

Aneuploidy can be an evolutionary diversion on the path to adaptation

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January 28, 2024

Classifications. Biological Sciences: Evolution, Genetics, Microbiology, Population Biology.

Keywords: whole-chromosome duplication, evolutionary model, adaptive evolution

Abstract

Aneuploidy is common in eukaryotes, often leading to decreased fitness. However, evidence from fungi and human tumour cells suggests that specific aneuploidies can be beneficial under stressful conditions and facilitate adaptation. In a previous evolutionary experiment with yeast, populations evolving under heat stress became aneuploid, only to later revert to euploidy after beneficial mutations accumulated. It was therefore suggested that aneuploidy is a “stepping stone” on the path to adaptation. Here, we test this hypothesis. We use Bayesian inference to fit an evolutionary model with both aneuploidy and mutation the experimental results. We then predict the genotype frequency dynamics during the experiment, demonstrating that most of the evolved euploid population likely did not descend from aneuploid cells, but rather from the euploid wild-type population. Our model shows how the beneficial mutation supply—the product of population size and beneficial mutation rate—determines the evolutionary dynamics: with low supply, much of the evolved population descends from aneuploid cells; but with high supply, beneficial mutations are generated fast enough to outcompete aneuploidy due to its inherent fitness cost. Our results suggest that despite its potential fitness benefits under stress, aneuploidy can be an evolutionary “diversion” rather than a “stepping stone”: it can delay, rather than facilitate, the adaptation of the population, and cells that become aneuploid may leave less descendants compared to cells that remain diploid.

34 Introduction

Aneuploidy is an imbalance in the number of chromosomes in the cell: an incorrect karyotype.

36 Evidence suggests aneuploidy is very common in eukaryotes, e.g. animals^{50,39,3}, and fungi^{42,75,46,64}. Aneuploidy has been implicated in cancer formation, progression, and drug resistance^{6,52,50,49,24,35}.

38 It is also common in protozoan pathogens of the *Leishmania* genus, a major global health concern³⁷, and contributes to the emergence of drug resistance⁵³ and virulence³⁸ in fungal pathogens, which

40 are under-studied⁴⁸, despite infecting a billion people per year, causing significant morbidity in >150 million and death in >1.5 million people per year^{53,48}.

42 Experiments with human and mouse embryos found that most germ-line aneuploidies are lethal. Aneuploidies are also associated with developmental defects and lethality in other multicellular organisms⁵⁶. For example, aneuploid mouse embryonic cells grow slower than euploid cells⁶⁹. Similarly, in unicellular eukaryotes growing in benign conditions, aneuploidy usually leads to slower growth and

46 decreased overall fitness, in part due to proteotoxic stress due to increased expression, gene dosage imbalance, and hypo-osmotic-like stress^{40,67,42,56,51,28,74,68,70,47}.

48 However, aneuploidy can be beneficial under stressful conditions due to the wide range of phenotypes it can produce, some of which are advantageous^{42,70}. Indeed, in a survey of 1,011 yeast strains, 50 aneuploidy has been detected in about 19%⁴³. Thus, aneuploidy can lead to rapid adaptation in unicellular eukaryotes^{18,66,22,45}, as well as to rapid growth of somatic tumour cells^{52,58}. For example,

52 aneuploidy in *Saccharomyces cerevisiae* facilitates adaptation to a variety of stressful conditions like heat and pH⁷², copper^{9,18}, salt^{12,47}, and nutrient limitation^{13,20,1}, with similar results in *Candida*

54 *albicans*⁷⁰. Importantly, aneuploidy can also lead to drug resistance in pathogenic fungi such as *C. albicans*^{55,54,17} and *Cryptococcus neoformans*⁵⁹, which cause candidiasis and meningoencephalitis,

56 respectively. Although we focus here on aneuploidy, a similar phenomena of adaptation via gene duplication or amplification has been observed in yeast³⁴, bacteria⁶², and DNA viruses¹⁴.

58 Yona et al.⁷² demonstrated experimentally the importance of aneuploidy in adaptive evolution. They evolved populations of *S. cerevisiae* under strong heat stress. The populations adapted to the heat stress

60 within 450 generations, and this adaptation was determined to be due a duplication of Chromosome III. Later on, after more than 1,500 generations, the populations reverted back to an euploid state, while

62 remaining adapted to the heat stress. Aneuploidy was therefore suggested to be a transient adaptive solution, because it can rapidly appear and take over the population under stressful conditions, and

64 can then be rapidly lost when the cost of aneuploidy outweighs its benefit—after the stress is removed, or after refined beneficial mutations appear and fixate⁷². Furthermore, it has been suggested that

66 aneuploidy is an evolutionary “stepping stone” that facilitates future adaptation by genetic mutations,
which require more time to evolve^{72,71}.

68 Here, we test the hypothesis that aneuploidy is a an evolutionary “stepping stone” that facilitates
adaptive evolution by genetic mutations Yona et al.⁷¹. We develop an evolutionary genetic model and
70 fit it to the experimental results of Yona et al.⁷² to predict the genotype frequency dynamics in the
experimental populations, thereby estimating the frequency of evolved euploid cells that descended
72 from aneuploid cells. Our results show that although aneuploidy reached high frequencies in the
experimental populations, the majority of cells in the evolved euploid population likely did not
74 descend from aneuploid cells, but rather directly from wild-type euploid cells. These suggests that at
the lineage level, aneuploidy may be an “evolutionary diversion”, rather than a “stepping stone”, on
76 the path to adaptation.

Results

78 In the heat-stress experiment of Yona et al.⁷², four populations of *S. cerevisiae* evolved under 39 °C.
Aneuploidy reached high frequency (>95%) in all four experimental repetitions in the first 450
80 generations. Two of the repetitions, marked *H2* and *H4*, carried no large-scale duplications other
than a Chromosome III trisomy. These two repetitions continued to evolve under the same conditions,
82 wherein aneuploidy was eliminated by generation 1,700 and 2,350 in *H4* and *H2*, respectively.

Evolutionary genetic model. To explore the dynamics during the evolutionary experiments, we
84 developed an evolutionary genetic model, fitted the model to empirical data, and used it to predict the
genotype frequency dynamics, or specifically, the fraction of the evolved euploid population descended
86 from aneuploid cells.

The model includes the effects of natural selection, genetic drift, aneuploidy, and mutation (i.e., other
88 genetic variants), and follows a population of cells characterized by their genotype: euploid wild-
type, $2n$, is the ancestral diploid genotype; euploid mutant, $2n^*$, has a diploid karyotype and a single
90 beneficial mutation; aneuploid wild-type, $2n + 1$, has an extra chromosome due to a chromosome
duplication event; and aneuploid mutant, $2n + 1^*$, has and extra chromosome (like $2n + 1$) and a
92 beneficial mutation (like $2n^*$). Note that ‘mutation’ here refers to point mutations and other genetic
variants unrelated to aneuploidy. Fitness values of the different genotypes are denoted by w_{2n} , w_{2n^*} ,
94 w_{2n+1} , and w_{2n+1^*} , and the rate of mutation and aneuploidy are denoted by μ and δ , respectively. See
Figure 1 for an illustration of the model.

96 We fitted this model to the experimental results⁷²—time for fixation (frequency >95%) and for loss
 98 (frequency <5%) of aneuploidy—using approximate Bayesian computation with sequential Monte Carlo
 100 (ABC-SMC)⁶⁰, thereby inferring the model parameters: rates of aneuploidy (i.e., mis-segregation,
 non-disjunction) and mutation and the fitness of all genotypes. We then sampled posterior predictions
 for the genotype frequency dynamics using the estimated parameter values and compared different
 versions of the model to test additional hypotheses about the evolutionary process.

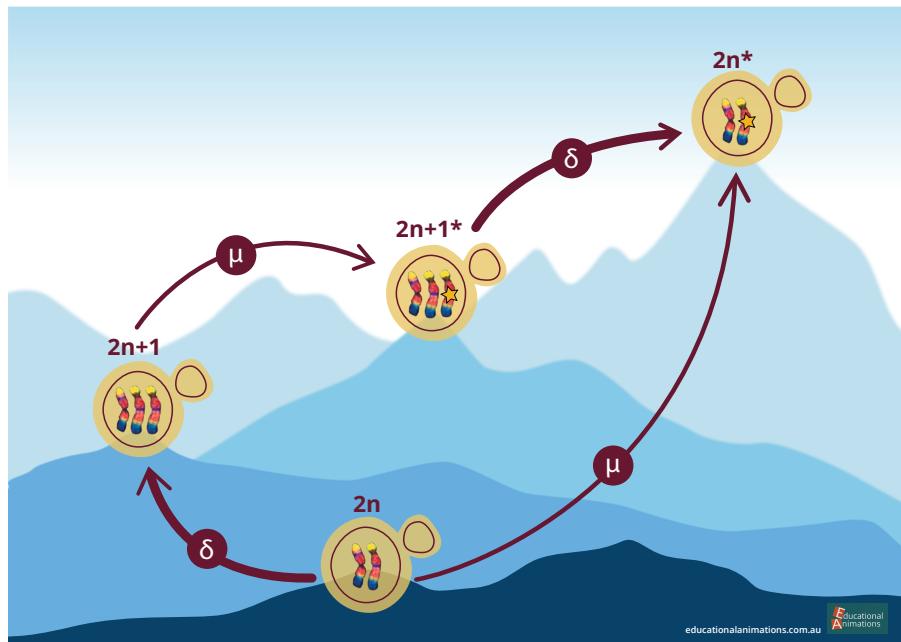


Figure 1: Model Illustration. There are four genotypes in our model: euploid wild-type, $2n$; euploid mutant, $2n^*$; aneuploid wild-type, $2n + 1$; and aneuploid mutant, $2n + 1^*$. Overall there are two possible trajectories from $2n$ to $2n^*$. Arrows denote transitions between genotypes, with transition rates μ for the beneficial mutation rate and δ for the aneuploidy rate. Elevation differences illustrate the expected, rather than the assumed, fitness differences between the genotypes.

102 **Estimated rates and fitness effects of aneuploidy and mutation.** We inferred the posterior distribution
 104 of model parameters (Figure 2). We report parameter estimates using the MAP (maximum a
 posteriori) and providing the 50% HDI (highest density interval) in square brackets. See Supplementary Material for sensitivity analysis.

106 The estimated beneficial mutation rate is $\mu = 2.965 \cdot 10^{-6}$ [$2.718 \cdot 10^{-7} - 3.589 \cdot 10^{-6}$] per genome
 per generation (that is, roughly 3 out of 10^6 cell divisions produce a mutant cell with a fitness
 108 advantage). From the literature, the mutation rate per base pair is roughly $2 - 3 \cdot 10^{-10}$ (refs.^{76,36}), but
 it may be higher under heat stress, as several stresses²¹, including heat²³, may cause hypermutation
 110 in yeast. If we assume a 10-fold increase over the mutation rate reported in the literature, then the
 estimated beneficial mutation rate can be explained by a genomic target size of 1,000 base pairs

112 (that is, 1,000 base pairs across the genome in which a mutation would provide a fitness advantage):
113 $3 \cdot 10^{-10} \times 10 \times 1,000 = 3 \cdot 10^{-6}$. Supporting this, Jarolim et al.²⁶ found 279 genes that contributed
114 to survival after a sudden shift from 30 °C to 50 °C, and Flynn et al.¹⁵ used a deep mutational
115 scan of a single protein, Hsp90, to find 465 amino-acid variants (out of 14,160) that significantly
116 increased growth rate in 37 °C. Furthermore, Yona et al.⁷² found at least 10 genes on Chromosome
117 III that increased heat tolerance when over-expressed. Assuming that other chromosomes also have
118 a similar number of heat-tolerance genes (and even more, as Chromosome III is one of the smallest
119 chromosomes¹⁹), we get a total of 160 heat-tolerance genes in the genome. Indeed, mutations were
120 found in 97 genes in an evolutionary experiment with yeast under heat stress²³. Thus, to get a genomic
121 target size of 1,000, it is enough that the average gene target size (number of base pairs in a gene in
122 which a mutation is beneficial) is 6.25 base pairs. For example, Kohn and Anderson³¹ found a target
size of 11 in a proton exporter gene (*PMA1*) that contributes to high-salt adaptation.

124 The estimated rate of aneuploidy (i.e., mis-segregation, non-disjunction), $\delta = 1.72 \cdot 10^{-3}$ [$1.47 \cdot 10^{-3} -$
125 $2.786 \cdot 10^{-3}$] is higher than in previous studies: for Chromosome III in diploid *S. cerevisiae*, Zhu
126 et al.⁷⁶ estimated $6.7 \cdot 10^{-6}$ chromosome gain events per generation, and Kumaran et al.³³ estimate
127 $3.0 \cdot 10^{-5} - 4.3 \cdot 10^{-5}$ chromosome loss events per generation (95% confidence interval). However,
128 this difference may be partly explained by an increased aneuploidy rate during heat stress: heat shock
can increase the rate of chromosome fragment loss by 2-3 orders of magnitude⁷.

130 The estimated fitness values are $w_{2n+1} = 1.022$ [1.021 – 1.023], $w_{2n+1^*} = 1.025$ [1.024 – 1.026],
131 $w_{2n^*} = 1.028$ [1.026 – 1.029], all relative to the fitness of $2n$, which is set to $w_{2n} = 1$. If we allow for
132 transitions (mutation, chromosome loss and gain) to less-fit genotypes (e.g., $2n^*$ to $2n + 1^*$), then we
infer similar but slightly different values, see Supplementary Material.

134 **Model comparison and goodness-of-fit.** To assess the fit of our model to the data, we use posterior
135 predictive checks, in which we simulate the frequency dynamics using MAP parameter estimates and
136 compare them to the data. Our model fits the data well: $2n^*$ fixed in 63% of simulations by generation
1,700 and in 100% of simulations by generation 2,350 (Figure 3).

138 However, a model without aneuploidy (where the aneuploidy rate is fixed at zero, $\delta = 0$), fails to
139 explain the experimental observations (Figure 3). The estimated mutation rate without aneuploidy is
140 $\mu = 7.98 \cdot 10^{-9}$ [$7.906 \cdot 10^{-9} - 8.138 \cdot 10^{-9}$], much lower compared to a model with aneuploidy. The
141 fitness of the mutant is also much lower at $w_{2n^*} = 1.013$ [1.012 – 1.013]. This is because, without
142 aneuploidy, a high mutation rate or fitness effect will lead to faster appearance and fixation of $2n^*$ than
in the experimental observations.

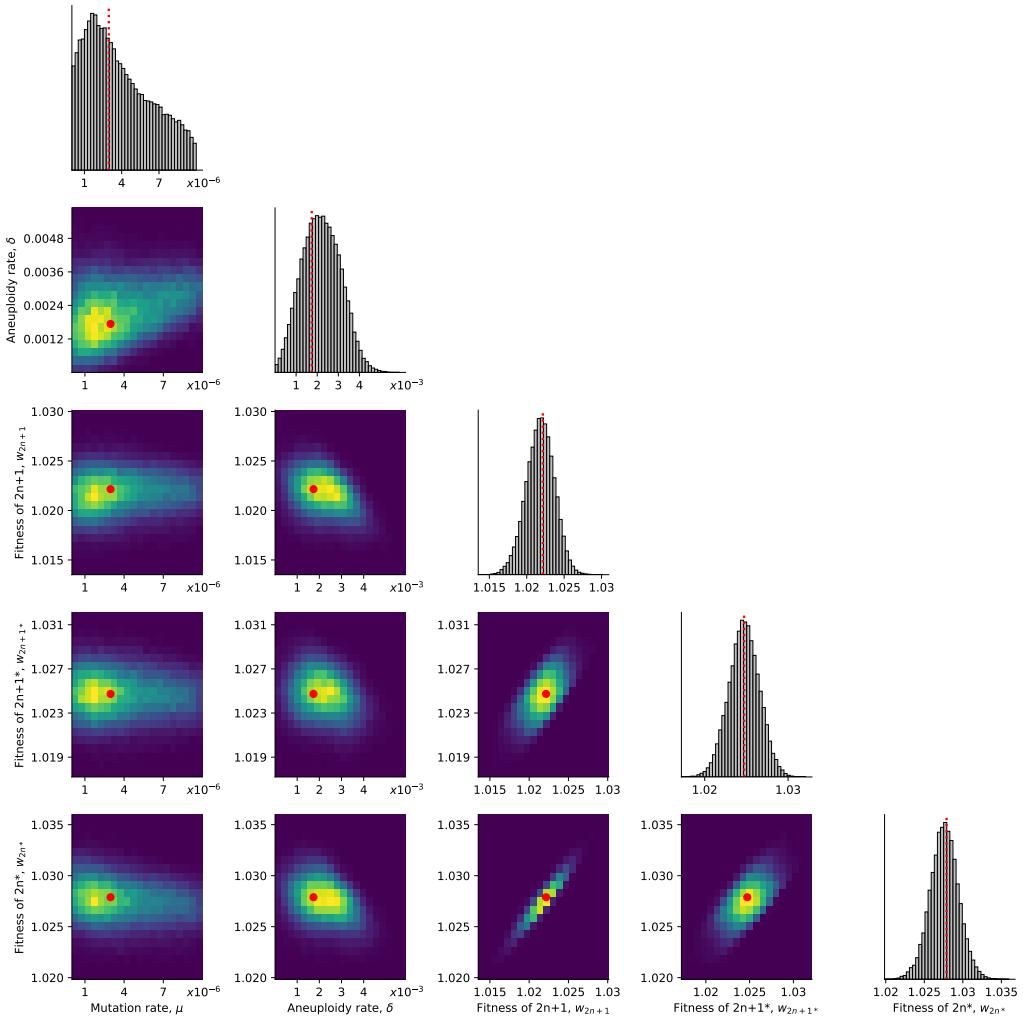


Figure 2: Posterior distribution of model parameters. On the diagonal, the marginal posterior distribution of each model parameter. Below the diagonal, the joint posterior distribution of pairs of model parameters (dark purple and bright yellow for low and high density, respectively). Red markers and orange lines for the joint MAP estimate (which may differ from the marginal MAP, as the marginal distribution integrates over all other parameters).

144 We also checked a model in which aneuploidy occurs but is adaptively neutral compared to the wild-type, that is, $w_{2n+1} = w_{2n}$ and $w_{2n+1}^* = w_{2n}^*$ but $\delta > 0$. This model fits the data better than the model
 146 with no aneuploidy (in which $\delta = 0$), but worse than a model with positive selection for aneuploidy, in which $w_{2n} < w_{2n+1} < w_{2n+1}^* < w_{2n}^*$ (Figure 3).

148 **Model predictions of genotype frequency dynamics.** We simulated 50 replicate genotype frequency dynamics using the MAP estimate parameters. Figure 4A shows the simulated frequencies of
 150 the four genotypes ($2n$, $2n + 1$, $2n + 1^*$ and $2n^*$), as well as the frequencies of $2n^*$ cells that arose from either $2n + 1$ cells via a sequences of mutation and chromosome loss events ($2n_A^*$), or directly from
 152 $2n$ cells via a mutation event ($2n_M^*$). We find that $2n + 1^*$ never reaches substantial frequency as it is

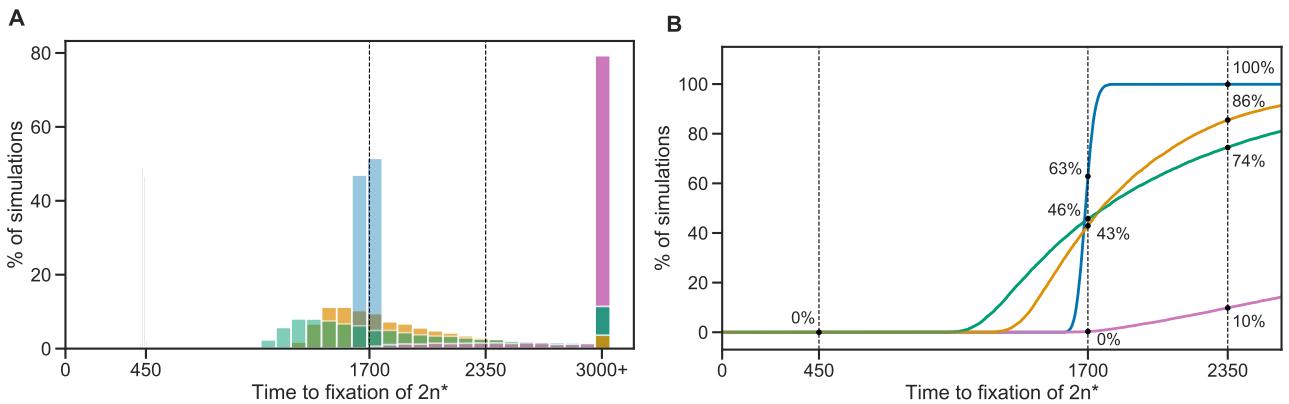


Figure 3: Model fit with and without aneuploidy. The distribution of time to fixation of $2n^*$ (i.e., adaptation time) in 10,000 simulations using MAP parameters of the model with beneficial aneuploidy (blue; $\delta > 0$, $w_{2n} < w_{2n+1} < w_{2n+1^*} < w_{2n^*}$) compared to alternative models: a model with the same parameter values but without aneuploidy (gray, $\delta = 0$, concentrated at $t = 450$); a model fitted to the data assuming no aneuploidy (green, $\delta = 0$); a model fitted to the data assuming neutral aneuploidy (yellow, $\delta > 0$, $w_{2n+1} = w_{2n}$, $w_{2n+1^*} = w_{2n^*}$); and a model with beneficial aneuploidy and an extended prior distribution (pink). In the experiment by Yona et al.⁷², one population lost aneuploidy by generation 1,700 and another by generation 2,350 (dashed lines) but not before generation 450. Thus, the blue distribution has a better fit compared to the other distributions (the gray distribution has a particularly poor fit). The MAP likelihood (eq. (4)) is 0.84, 0.78, 0.67, and 0.14 for the models represented by blue, yellow, green, and pink distributions, respectively. **(A)** Histogram of the time to fixation of $2n^*$. The last bin contains all values equal or greater than 3,000. **(B)** Cumulative distribution of the time to fixation.

quickly replaced by $2n^*$ in a process similar to *stochastic tunneling*^{25,32}.

- 154 To test the hypothesis that aneuploidy facilitates adaptation, we estimated F_A , the expected frequency
 155 of $2n^*$ that arose from $2n + 1$, computed as the average frequency of such $2n_A^*$ cells at the end of
 156 simulations using the MAP estimate parameters. Surprisingly, we observe that the majority of $2n^*$
 157 cells are $2n_M^*$, a product of a direct mutation in $2n$ cells, rather than descending from $2n + 1$ cells
 158 ($F_A^{MAP} = 0.106$, average end point of 50 purple lines in Figure 4A). This is despite the fact that the
 159 $2n + 1$ genotype reaches high frequencies in the population (at least 0.98, Figure 4A).
- 160 This result is not unique to the MAP parameter estimate. We simulated genotype frequency dynamics
 161 using parameter samples from the posterior distribution, and computed the posterior distribution of
 162 F_A (Figure 4B). The posterior mode F_A was just 0.147 [0.0154-0.370 95% CI] and only in 489 of
 163 100,000 posterior samples (0.489%) F_A was larger than 0.5 (see Supporting Material for results when
 164 transitions to less-fit genotypes are allowed, such as $2n^*$ to $2n + 1^*$). Thus, if we sample a random

cell from the evolved $2n^*$ population, it is more likely to have descended directly from an euploid cell than from an aneuploid cell. The probability of $2n^*$ descending from $2n + 1$ (F_A) increases with the aneuploidy rate, δ , and decreases with both the population size N and the mutation rate, μ (Figure 4C,D). In some cases it can also be affected by the fitness parameters (Figure S10).

Genetic instability in aneuploid cells. It has been suggested that aneuploidy increases genetic instability: Sheltzer et al.⁵⁷ have demonstrated a fold increase of between 2.2 and 7.1 in the mutation rate of disomic yeast (rather than trisomic yeast, the focus of our analysis). Therefore, we inferred model parameters under the assumption that the mutation rate increases in aneuploid cells by a factor $\tau = 1, 33/32$ (due to an additional chromosome), 2, 5, 10, or 100 (due to genetic instability). We found that the posterior distribution was similar for $\tau = 1, 33/32, 2$, and 5 (Figure S4). Furthermore, we computed the WAIC, a criterion for model selection (Methods). The WAIC values were similar for all τ values (Table S1).

Assuming a strong increase of the mutation rate in aneuploid cells, i.e. $\tau = 100$, the inferred a mutation rate was $\mu = 4.094 \cdot 10^{-7}$ [$6.252 \cdot 10^{-8} - 6.046 \cdot 10^{-7}$]), and the inferred aneuploidy rate that was $\delta = 0.744 \cdot 10^{-3}$ [$0.506 \cdot 10^{-3} - 1.827 \cdot 10^{-3}$]. Compared to inference made assuming no effect of aneuploidy on the mutation rate, these rates were about 7-8-fold and 2-3-fold lower for μ and δ , respectively. Assuming $\tau = 10$, the inferred a mutation rate was only slightly lower compared to $\tau = 1$ ($\mu = 1.67 \cdot 10^{-6}$ [$2.836 \cdot 10^{-8} - 2.245 \cdot 10^{-6}$]).

Therefore, we do not find evidence of an increase in mutation rate in aneuploid cells. This may be because, unless the increase is strong ($\tau \geq 10$), it does not seem to affect our inference; or because Chromosome III is one of the smallest chromosomes¹⁹. We also checked the differences in genotype frequency dynamics for different τ values. We observe $\tau = 100$ could be distinguished if accurate data was available for the waiting time until the frequency of $2n$ to decrease below 95% (Figure S5A) or for waiting time for the frequency of $2n + 1$ to either reach or go below 95% (Figure S5B).

Similarly, we did not find evidence for an increase in the rate of chromosome loss in aneuploid cells⁵⁷, probably due to lack of statistical power. Nevertheless, increasing the rate of chromosome loss (transitions from $2n + 1^*$ to $2n^*$) without increasing the rate of chromosome gain (transitions from $2n$ to $2n + 1$) increases F_A (Figure S11B), but not to the same extent as increasing the rate of chromosome gain. In contrast, increasing the mutation rate in aneuploid cells can have a marked effect on the dynamics: when using the MAP parameter estimates, F_A increases to 0.52 when the mutation rate in aneuploid cells increases 10-fold (Figure S11C).

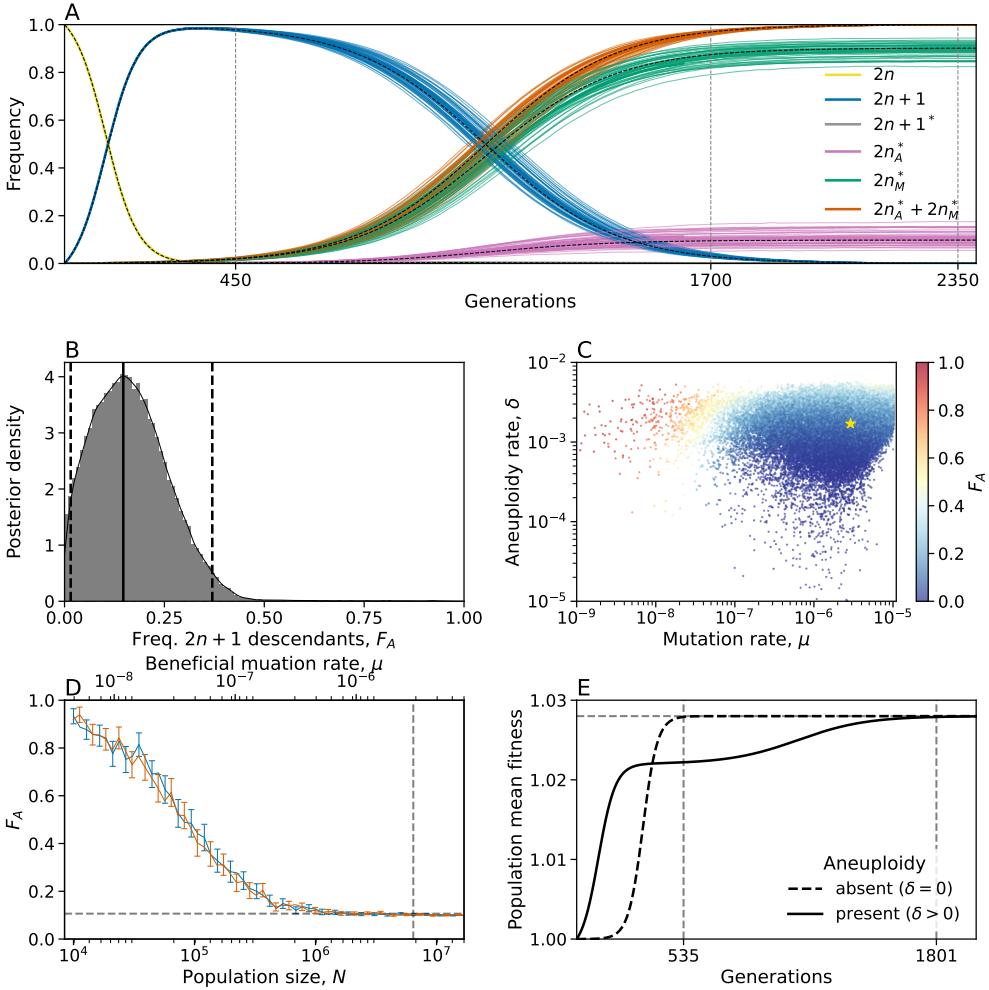


Figure 4: Predicted frequency of aneuploid-descended cells. (A) Posterior predicted genotype frequencies over time, including the source of $2n^*$: $2n_A^*$ arose from $2n+1$, whereas $2n_M^*$ arose directly from $2n$. Colored curves are 50 simulations using the MAP estimate parameters. Black dashed curves are the expected genotype frequencies without genetic drift (from a deterministic model). See Figure S9 for log-log scale, in which the sequence of events is easier to observe. (B) Posterior distribution of F_A , the expected frequency of $2n^*$ cells descended from $2n+1$ cells, computed as the average frequency at the end of 100 simulations for 100,000 samples from the parameter posterior distribution. Solid and dashed lines show the mode and 95% CI. (C) F_A values (color coded) from panel B, with their corresponding mutation rate μ on x-axis and aneuploidy rate δ on the y-axis. Yellow star shows the MAP estimate. See also Figure S10. (D) F_A as a function of the population size (N , bottom x-axis) and the beneficial mutation rate (μ , top x-axis) in posterior predictions with MAP parameters. Markers show F_A in 250 simulations per population size or mutation rate value. Error bars show mean F_A with 95% CI (bootstrap, $n = 10,000$). Blue and red bars for varying population size and mutation rate, respectively. Vertical dashed line for population size in the experiment, $6.425 \cdot 10^6$, and the MAP mutation rate, $2.965 \cdot 10^{-6}$. Horizontal line for $F_A^{MAP} = 0.106$. (E) Population mean fitness in a model without drift using MAP estimate parameters. Solid lines for mean fitness with aneuploidy ($\delta > 0$), where the population reaches adaptation (mean fitness at 99.99% of maximum value) at generation 1,802. Dashed lines for mean fitness without aneuploidy ($\delta = 0$), where the population adapts much earlier, at generation 535.

196 **Discussion**

In a study on the role of chromosome duplication in adaptive evolution, Yona et al.⁷² found that a
198 Chromosome III trisomy was acquired by *S. cerevisiae* populations evolving under heat stress, only
to be later replaced by euploid mutant cells that carry "refined" solutions to the stress. Additionally,
200 such a replacement also occurred when they initiated evolutionary experiments with a population in
which all cells carry a Chromosome III trisomy. They hypothesized that aneuploidy is a "useful yet
202 short-lived intermediate that facilitates further adaptation", suggesting that the euploid mutant cells
evolved by heat-resistance mutations in aneuploid cells followed by reversion of trisomy due to a
204 chromosome loss event.

We developed an evolutionary genetic model of adaptive evolution by aneuploidy and mutation
206 (Figure 1), fitted it to the experimental results of Yona et al.⁷², and used it to predict the genotype
frequency dynamics. The model predicted that only about 10-15% of the evolved euploid population
208 descended from aneuploid cells by acquiring a mutation and losing the extra chromosome—that is,
the majority of the euploid population are not descended from aneuploid cells, but rather are direct
210 descendants of the ancestral wild-type population (Figure 4).

This happens despite aneuploidy reaching a high frequency in the population (>95%). Conventional
212 wisdom might suggest that once the aneuploid genotype $2n + 1$ reaches high frequency, it will have a
better chance at producing "refined" solutions via mutations, and its descendants will come to dominate
214 the population: the frequency of $2n_A^*$ (which arises from $2n + 1^*$) will be higher than the frequency of
 $2n_M^*$ (which arises directly from $2n$).

216 So how does $2n_M^*$ prevail? Initially, the supply rates of $2n + 1$ and $2n_M^*$ are $N\delta \approx 11,000$ and $N\mu \approx 19$,
respectively (assuming MAP parameter estimates). Therefore, both genotypes are expected to appear
218 immediately at the beginning of the experiment (Figure S9). However, $2n + 1$ appears at a much
higher frequency as $\delta \gg \mu$ by 2-3 orders of magnitude. After they first appear, $2n_M^*$ has higher
220 fitness. But as long as the frequency of $2n$ is high, the supply rate of $2n + 1$ is higher than that of
 $2n_M^*$, again due to $\delta \gg \mu$. However, supply rates of both genotypes decrease with the frequency
222 of $2n$. Therefore, when the latter decreases, mainly due to the increase in the frequency of $2n + 1$,
both supply rates diminish. At this stage, the higher fitness of $2n_M^*$ comes into play and it starts to
224 take over the population, which is mainly composed of $2n + 1$. For the aneuploid lineage to compete
with the mutant lineage, it must produce $2n_A^*$ via a mutation followed by chromosome loss. Although
226 this is a stochastic process (due to drift), our results show that the time until $2n_A^*$ reaches a frequency
of 0.1% is roughly 450 generations, without much variation (intersection of purple lines and vertical

228 dashed line in Figure S9). However, by that time $2n_M^*$ is already at a roughly 10-fold higher frequency
(1.86%), and since both mutants have the same fitness, their relative frequency remains roughly the
230 same until the end of the experiment.

232 **Predictions for small populations and low mutation rates.** We examined the effect of the popula-
234 tion size, N , and the beneficial mutation rate, μ , on the frequency of $2n + 1$ descendants in the evolved
236 population, F_A . We found that F_A is expected to decrease as the population size or mutation rate
238 increase (Figure 4D), ranging from >90% when the population size is 10,000 or the mutation rate is
240 $6 \cdot 10^{-9}$, to about 10% when the population size is above 1,000,000 (less than the experimental popu-
lation size, which was 6,425,000) or the mutation rate is above $2 \cdot 10^{-6}$ (less than the inferred mutation
rate, which is $2.965 \cdot 10^{-6}$). Thus, our model provides a testable prediction: if the experiment was
repeated under a lower population size (via stronger daily dilutions or in a smaller volume) or a lower
mutation rate (via a non-mutagenic stress or stress with a smaller target size such as drug resistance),
then the fraction of the population descending from aneuploid cells would be much higher.

242 **Aneuploidy delays rather than facilitates adaptation.** An additional interesting result of our
244 study is that aneuploidy increases, rather than decreases, the adaptation time (Figure 4E). This
happens despite the fact that the mean fitness initially increases faster in the presence of aneuploidy
246 (Figure 4E). This is because once $2n + 1$ is common, selection for the mutant strain ($2n + 1^*$ or $2n^*$) is
weaker compared to when $2n^*$ competes directly with $2n$.

246 **Rate and fitness effect of aneuploidy and mutation.** We inferred the rates of aneuploidy and
248 mutation and their effects on fitness. We estimate that the aneuploidy rate (i.e., number of chromosome
gains per generation) is $1.7 \cdot 10^{-3}$, higher than a previous estimate of $6.7 \cdot 10^{-6}$ (ref. 75). This may
250 be due to genetic instability caused by heat stress⁷, but we note that there is a general scarcity of
empirical data on aneuploidy rates. In addition, we did not find evidence for increased mutation rates
252 in aneuploid cells. Previous empirical studies have suggested that genetic instability (e.g., elevated
mutation rates) in aneuploid cells is due to stress associated with the aneuploid state^{5,8,73,24}. However,
254 in the experiment of Yona et al.⁷², both the wild-type and the aneuploid were under heat stress, which
may explain why we didn't find evidence for an increased mutation rate specifically in aneuploid
cells.

256 **Effect of ploidy.** The evolutionary dynamics may change in haploid yeast, in which aneuploidy
258 results in a second, rather than third, chromosome copy. For example, it has been demonstrated that
drug resistance mainly evolves via recessive mutations and aneuploidy in haploid yeast⁶¹, whereas in

diploid yeast it evolves via dominant mutations, aneuploidy, and gene/segmental duplications⁴. Thus,
260 the second chromosome copy of disomic yeast may facilitate further adaptation via duplications,
rearrangements, and increased mutational tolerance², while decreasing the chance for adaptation via
262 recessive mutations. Future models and experiments can consider how ploidy and other genomic
contexts affect the role of aneuploidy in adaptive evolution.

264 **Conclusions.** Here, we tested the hypothesis that aneuploidy cells are an evolutionary “stepping
stone”, or adaptive intermediate, between wild-type euploid cells and mutant euploid cells⁷¹. Our
266 results suggest that, although it seems the population goes from euploid to aneuploid and back, this is
not the case at the individual level. We estimate that only about 10-15% of the euploid cells descended
268 from aneuploid cells, whereas the rest are direct descendants of the wild-type euploid cells. Thus,
aneuploidy can delay, rather than accelerate, adaptation, and cells that become aneuploid may leave
270 less descendants than cells that remain euploid. This surprising result reinforces the importance of
mathematical models when interpreting evolutionary dynamics. Moreover, our study emphasizes the
272 unintuitive outcomes of clonal interference between mechanisms for generation of variation that differ
in their rate of formation and distribution of fitness effects, including mutation, copy number variation,
274 horizontal gene transfer, and epigenetic modifications.

Models and Methods

276 **Evolutionary genetic model.** We model the evolution of a population of cells using a Wright-Fisher
model⁴¹, assuming a constant effective population size N , non-overlapping generations, and including
278 the effects of natural selection, genetic drift, aneuploidy, and mutation. We focus on beneficial ge-
netic modifications, neglecting the effects of deleterious and neutral mutations or karyotypic changes.
280 The model allows for a single aneuploid karyotype (e.g., Chromosome III duplication) and a single
mutation to accumulate in the genotype. Thus, the model follows four genotypes (Figure 1): euploid
282 wild-type, $2n$, the initial genotype; euploid mutant, $2n^*$, with the standard karyotype and a single
beneficial mutation; aneuploid wild-type, $2n + 1$, with an extra chromosome, i.e., following chro-
284 mosome duplication; and aneuploid mutant, $2n + 1^*$, with an extra chromosome and a beneficial
mutation.

286 Transitions between the genotypes occur as follows (Figure 1): Beneficial mutations from $2n$ to $2n^*$
and from $2n+1$ to $2n+1^*$ occur with probability μ , the mutation rate. We neglect back-mutations (i.e.,
288 from $2n^*$ to $2n$ and from $2n + 1^*$ to $2n + 1$). Aneuploidy is formed by chromosome mis-segregation,
so that cells transition from $2n$ to $2n + 1$ and from $2n + 1^*$ to $2n^*$ with probability δ , the aneuploidy

rate. That is, we assume chromosomes are gained and lost at the same rate, and we neglect events that form a less-fit genotype (i.e., $2n + 1$ to $2n$ and $2n^*$ to $2n + 1^*$). A model that assumes an increased rate of chromosome loss in aneuploid cells (as in Sheltzer et al.⁵⁷) did not perform well, probably due to lack of statistical power, and was abandoned.

In the experiment by Yona et al.⁷², the population was grown every day from $1.6 \cdot 10^6$ cells until reaching stationary phase and then diluted 1:120. Thus, we set the population size to $N = 6.425 \cdot 10^6$, the harmonic mean of $\{2^k \cdot 1.6 \cdot 10^6\}_{k=0}^7$ ¹¹. The initial population has N cells with genotype $2n$. The effect of natural selection on the frequency f_i of genotype $i = 2n, 2n + 1, 2n + 1^*$, or $2n^*$ is given by

$$f_i^s = \frac{f_i w_i}{\bar{w}}, \quad (1)$$

where w_i is the fitness of genotype i and $\bar{w} = \sum_j f_j w_j$ is the population mean fitness. The effect of mutation and aneuploidy on genotype frequencies is given by

$$\begin{aligned} f_{2n}^m &= (1 - \delta - \mu) f_{2n}^s, \\ f_{2n+1}^m &= \delta f_{2n}^s + (1 - \mu) f_{2n+1}^s, \\ f_{2n+1^*}^m &= \mu f_{2n+1}^s + (1 - \delta) f_{2n+1^*}^s, \\ f_{2n^*}^m &= \mu f_{2n}^s + \delta f_{2n+1^*}^s + f_{2n^*}^s. \end{aligned} \quad (2)$$

Finally, random genetic drift is modeled using a multinomial distribution⁴¹,

$$\mathbf{f}' \sim \frac{1}{N} \cdot \text{Mult}(N, \mathbf{f}^m), \quad (3)$$

where $\mathbf{f}^m = (f_{2n}^m, f_{2n+1}^m, f_{2n+1^*}^m, f_{2n^*}^m)$ are the frequencies of the genotypes after mutation and aneuploidy, \mathbf{f}' are the genotype frequencies in the next generation, and $\text{Mult}(N, \mathbf{f})$ is a multinomial distribution parameterized by the population size N and the genotype frequencies \mathbf{f} . Overall, the change in genotype frequencies from one generation to the next is given by the transformation $f_i \rightarrow f'_i$.

Empirical data for model inference. We use the results of evolutionary experiments reported by Yona et al.⁷². In their heat-stress experiment, four populations of *S. cerevisiae* evolved under 39 °C. Aneuploidy fixed (frequency >95%) in all four population in the first 450 generations. Hereafter, fixation or elimination of a genotype by generation t means that more than 95% or less than 5% of the population carry the genotype at generation t , and possibly earlier. In the original analysis of Yona et al.⁷², samples were routinely extracted from the evolving populations and tested for indication of heat-shock tolerance. The first generation in which such indication was found was generation 200. Therefore, we determine that aneuploidy did not reach high frequency before generation 200. The

experiment continued with two populations, in which aneuploidy was eliminated by generation 1,700
 318 and 2,350 while still under the same conditions of elevated heat (39 °C).

Likelihood function. Because our model, just like the Wright-Fisher model, is non-linear and
 320 stochastic, computing the distribution of fixation time $T(g)$ of genotype g for use in the likelihood
 function is intractable (it is even hard to use a diffusion-equation approximation due to the model having
 322 multiple genotypes, rather than just two). We overcome this problem by approximating the likelihood
 using simulations. We simulate 1,000 experiments per parameter vector $\theta = (\mu, \delta, s, b, c)$, resulting
 324 in a set of simulated observations $\tilde{\mathbf{X}} = \{\tilde{X}_i\}_{i=1}^{1000}$. We then compute the approximate likelihood,

$$\begin{aligned}\mathcal{L}(\theta) = P^4(200 \leq T(2n+1) \leq 450) \cdot & \left[1 - \right. \\ & P_{\tilde{\mathbf{X}}}^4(\{\{T(2n^*) < 1700\} \mid 200 \leq T(2n+1) \leq 450\}) - \\ & P_{\tilde{\mathbf{X}}}^4(\{\{1700 < T(2n^*) < 2350\} \mid 200 \leq T(2n+1) \leq 450\}) + \\ & \left. P_{\tilde{\mathbf{X}}}^4(\{\{T(2n^*) < 1700\} \wedge \{1700 < T(2n^*) < 2350\} \mid 200 \leq T(2n+1) \leq 450\}) \right],\end{aligned}\quad (4)$$

326 where $!\{\dots\}$ is the "logical not" operator, $P^4(\dots)$ is the 4th power of $P(\dots)$, and all probabilities
 $P_{\tilde{\mathbf{X}}}(\dots)$ are approximated from the results of the simulations $\tilde{\mathbf{X}}$. For example, $P_{\tilde{\mathbf{X}}}(\{\{T(2n^*) < 1700\} \mid$
 328 $200 \leq T(2n+1) \leq 450\})$ is approximated by taking simulations in which $2n+1$ fixed (reached >95%)
 before generation 450 but not before generation 200, and computing the fraction of such simulations
 330 in which $2n^*$ did not fix by generation 1,700, and hence aneuploidy did not extinct (reach <5%)
 before generation 1,700. Figure S1 compares results with less and more simulated experiments,
 332 demonstrating that 1,000 simulations are likely sufficient.

For a model without aneuploidy (that is, when the aneuploidy rate is fixed at zero, $\delta = 0$), we disregard
 334 the increased expression in Chromosome III and the growth advantage measured in generation 450,
 and focus on the growth advantage measured in later generations, presumably due to a beneficial
 336 mutation. Therefore, the likelihood is approximated by

$$\begin{aligned}\mathcal{L}_!(\theta) = 1 - & P_{\tilde{\mathbf{X}}}^4(\{\{T(2n^*) < 1700\}\}) - \\ & P_{\tilde{\mathbf{X}}}^4(\{\{1700 < T(2n^*) < 2350\}\}) + \\ & P_{\tilde{\mathbf{X}}}^4(\{\{T(2n^*) < 1700\} \wedge \{1700 < T(2n^*) < 2350\}\}).\end{aligned}\quad (5)$$

338 **Parameter inference.** To infer model parameters, we use approximate Bayesian computation with
 a sequential Monte-Carlo scheme, or ABC-SMC⁶⁰, implemented in the pyABC Python package³⁰
 340 pyabc.readthedocs.io. This approach uses numerical stochastic simulations of the model to infer a
 posterior distribution over the model parameters. It is a method of likelihood-free, simulation-based

342 inference¹⁰, that is, for estimating a posterior distribution when a likelihood function cannot be
343 directly computed. It is therefore suitable in our case, in which the likelihood function can only be
344 approximated from simulations, and cannot be directly computed.

The ABC-SMC algorithm employs sequential importance sampling over multiple iterations^{65,29,63}. In
346 iteration t of the algorithm, a set of parameter vectors, $\{\theta_{i,t}\}_{i=1}^{n_t}$, also called *particles*, are constructed
in the following way. A proposal particle, θ^* , is sampled from a proposal distribution, and is either
348 accepted or rejected, until n_t particles are accepted. The number of particles, n_t , is adapted at every
iteration t using the adaptive population strategy³⁰ pyabc.readthedocs.io. For $t = 0$, the proposal
350 particle is sampled from the prior distribution, $p(\theta)$. For $t > 0$, the proposal particle is sampled from
the particles accepted in the previous iteration, $\{\theta_{i,t-1}\}_{i=1}^{n_{t-1}}$, each with a probability relative to its weight
352 $W_{t-1}(\theta_{i,t-1})$ (see below). The proposal particle is then perturbed using a kernel perturbation kernel,
 $K_t(\theta^* | \theta)$ where θ is the sample from the previous iteration. Then, a set of synthetic observations
354 $\tilde{\mathbf{X}}^*$ is simulated, and the proposal particle θ^* is accepted if its approximate likelihood (eq. (4)) is high
enough, $\mathcal{L}(\theta^*) > 1 - \epsilon_t$ (or more commonly, if $1 - \mathcal{L}(\theta^*) < \epsilon_t$), where $\epsilon_t > 0$ is the *acceptance*
356 *threshold*, as higher values of ϵ_t allow more particles to be accepted. The acceptance threshold ϵ_t is
chosen as the median of the $1 - \mathcal{L}(\theta)$ of the particles accepted in the previous iteration, $t - 1$, and
358 $\epsilon_0 = 0.01$. For each accepted particle $\theta_{i,t}$ a weight $W_t(\theta_{i,t})$ is assigned: for $t = 0$, $W_0(\theta_{i,0}) = 1$,
and for $t > 0$, $W_t(\theta_{i,t}) = p(\theta_{i,t}) / \sum_{i=1}^{n_{t-1}} W_{t-1}(\theta_{i,t-1}) K_t(\theta_{i,t}, \theta_{i,t-1})$, where $p(\theta)$ is the prior density of
360 θ and $K_t(\theta', \theta)$ is the probability of a perturbation from θ to θ' . $K_t(\theta' | \theta)$ is a multivariate normal
distribution, fitted at iteration t to the particles from the previous iteration, $\{\theta_{i,t-1}\}_{i=1}^{n_{t-1}}$, and their
362 weights, $\{W(\theta_{i,t-1})\}_{i=1}^{n_{t-1}}$.

Acceptance is determined according to the approximate likelihood (eq. (4)), which has a maximum
364 value of $\mathcal{L}_{max} = 0.875$ (giving a minimal value of $\epsilon_{min} = 0.125$). We terminated the inference
iterations when the change in ϵ value from one iteration to the next was small. With our standard prior
366 and model, we reached $\epsilon = 0.13$ (or $\mathcal{L} = 0.87$) after six iterations, with $n_6 = 982$ accepted parameter
vectors and effective sample size ESS=651 (Figure S2). Running the inference algorithm with different
368 initialization seeds and less or more simulations for approximating the likelihood produced similar
posterior distributions (Figure S1).

370 After producing a set of weighted particles from the the posterior distribution using the above ABC-
371 SMC algorithm, we approximate the posterior using kernel density estimation (KDE) with Gaussian
372 kernels. We truncate the estimated posterior to avoid positive posterior density for values with zero
prior density. The MAP (maximum a posteriori) estimate is computed as the the maximum of the
374 estimated joint posterior density. We then draw 5,000,000 samples from the posterior distribution

to compute the HDI (highest density interval) and draw 50,000 samples to visualize the posterior
376 distribution with histograms.

Model comparison. We examine several versions of our evolutionary models, e.g. without aneuploidy or with increased mutation rate in aneuploid cells, as well as several different prior distributions (see below). To compare these, we plot posterior predictions: for each model we execute 10,000
378 simulations using the MAP parameter estimates and plot the distributions of time to fixation of $2n^*$,
380 one of key properties of the model likelihood. These plots visualize the fit of each model to the
382 data. Also, for similar models we plot the marginal and joint posterior distributions of the parameters;
384 if these are similar, we consider the models interchangeable. We validate this by comparing HDI
(highest density interval) of posterior distributions.

Where posterior plots are very similar and the number of parameters is the same, we use WAIC, or
386 the widely applicable information criterion¹⁶, defined as

$$WAIC(\theta) = -2 \log \mathbb{E}[\mathcal{L}(\theta)] + 2\mathbb{V}[\log \mathcal{L}(\theta)] \quad (6)$$

388 where θ is a parameter vector, and $\mathbb{E}[\cdot]$ and $\mathbb{V}[\cdot]$ are the expectation and variance taken over the
posterior distribution, which in practice are approximated using 50,000 samples from the posterior
390 KDE. We validated that upon resampling WAIC values do not significantly change and that differences
in WAIC between models are preserved. WAIC values are scaled as a deviance measure: lower values
392 imply higher predictive accuracy.

Prior distributions. We used informative prior distributions for w_{2n+1} , w_{2n+1^*} and w_{2n^*} (we set
394 $w_{2n} = 1$), which we estimated from growth curves data from mono-culture growth experiments
previously reported by Yona et al.⁷², Figs. 3C, 4A, and S2. We used Curveball, a method
396 for predicting results of competition experiments from growth curve data⁴⁴ curveball.yoavram.com.
Briefly, Curveball takes growth curves of two strains growing separately in mono-culture and predicts
398 how they would grow in a mixed culture, that is, it predicts the results of a competition assay. From these
predictions, relative fitness values can be computed. Because Curveball uses a maximum-likelihood
400 approach to estimate model parameters, we were able to estimate a distribution of relative fitness
values to be used as a prior distribution by sampling 10,000 samples from a truncated multivariate
402 normal distribution defined by the maximum-likelihood covariance matrix (Figure S3).

We used growth curves of $2n$ and $2n + 1$ in 39 °C to estimate an informative prior distribution for
404 w_{2n+1} (Figure S3D, assuming $w_{2n} = 1$). In this prior distribution, we used the same prior for w_{2n+1^*}
and w_{2n^*} . To increase computational efficiency, we also assumed $w_{2n^*} > w_{2n+1^*} > w_{2n+1} > w_{2n}$;

406 running the inference without this assumption produced similar results. See *supporting material* for
an extended informative prior distribution that uses growth curves of $2n^*$ and $2n+1$ growing in 39 °C;
408 this prior distribution proved to be less useful.

As a control, we tested an uninformative uniform prior with $U(1, 6)$, for (i) all w_{2n+1} , w_{2n+1^*} , w_{2n^*} ,
410 or (ii) only for w_{2n+1^*} , w_{2n^*} , using the above informative prior for w_{2n+1} . In these cases the inference
algorithm failed to converge.

412 For the mutation rate, μ , and aneuploidy rate, δ , we used uninformative uniform priors, $\mu \sim U(10^{-9}, 10^{-5})$ and $\delta \sim U(10^{-6}, 10^{-2})$. A wider mutation rate prior, $\mu \sim U(10^{-9}, 10^{-3})$, produced
414 similar results.

Acknowledgements

416 We thank Lilach Hadany, Judith Berman, David Gresham, Shay Covo, Martin Kupiec, Uri Obolski, Daniel
Weissman, and Tal Simon for discussions and comments. This work was supported in part by the Israel
418 Science Foundation (ISF, YR 552/19), the US–Israel Binational Science Foundation (BSF, YR 2021276),
Minerva Center for Live Emulation of Evolution in the Lab (YR, YP), Minerva Stiftung short-term research
420 grant (MP).

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Supplementary Material

620 Supplementary Analysis

Sensitivity analysis. Changing a single parameter while keeping the rest fixed at the MAP estimate produces a worse fit to the data (Figure S6). Furthermore, we fitted models with a mutation rate fixed at $\mu = 10^{-5}$, 10^{-6} and 10^{-7} . We inferred similar parameters estimates for the model with $\mu = 10^{-6}$ compared to the model with a free μ parameter, in which the inferred mutation rate is $\mu \approx 3 \cdot 10^{-6}$. Inference assuming $\mu = 10^{-5}$ or $\mu = 10^{-7}$ produced similar estimates except that the estimated aneuploidy rate, δ , was higher, and assuming $\mu = 10^{-7}$, the estimated fitness of $2n + 1$ was lower (Figure S7).

628 Extended informative prior distribution. In an extended informative prior distribution, we used additional growth curves of $2n^*$ (*refined* strain from Yona et al.⁷²) and $2n + 1$ in 39 °C to estimate w_{2n^*}/w_{2n+1} (Figure S3H). The same distribution was used for w_{2n^*}/w_{2n+1^*} . Thus, our main informative prior uses a single prior distribution for fitness values of $2n + 1$, $2n + 1^*$, and $2n^*$, whereas the extended informative prior uses one distribution for $2n + 1$, and another distribution for both $2n + 1^*$ and $2n^*$.

634 We estimated the parameters under this extended informative prior. Inference took much longer to run but the posterior distribution seemed to converge, as it did not change much in the final **636** iterations. The posterior predictive plot shows that inference with this extended prior produces a posterior distribution that fails to explain the empirical observations (pink in Figure 3). However, **638** the inferred posterior distribution is considerably narrower (compare Figures 2 and S8) and therefore parameter estimates are less variable. The estimated mutation rate was much lower compared to **640** the main informative prior, with $\mu = 2.474 \cdot 10^{-9}$ [$2.423 \cdot 10^{-9} - 2.612 \cdot 10^{-9}$]. Other parameter estimates are: $\delta = 2.705 \cdot 10^{-3}$ [$2.094 \cdot 10^{-3} - 3.094 \cdot 10^{-3}$], $w_{2n+1} = 1.022$ [$1.021 - 1.024$], **642** $w_{2n+1^*} = 1.052$ [$1.05 - 1.054$], $w_{2n^*} = 1.053$ [$1.051 - 1.055$], the latter two being much higher compare to the main informative prior. Notably, the mode of the posterior ratio $w_{2n^*}/w_{2n+1} = 1.0009$ **644** is much lower than the mode of the prior ratio of 1.033 (Figure S3H) and closer to the ratio of 1 that we assume in the main informative prior. Together with the posterior predictive results, we conclude **646** that the main informative prior is preferable over the extended informative prior.

Model with transitions to less-fit genotypes We also estimated the parameters of a version of the **648** model that includes transitions (mutation, chromosome loss and gain) to less-fit genotypes (e.g., $2n^*$

to $2n + 1^*$),

$$\begin{aligned}
 f_{2n}^m &= (1 - \delta - \mu)f_{2n}^s + \delta f_{2n+1}^s + \mu f_{2n^*}^s , \\
 f_{2n+1}^m &= \delta f_{2n}^s + (1 - \delta - \mu)f_{2n+1}^s + \mu f_{2n+1^*}^s , \\
 f_{2n+1^*}^m &= \mu f_{2n+1}^s + (1 - \delta - \mu)f_{2n+1^*}^s + \delta f_{2n^*}^s , \\
 f_{2n^*}^m &= \mu f_{2n}^s + \delta f_{2n+1^*}^s + (1 - \delta - \mu)f_{2n^*}^s .
 \end{aligned} \tag{7}$$

The inferred values are slightly different. The estimated mutation rate, $\mu = 1.036 \cdot 10^{-7}$ [8.01 · 650 $10^{-8} - 1.339 \cdot 10^{-7}$], corresponds to a mutation target size of $\sim 300 - 500$, assuming the mutation rate per base pair is roughly $2 \cdot 10^{-10}$ (ref.⁷⁶) or $3.3 \cdot 10^{-10}$ (ref.³⁶). The estimated aneuploidy 652 rate, $\delta = 2.358 \cdot 10^{-4}$ [$1.766 \cdot 10^{-4} - 2.837 \cdot 10^{-4}$] is 5-35-fold higher than in previous studies: 654 for Chromosome III in diploid *S. cerevisiae*, Zhu et al.⁷⁶ estimated $6.7 \cdot 10^{-6}$ chromosome gain 656 events per generation, and Kumaran et al.³³ estimate $3.0 - 4.3 \cdot 10^{-5}$ chromosome loss events per generation (95% confidence interval). The estimated fitness values are $w_{2n+1} = 1.024$ [1.023–1.025], 658 $w_{2n+1^*} = 1.025$ [1.024–1.026], $w_{2n^*} = 1.032$ [1.031–1.033], all relative to the fitness of $2n$, which 660 is set to $w_{2n} = 1$.

We simulated genotype frequency dynamics using parameter samples from the posterior distribution, 662 and computed the posterior distribution of F_A . The mean F_A in this case is just 0.0189 [0.0004 - 664 0.1214 95% CI], lower than without the transitions to less-fit genotypes. Here, F_A is the sum of frequencies of both $2n_A^*$ and $2n + 1^*$, which reaches a frequency of 0.0007. Out of 100,000 posterior samples, none had F_A above 0.05 (i.e., 5% of the population).

Supplementary Figures & Tables

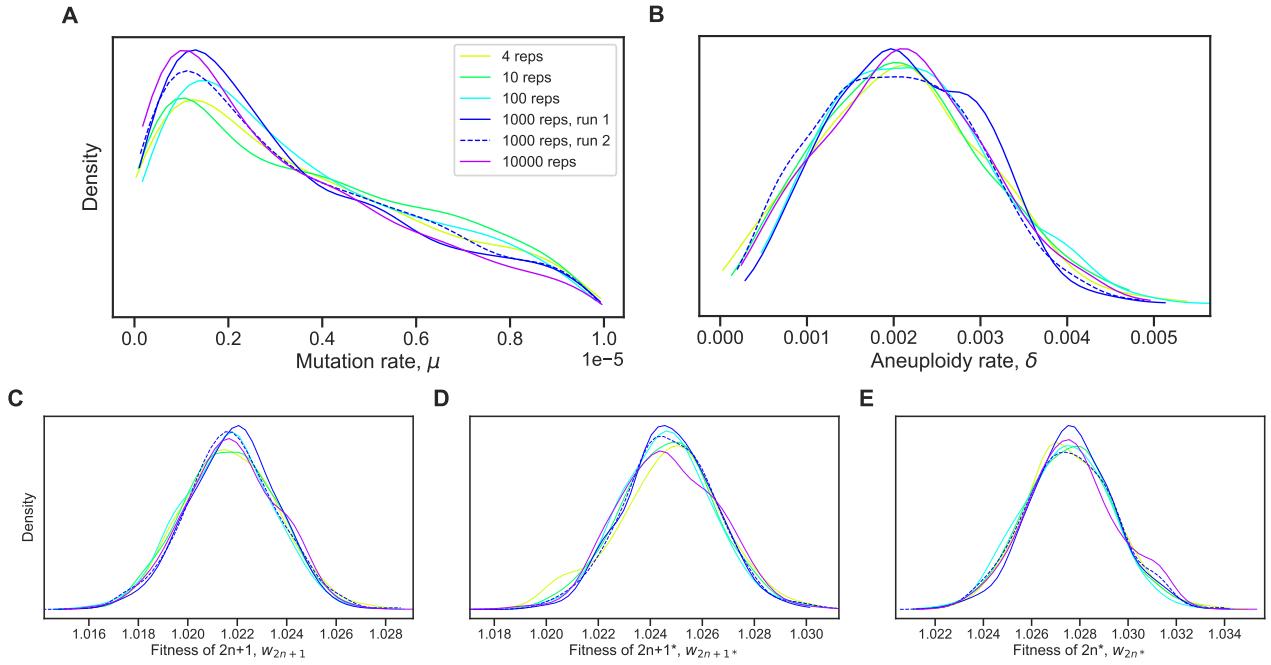


Figure S1: Posterior distribution validation. The posterior distribution of model parameters is roughly the same regardless of the number of simulations (4-10,000 replicates) used to approximate the likelihood (eq. (4)).

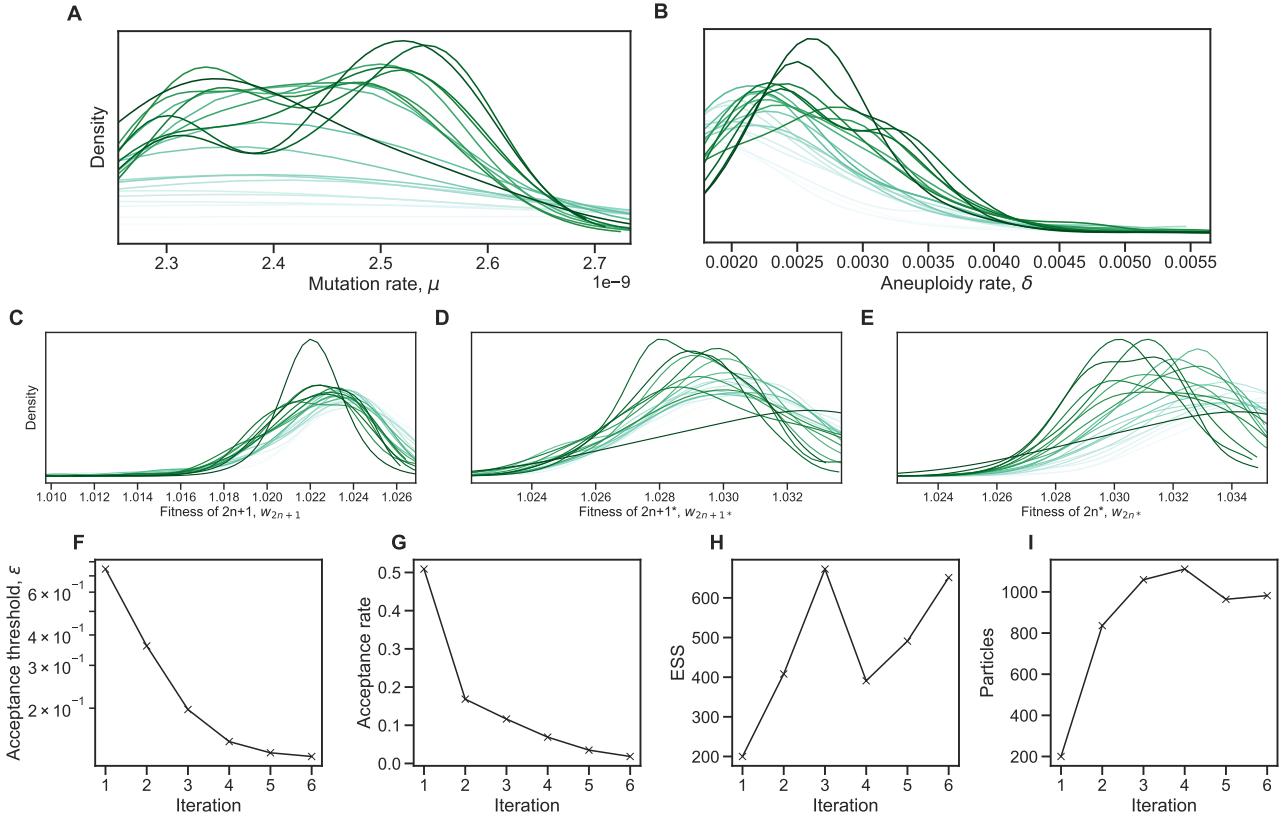


Figure S2: Inference convergence. The ABC-SMC algorithm was used to infer the model parameters. **(A-E)** The approximate posterior distributions of model parameters at each iteration of the ABC-SMC algorithm demonstrates convergence, as the posterior did not significantly change after the first iteration, $t = 1$. **(F-I)** ABC-SMC measures of convergence. After iteration number 6, the acceptance threshold was $\epsilon = 0.13$ (i.e., $\mathcal{L} = 0.87$, eq. (4)), the acceptance rate was 0.018, the number of particles was 982, and the effective sample size ESS=651.

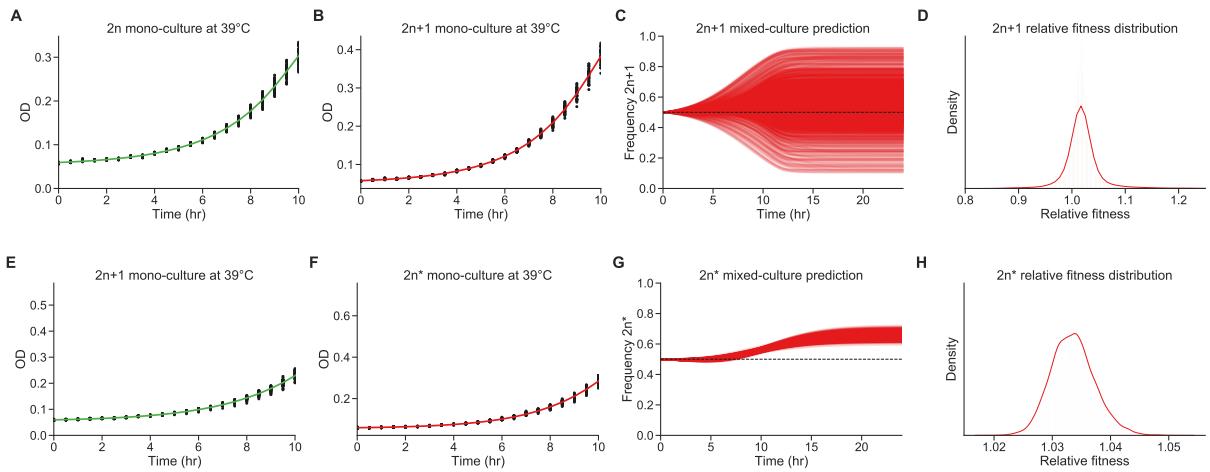


Figure S3: Fitness estimation from growth curves. **(A-D)** Fitness estimation from growth curves of $2n$ and $2n+1$ at 39°C . $w_{2n+1}/w_{2n}=1.024$ (95% CI: 0.959 - 1.115). Curveball **(E-H)** Fitness estimation from growth curves of $2n+1$ and $2n^*$ at 39°C . $w_{2n^*}/w_{2n+1}=1.033$ (95% CI: 1.027 - 1.041). Growth curves previously described in Yona et al.⁷², Figs. 3C, 4A, and S2. Fitness estimated from growth curves using Curveball, a method for predicting results of competition experiments from growth curve data⁴⁴ curveball.yoavram.com. See *Models and Methods, Prior distributions* for more details. **(A,B;E,F)** Mono-culture growth curve data (markers) and best-fit growth models (lines). **(C,G)** The mixed-culture prediction for the strains from A,B and E,F respectively, 6,375 generated curves. **(D,H)** The relative fitness distribution for $2n+1$ relative to $2n$ (panel D) and $2n^*$ relative to $2n+1$ (panel H). Figures generated by Curveball.

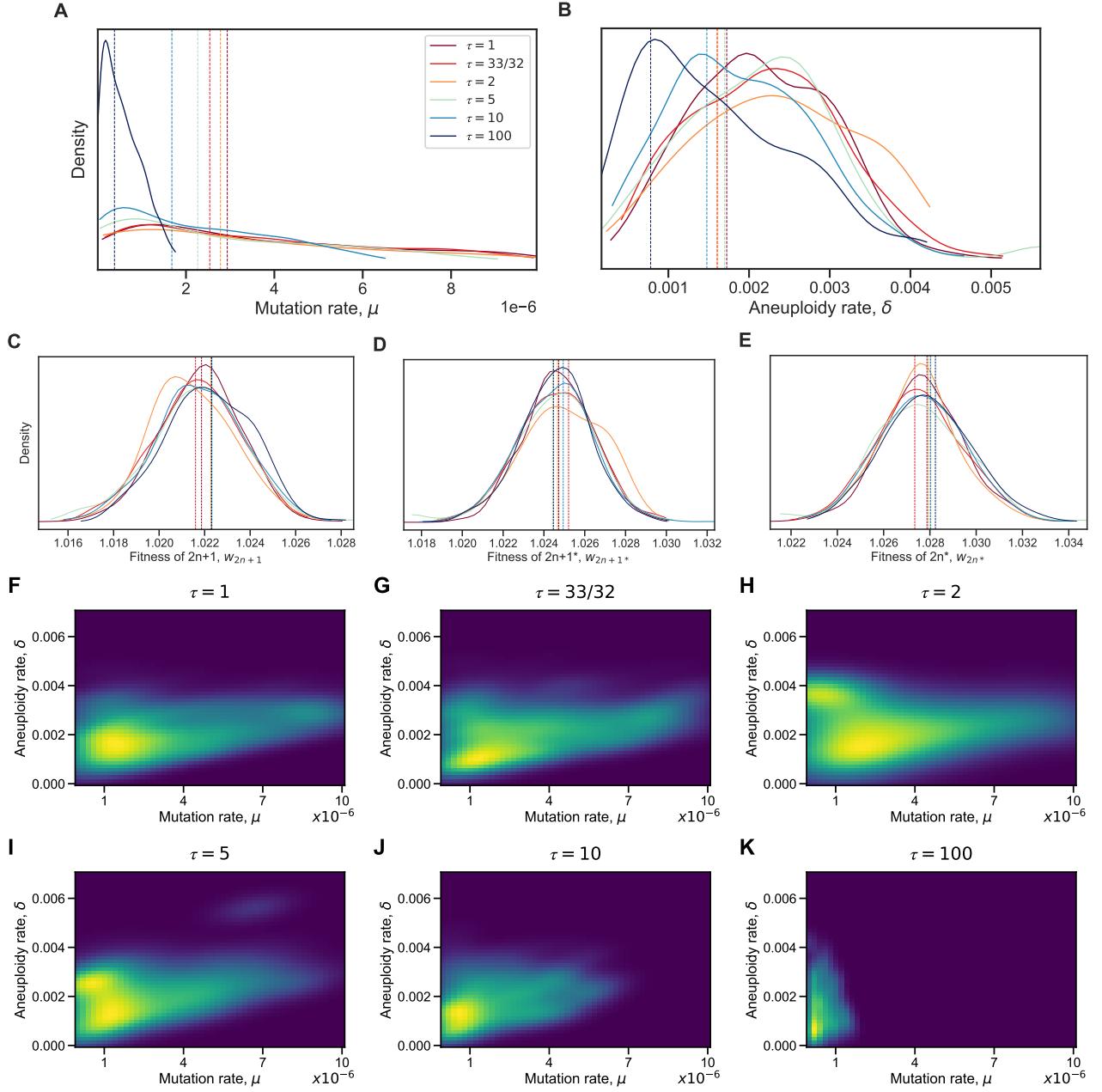


Figure S4: Model with elevated mutation rate in aneuploid cells. (A-E) The inferred posterior distributions for models with different values of τ , the fold-increase in mutation rate in aneuploid cells ($2n + 1$ and $2n + 1^*$). Vertical dashed lines represent the MAP (maximum a posteriori) of each distribution. When the increase in mutation rate is high, $\tau = 10$ and $\tau = 100$, the inferred mutation (A) and aneuploidy (B) rates tend to be lower. (F-K) The inferred joint posterior distribution of mutation rate (μ) and aneuploidy rate (δ) with different τ values (dark purple and bright yellow for low and high density, respectively).

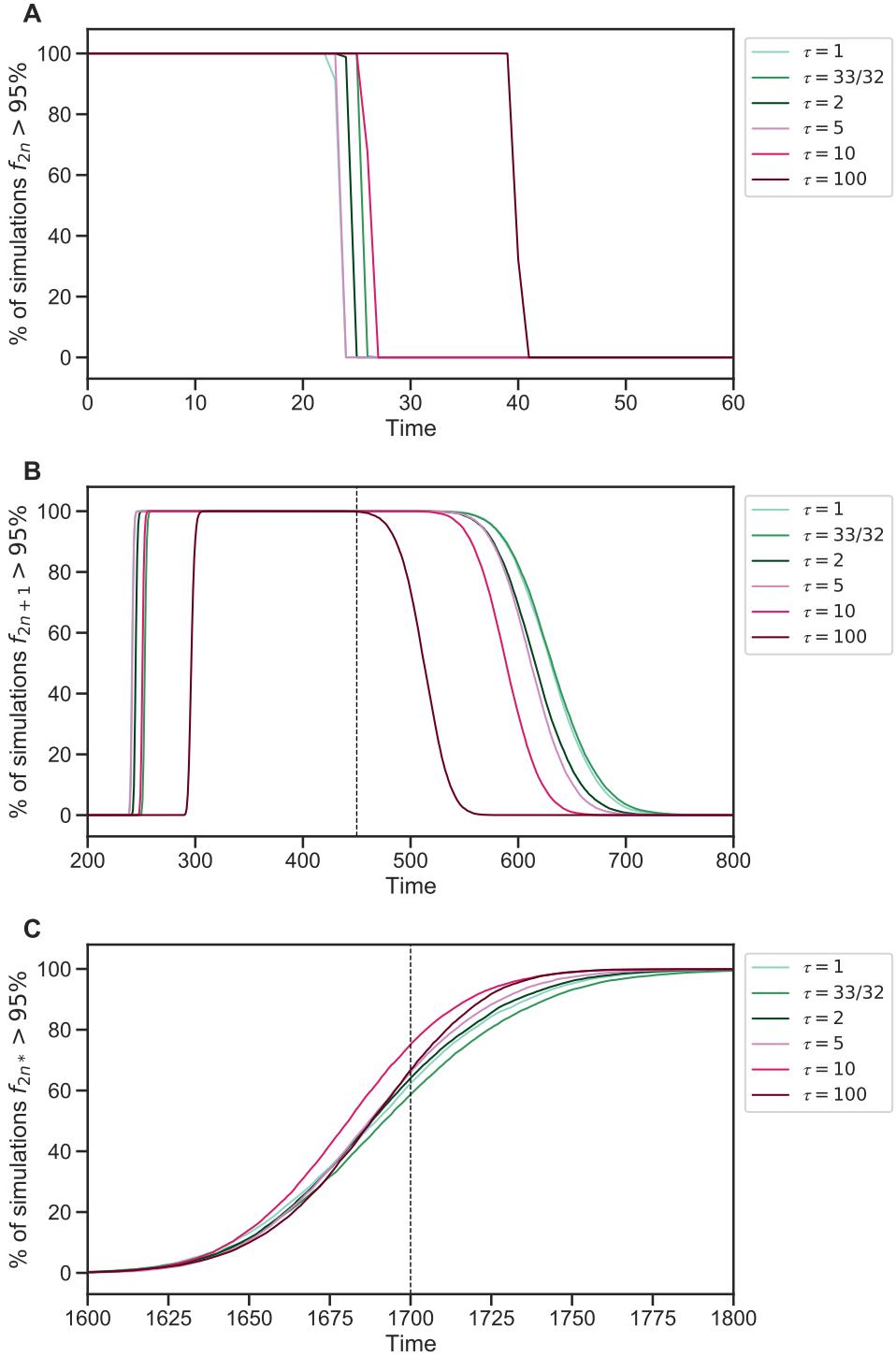


Figure S5: Genotype fixations for models with increased genetic instability. We estimated the parameters for different models, each assuming a different value of τ , the fold-increase in mutation rate in aneuploid cells. We then generated 10,000 simulations using the MAP estimate of each model and evaluated the fraction of simulations in which the frequency of genotype $2n$ (**A**), $2n + 1$ (**B**), and $2n^*$ (**C**) is above 95% (y-axis) at each generation (x-axis). Note that $2n + 1^*$ did not fix. We can see that $\tau = 100$ can be distinguished if the waiting time for $f_{2n} < 95\%$ is known (panel A) or if the waiting time for $f_{2n+1} > 95\%$ or $f_{2n+1} < 95\%$ is known (panel B). It is harder to distinguish between $1 \leq \tau \leq 10$.

Table S1: WAIC values for different τ values. Differences of less than 6 are considered of weak significance²⁷.

Model	WAIC
$\tau = 1$	-9
$\tau = 33/32$	-9
$\tau = 2$	-8
$\tau = 5$	-12
$\tau = 10$	-9
$\tau = 100$	-12

WAIC defined in eq. (6).

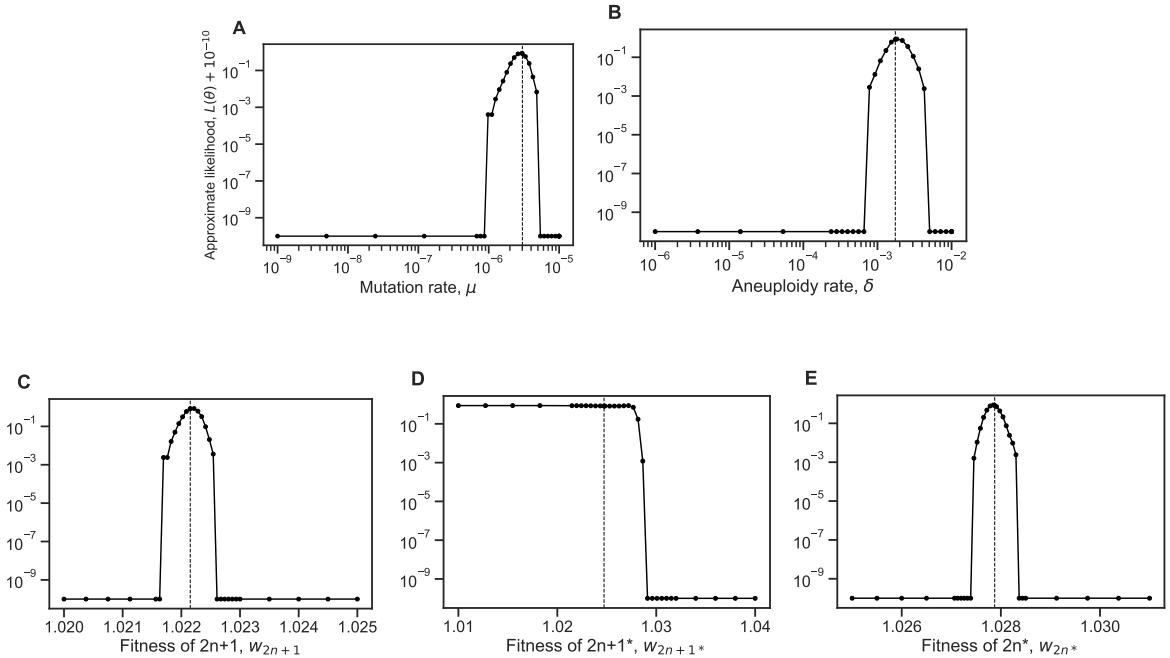


Figure S6: Likelihood profiles. Sensitivity of the model approximate likelihood, $\mathcal{L}(\theta)$, to changing a single parameter while the other parameters remain fixed at their MAP estimates. Dashed vertical line represents the MAP value. The prior distributions for the mutation rate and aneuploidy rate are $\mu \sim U(10^{-9}, 10^{-5})$ and $\delta \sim U(10^{-6}, 10^{-2})$, respectively.

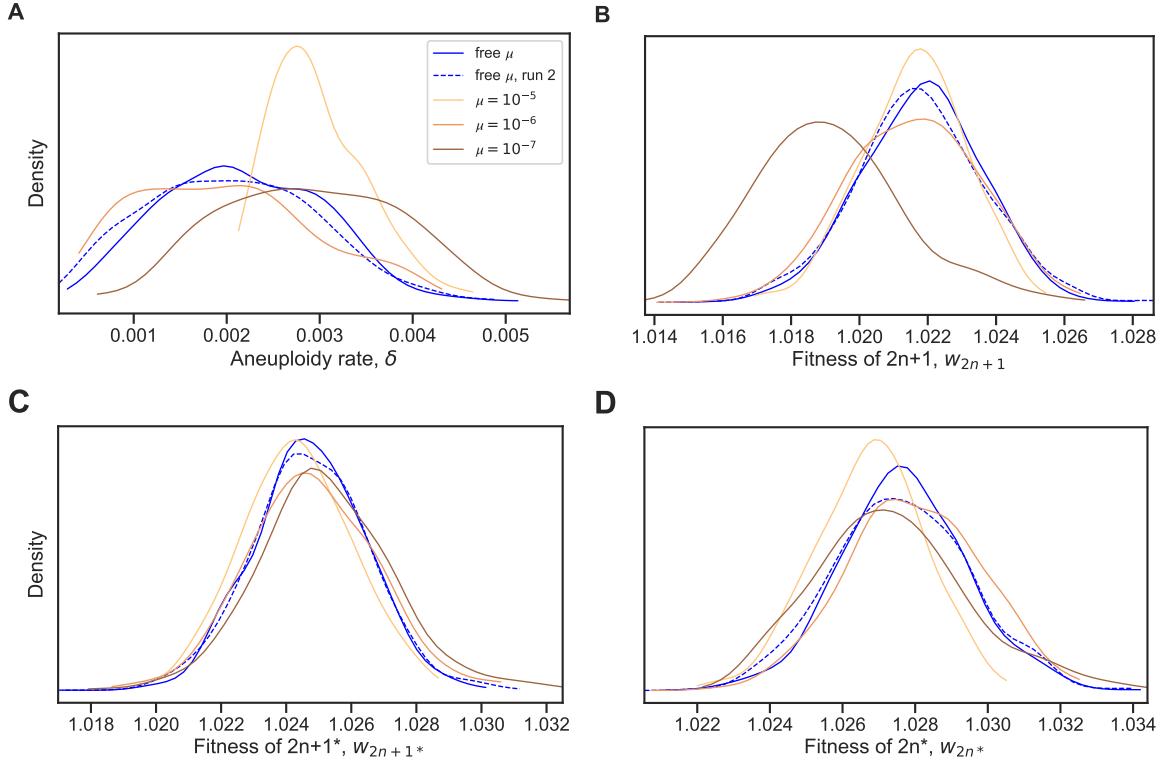


Figure S7: Model with fixed mutation rate. (A-D) The inferred posterior distributions for models with free and fixed mutation rate, μ . The MAP (maximum a posteriori) and 50% HDI (highest density interval) for each model are: **free μ , run 1:** $\delta = 1.720 \cdot 10^{-3}$ [$1.470 \cdot 10^{-3} - 2.786 \cdot 10^{-3}$], $w_{2n+1} = 1.022$ [1.021 – 1.023], $w_{2n+1^*} = 1.025$ [1.024 – 1.026], $w_{2n^*} = 1.028$ [1.026 – 1.029]; **free μ , run 2:** $\delta = 2.129 \cdot 10^{-3}$ [$1.334 \cdot 10^{-3} - 2.695 \cdot 10^{-3}$], $w_{2n+1} = 1.022$ [1.02 – 1.023], $w_{2n+1^*} = 1.025$ [1.023 – 1.026], $w_{2n^*} = 1.028$ [1.026 – 1.029]; **$\mu = 10^{-5}$:** $\delta = 2.903 \cdot 10^{-3}$ [$2.399 \cdot 10^{-3} - 3.156 \cdot 10^{-3}$], $w_{2n+1} = 1.022$ [1.021 – 1.023], $w_{2n+1^*} = 1.024$ [1.023 – 1.025], $w_{2n^*} = 1.027$ [1.026 – 1.028]; **$\mu = 10^{-6}$:** $\delta = 1.917 \cdot 10^{-3}$ [$9.624 \cdot 10^{-4} - 2.447 \cdot 10^{-3}$], $w_{2n+1} = 1.022$ [1.02 – 1.023], $w_{2n+1^*} = 1.025$ [1.023 – 1.026], $w_{2n^*} = 1.028$ [1.027 – 1.029]; **$\mu = 10^{-7}$:** $\delta = 2.901 \cdot 10^{-3}$ [$2.139 \cdot 10^{-3} - 3.671 \cdot 10^{-3}$], $w_{2n+1} = 1.019$ [1.017 – 1.02], $w_{2n+1^*} = 1.025$ [1.024 – 1.026], $w_{2n^*} = 1.027$ [1.026 – 1.029].

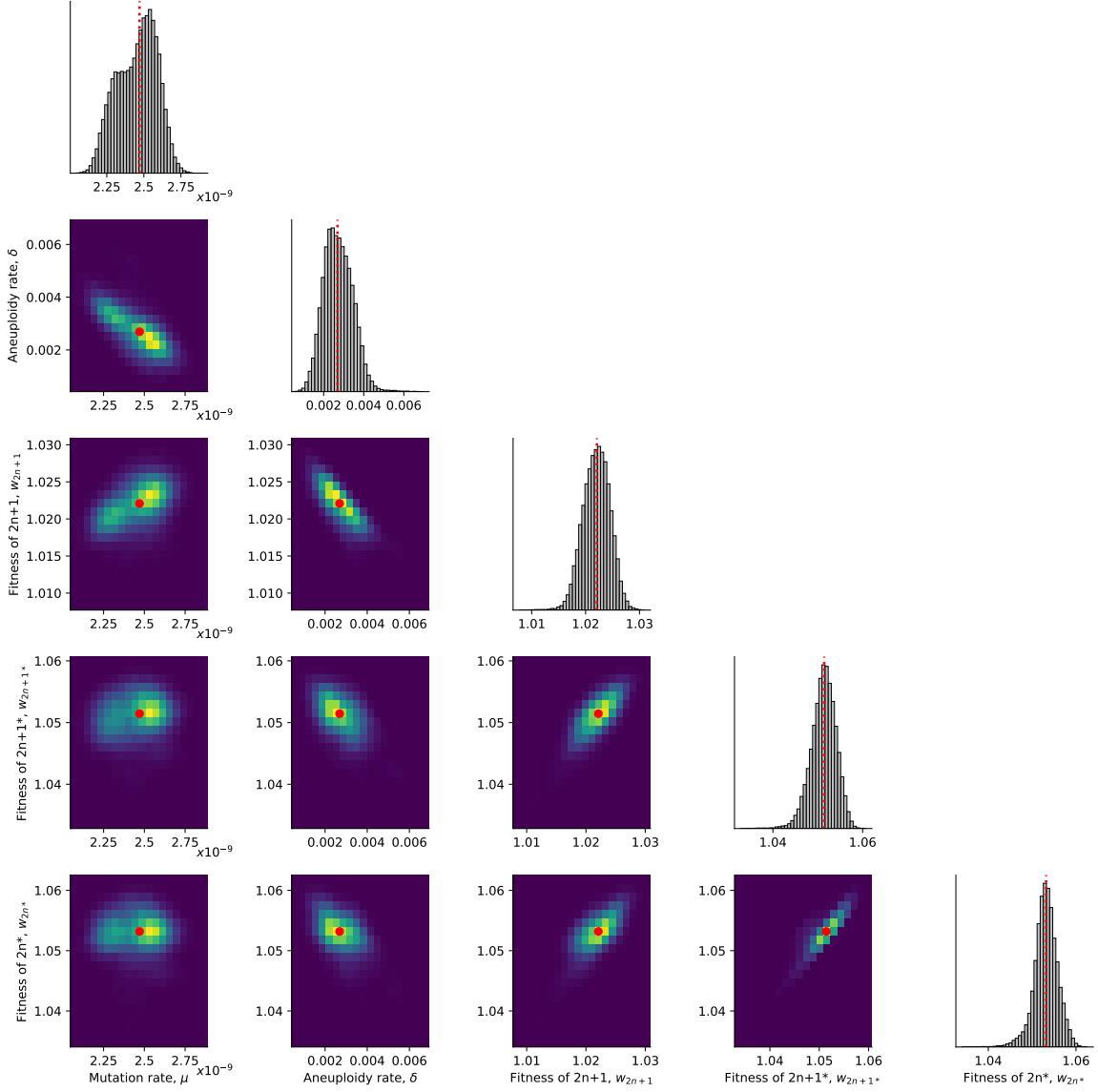


Figure S8: Posterior distribution of parameters inferred with the extended prior distribution. On the diagonal, the inferred posterior distribution of each model parameter. Below the diagonal, the inferred joint posterior distribution of pairs of model parameters (dark purple and bright yellow for low and high density, respectively). Red markers and orange lines for the joint MAP estimate (which may differ from the marginal MAP, as the marginal distribution integrates over all other parameters).

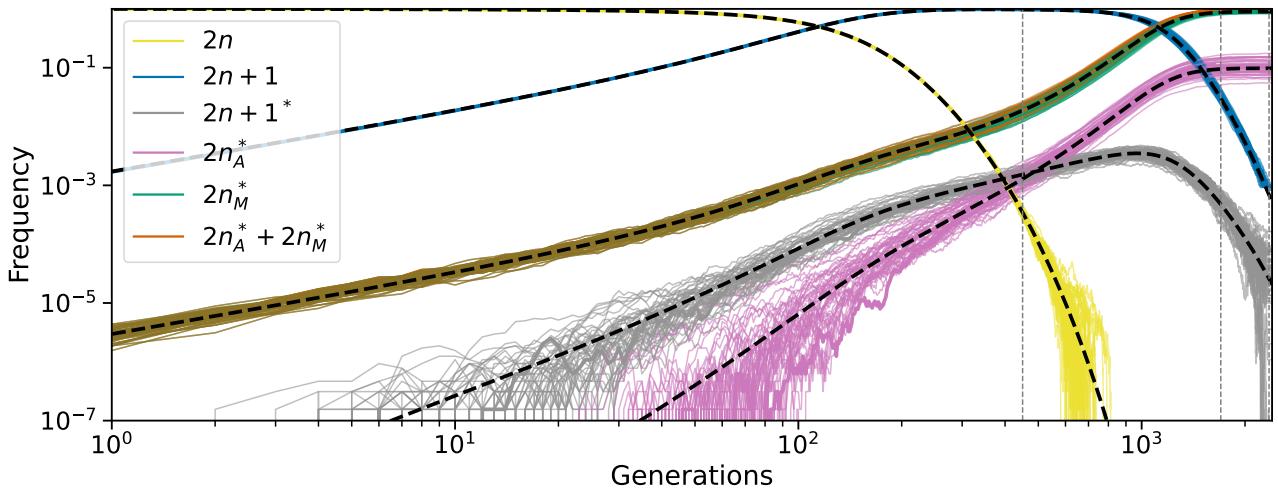


Figure S9: Posterior predicted genotype frequencies in log-log scale. Frequency dynamics of the different genotypes with MAP parameter estimates, same as Figure 4A, but in log-log scale. Black dashed curves for a deterministic model without genetic drift. Clearly, appearance of $2n + 1$ and $2n_M^*$ is deterministic. Appearance of $2n + 1^*$, and therefore $2n_A^*$, is stochastic, however, the frequency dynamics are deterministic above a frequency of roughly 0.001. Note that the $2n_M^*$ and the $2n_A^* + 2n_M^*$ lines are overlapping for much of their trajectories.

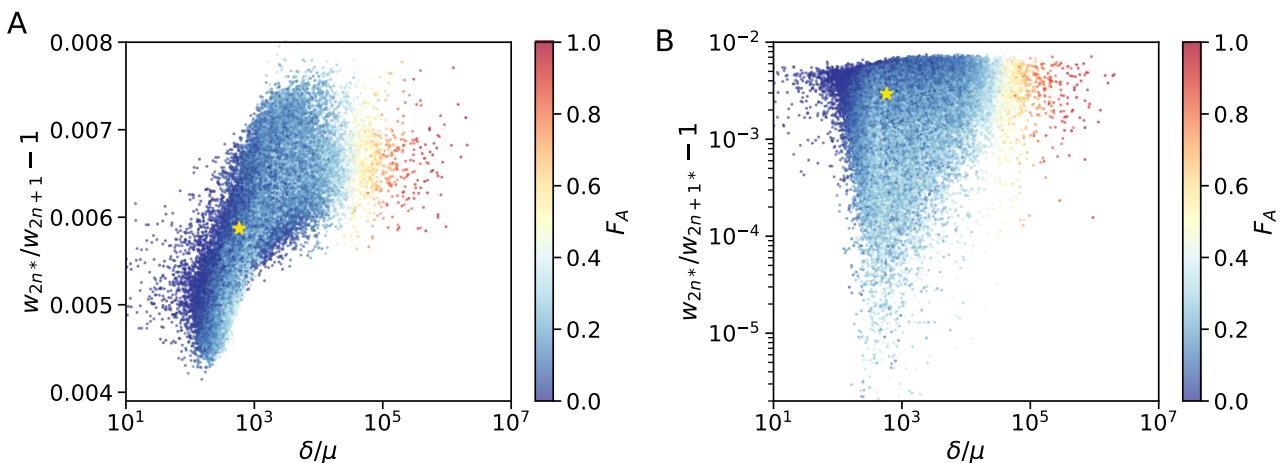


Figure S10: Posterior distribution of F_A . (A,B) F_A values (color coded) as in Figure 4 for different parameter choices on the x- and y-axes. White star denotes the MAP estimate.

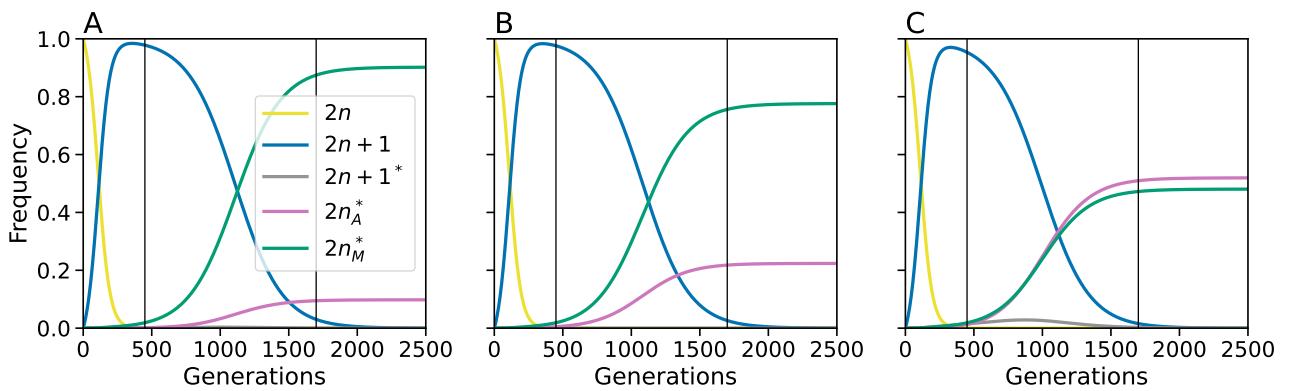


Figure S11: Effect of genomic instability on genotype frequencies. Genotype frequencies in the deterministic model without drift and with MAP parameter estimates (**A**), with 100-fold increase in rate of chromosome loss (transition from $2n+1^*$ to $2n^*$) (**B**), and with 10-fold increase in mutation rate in aneuploid cells (**C**). Corresponding F_A values (purple line at generation 2,500) are 0.098, 0.223, and 0.519.