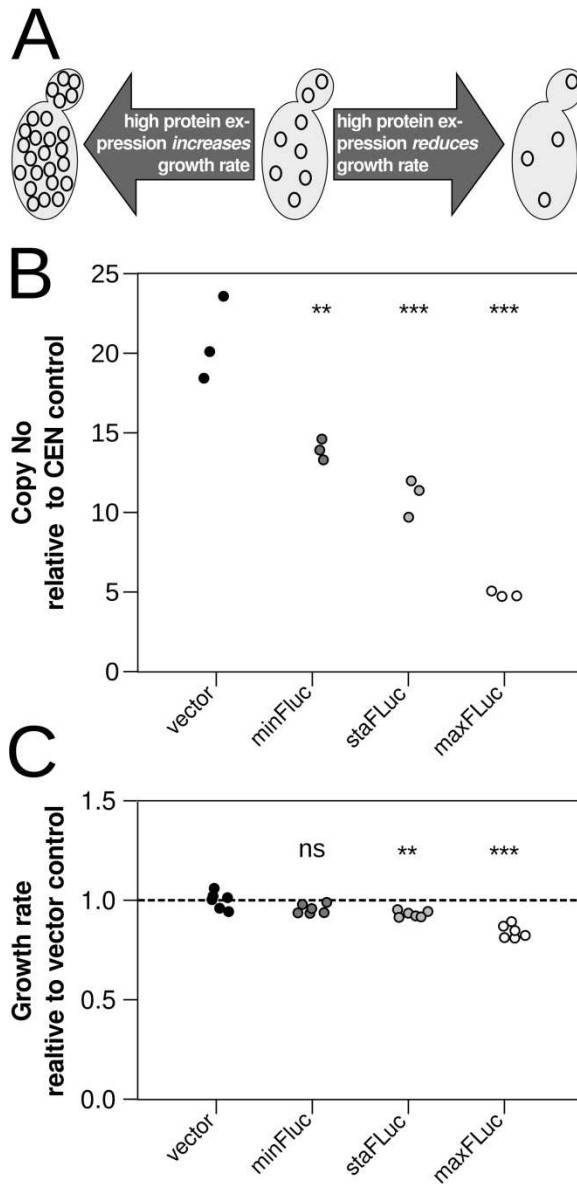


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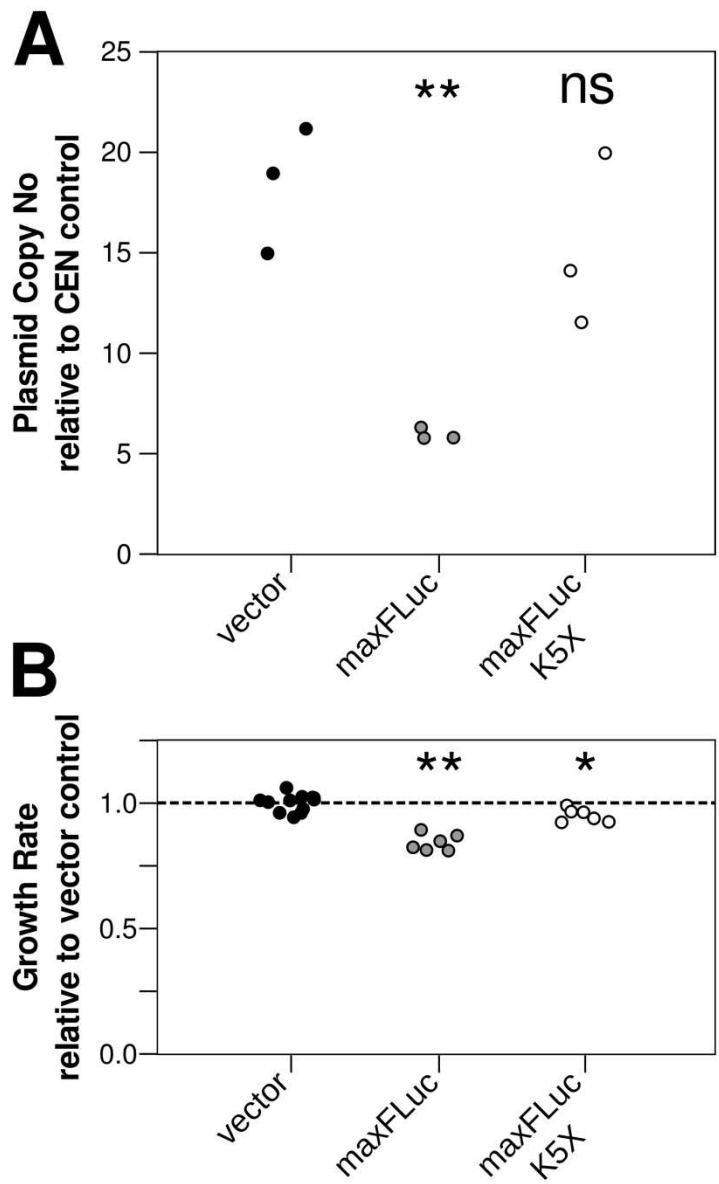
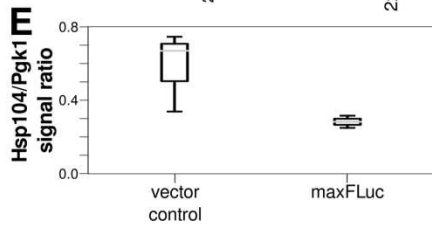
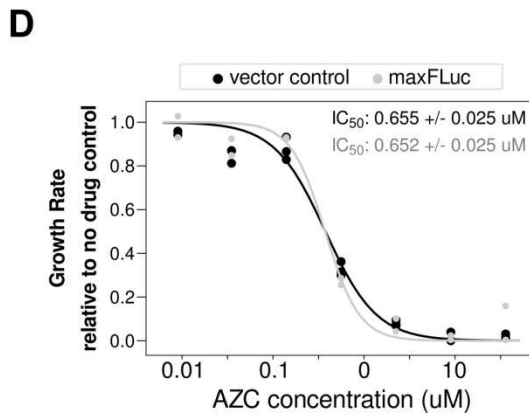
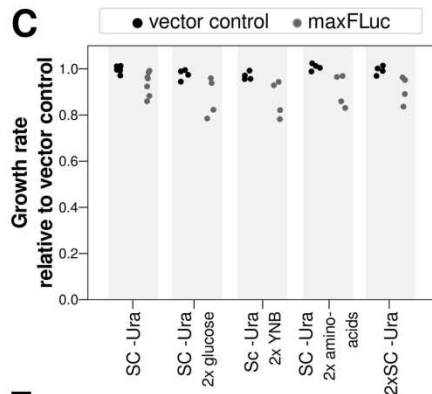
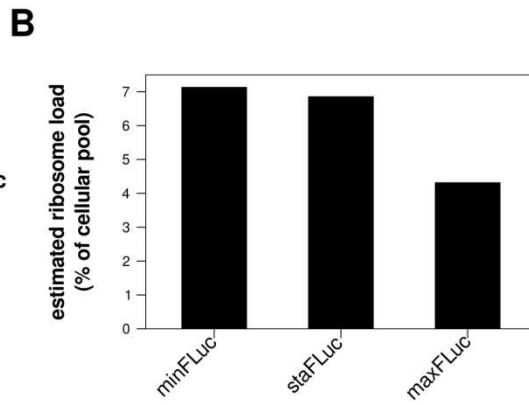
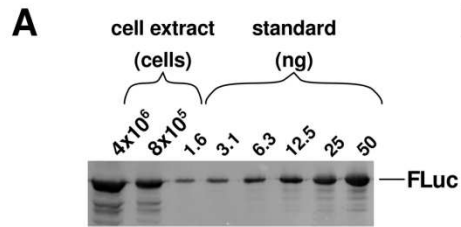


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178x152mm (300 x 300 DPI)

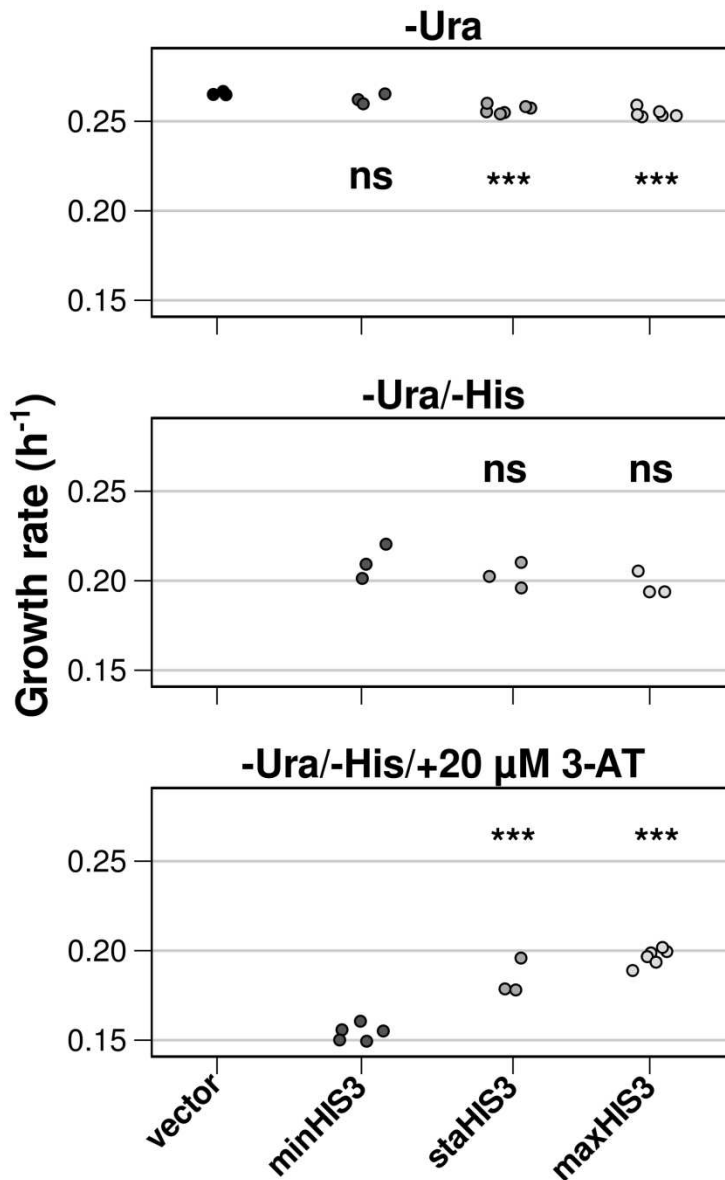


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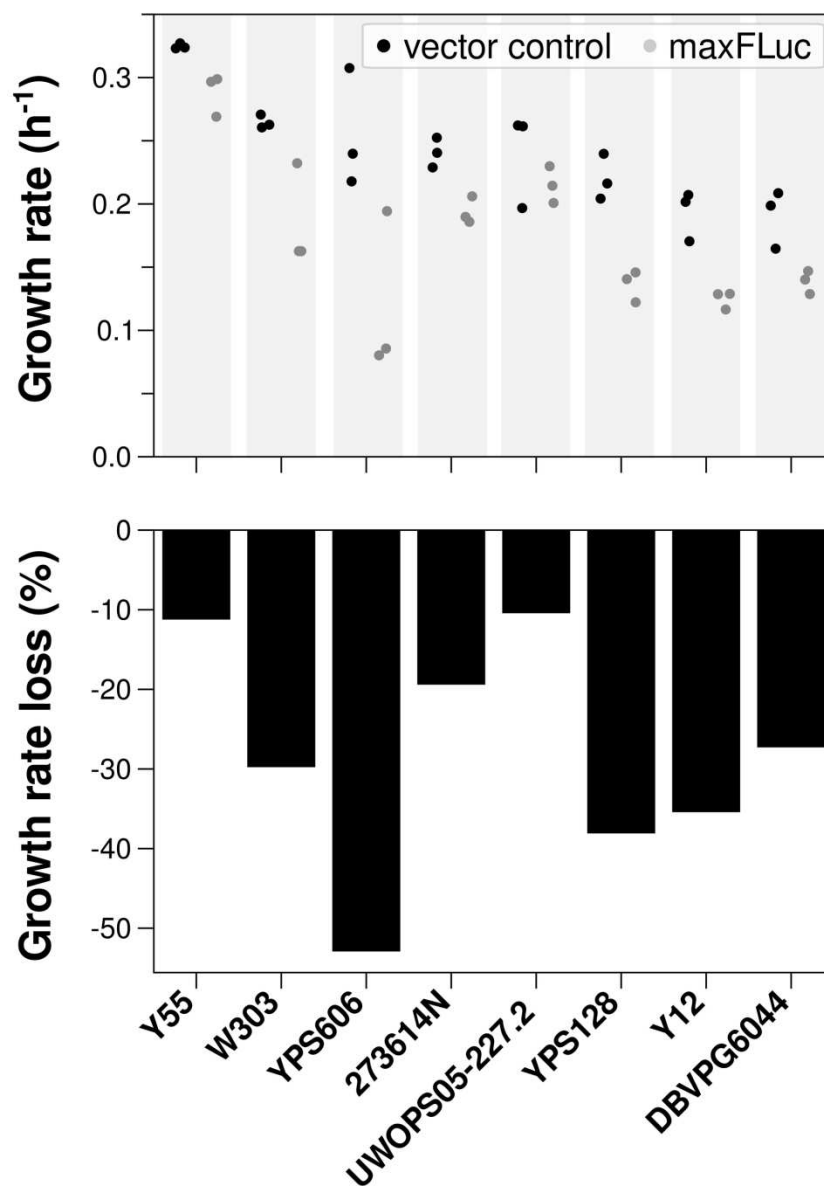


Figure 6. Fitness defects caused by expression of codon optimized firefly luciferase are strain dependent. Top, growth rates of strains containing the maxFLuc vector (grey dots) or a control vector (black dots). Strains are shown in descending order of average growth rate when containing the vector control construct (n = 3). Bottom, average growth rate loss in % for strains containing the firefly luciferase gene compared to the vector control.

79x113mm (600 x 600 DPI)

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3 **Table 1. Plasmids used in this study.**
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Plasmid	Description	Addgene ID	Reference
pBEVY-U	2 micron <i>URA3</i> vector containing <i>TDH3</i> and <i>ADH1</i> -based promoter sequences	51230	(Miller et al., 1998)
pTH646	pBEVY-U expressing <i>Renilla</i> luciferase	40600	this study
pTH731	pTH646 also expressing minCFLuc	40601	this study
pTH732	pTH646 also expressing staCFLuc	40607	this study
pTH733	pTH646 also expressing maxCFLuc	40608	this study
pTH734	pTH646 also expressing maxCFLuc (K5X)	40609	this study
pTH719	pBEVY-U expressing min <i>HIS3</i>	40610	this study
pTH720	pBEVY-U expressing sta <i>HIS3</i>	40611	this study
pTH721	pBEVY-U expressing max <i>HIS3</i>	40612	this study