Rampant transposition following RNAi loss causes hypermutation and antifungal drug resistance in clinical isolates of a human fungal pathogen

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Short title: RNAi loss and transposition cause hypermutation in *Cryptococcus*

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Abstract

Microorganisms survive and compete within their environmental niches and avoid evolutionary stagnation by stochastically acquiring mutations that enhance fitness. Although increased mutation rates are often deleterious in multicellular organisms, hypermutation can be beneficial for microbes in the context of strong selective pressures. To explore how hypermutation arises in nature and elucidate its consequences, we employed a collection of 387 sequenced clinical and environmental isolates of *Cryptococcus neoformans*. This fungal pathogen is responsible for ~15% of annual AIDS-related deaths and is associated with high mortality rates, attributable to a dearth of antifungal drugs and increasing drug resistance. Isolates were screened for the ability to rapidly acquire antifungal drug resistance, and two robust hypermutators were identified. Insertion of the non-LTR Cnl1 retrotransposon was found to be responsible for the majority of drug-resistant isolates. Long-read whole-genome sequencing revealed both hypermutator genomes have two unique features: 1) hundreds of Cnl1 copies organized in subtelomeric arrays on both ends of almost all chromosomes, and 2) a nonsense mutation in the first exon of *ZNF3*, a gene encoding an RNAi component involved in silencing transposons. Quantitative trait locus mapping identified a significant genetic locus associated with hypermutation that includes the mutant *znf3* allele, and CRISPR-mediated genome editing of the *znf3* single-base pair nonsense mutation abolished the hypermutation phenotype and restored siRNA production. In sum, hypermutation and drug resistance in these isolates results from loss of RNAi combined with subsequent accumulation of a large genomic burden of a novel transposable element in *C. neoformans*.

Introduction

Stochastic mutations and genomic rearrangements provide variation in populations for natural selection to act upon and enable evolution. However, genetic changes are a double-edged sword: too little variation can lead to evolutionary stagnation, while too much variation can lead to a lethal accumulation of deleterious mutations. Hypermutation, one extreme of this mutational spectrum, can lead to adaptation, disease, or eventual extinction if left unchecked.

Microbes are known to adopt highly mutable states that would normally be viewed as deleterious from the perspective of multicellular organisms. Studies have found that microorganisms with defects in pathways associated with maintaining genomic integrity, such as those involved in chromosome stability, DNA mismatch repair, DNA damage repair, and cell cycle checkpoints associated with recognizing DNA damage, accelerate adaptation to environmental stressors^{1–3}. These defects can be beneficial in the short term, yet deleterious in the long term as mutations continue to accumulate. Defects in DNA mismatch repair resulting in increased mutation rates have been reported in fungi, including the model yeast *Saccharomyces cerevisiae*, the human pathogen *Candida glabrata*, in an outbreak strain of *Cryptococcus deuterogattii*, and in several clinical isolates of the model basidiomycete human fungal pathogen *Cryptococcus neoformans*^{4–10}. Genomic stability in pathogenic *Cryptococcus* species is also significantly affected by karyotypic changes and transposable elements, both of which can mediate antifungal drug resistance^{11–14}.

Cryptococcus is an environmentally ubiquitous haploid basidiomycete and facultative human pathogen¹⁵. Approximately 95% of cryptococcal infections are attributable to the serotype A group, *C. neoformans* var. *grubii*, now known as *C. neoformans*, which is divided into four lineages: VNI, VNII, VNBI, and VNBII^{16–18}. This species infects immunocompromised

individuals and accounts for ~15% of HIV/AIDS-related deaths¹⁹. The threat of cryptococcal infections is exacerbated because the arsenal of antifungal drugs is limited. Amphotericin B, a fungicidal polyene, is often used in combination with 5-flucytosine (5-FC), an antimetabolite, to treat cryptococcal infections²⁰. Unfortunately, amphotericin B and 5-FC have undesirable side effects, and 5-FC monotherapy frequently leads to resistance^{21,22}. Fluconazole is used to treat asymptomatic patients with isolated cryptococcal antigenemia, those with disease limited to lung nodules or central nervous system infections after clearance of cerebrospinal fluid cultures, or for chronic maintenance therapy²⁰. However, *C. neoformans* frequently develops resistance to fluconazole via aneuploidy or mutations in the sterol biosynthesis pathway, contributing to recurrent infections^{11,23–25}. The limited number of drugs available to treat cryptococcosis, prevalence of resistance and recurrent infections, and difficulty in developing novel antifungal therapies combine to make *C. neoformans* drug resistance an important clinical problem.

Transposons in the *C. neoformans* H99 reference strain and the sister species *Cryptococcus deneoformans* JEC21 reference strain have been characterized^{13,26,27}. The genomes of these species encode many retrotransposons with and without long-terminal repeats, known as LTR retrotransposons and non-LTR retrotransposons, respectively, which move via a copy-and-paste mechanism, allowing them to proliferate throughout the genome if unchecked. The most well-characterized *Cryptococcus* LTR-retrotransposons are Tcn1 through Tcn6, which are primarily located in centromeric regions²⁸. The *C. deneoformans* JEC21 genome also encodes three types of DNA transposons (T1, T2, and T3), as well as ~25 copies of the non-LTR retrotransposon Cnl1 (*C. neoformans* LINE-1), which is thought to associate with telomeric repeat sequences^{13,14,29}. In the *C. neoformans* H99 genome, there are no full-length copies of Cnl1, and DNA transposons are rare¹³.

Studies have illustrated that transposon silencing in *Cryptococcus* is governed by RNAi through three primary lines of evidence: 1) siRNAs map predominantly to transposable elements, 2) RNAi mutants show increased transposon expression, and 3) spliceosomes stall on transposable element transcripts at an unusually high rate, triggering RNAi^{13,30–34}. Other mechanisms thought to regulate *Cryptococcus* transposons include 5-methylcytosine DNA methylation^{28,35,36} and heterochromatic marks³⁷. Interestingly, the outbreak species *C. deuterogattii* is RNAi deficient because many genes encoding RNAi components are severely truncated or absent entirely³². This loss of RNAi has been shown to be associated with loss of all functional transposable elements, consequently shorter centromeres, and higher rates of intron retention^{28,38}.

Here, we identified two clinical, hypermutator *C. neoformans* isolates with significantly increased mutation rates on antifungal drug media. The majority of drug resistance in these two strains is mediated by insertions of the Cnl1 transposon into genes whose mutation confers resistance. Genetic backcrossing, quantitative trait loci mapping, and CRISPR-mediated gene editing all confirmed that a nonsense mutation in the RNAi component *ZNF3*, resulting in loss of RNAi, is the cause of hypermutation in these two strains. Small RNA sequencing confirmed the role of Znf3 in silencing Cnl1, and whole-genome sequencing revealed both hypermutator genomes encode >800 copies or fragments of Cnl1. This is the first time full-length copies of Cnl1 have been identified in *C. neoformans*, and the massive Cnl1 burden in these hypermutators is substantially higher than previously observed in any other *Cryptococcus* strain. Our results demonstrate the hypermutator phenotype described here is attributable to loss of RNAi, allowing rampant transposition of Cnl1. These transposition events lead to Cnl1 accumulation at subtelomeres and movement to novel genomic locations, which can result in drug resistance.

Results

Identification of two clinical, hypermutator *C. neoformans* isolates

To identify natural isolates of *C. neoformans* with increased mutation rates, we screened strains in the Strain Diversity Collection (SDC) for an increased ability to produce colonies resistant to various classes of antifungal drugs. The SDC contains 387 strains from all four *C. neoformans* lineages (VNI, VNII, VNBI and VNBII), including geographically diverse clinical and environmental isolates of both mating types. For each isolate in this collection, whole-genome sequencing (WGS) and phylogenetic relationships are available¹⁸. To screen isolates for increased mutation rates in a relatively high-throughput manner, strains were grown in liquid cultures, swabbed onto agar plates supplemented with either 5-fluorocytosine (5-FC) or a combination of FK506 and rapamycin (immunosuppressants that bind FKBP12 to form complexes that inhibit activity of calcineurin and TOR, respectively), and qualitatively evaluated for their ability to generate resistant colonies (Figure 1A)^{39–42}. Strains that produced more spontaneously resistant colonies on average than the control strain H99 were categorized as hypermutator candidates. We screened 186 strains and identified 40 hypermutator candidates (Table S1). Interestingly, all but one of fourteen (93%) environmental isolates screened were identified as hypermutator candidates (as compared to only 16% (27/170) clinical isolates, *p*-value < 0.001, Fisher's exact test). Two previously identified hypermutator strains with mismatch repair defects, C23 and C45, were identified as hypermutator candidates as well⁹.

We chose to focus on two clinical strains, Bt65 and Bt81, that produced the most rapamycin + FK506-resistant (R+F^R) colonies (Figure 1A). Bt65 and Bt81 are both VNBII *MATa* strains isolated from different HIV-positive individuals in Botswana^{18,43}. To quantify the mutation rates of Bt65 and Bt81, we performed fluctuation assays on YPD + rapamycin +

Figure 1. Hypermutation in Bt65 and Bt81 is driven primarily by the insertion of Cn1 into *FRR1*. (A) Generation of spontaneously resistant colonies on YPD + rapamycin + FK506 medium was utilized to identify hypermutator candidates; pictures of representative plates are shown. Strains include the phylogenetically closely related strains involved in fluctuation assay in B as well as positive (*msh2Δ*) and negative (H99) controls. (B) Mutation rates of closely related VNBII strains and controls on YPD + rapamycin + FK506. Bars represent the mutation rate and error bars represent 95% confidence intervals; mutation rates represent the number of mutations per cell per generation. Schematic depicts the phylogenetic relationships of all strains included in fluctuation analyses based on Desjardins et al. 2017¹⁸. Mutational spectra in *FRR1* in YPD + rapamycin + FK506-resistant colonies of H99, Bt65, and Bt81 as characterized by (C) gel electrophoresis and Sanger sequencing of *FRR1* PCR products. MicroINDELs are defined as insertions or deletions < 50 bp. All mutations are relative to the appropriate rapamycin + FK506-sensitive parental strain.

FK506, YNB + 5-fluoroorotic acid (5-FOA), and YNB + 5-FC media. Both Bt65 and Bt81 produced significantly higher mutation rates on YPD + rapamycin + FK506 compared to H99 as well as eleven of the most closely phylogenetically related strains (Figure 1B). On 5-FC, only the strain NRHc5014.ENR and the KN99 α *msh2* Δ positive control had significantly higher mutation rates compared to H99 (Figure S1A); on 5-FOA medium (Figure S1B), only KN99 α *msh2* Δ produced a significantly higher mutation rate.

A recent study illustrated how incubation at an elevated temperature of 37°C results in increased mutation rates due to transposon mobilization in the closely related species *C. deneoformans*¹⁴. To determine if elevated temperature contributed to hypermutation in Bt65 and Bt81, we concurrently grew these strains as well as wild-type H99 and *msh2* Δ , *ago1* Δ , and *rdp1* Δ genetic deletion mutants in the H99 genetic background overnight at 30°C and 37°C, and then performed a fluctuation analysis on YPD + rapamycin + FK506 medium. Fluctuation analysis revealed H99, Bt65, and Bt81 had lower mutation rates when grown overnight at 37°C compared to growth at 30°C (Figure S2). Interestingly, all of the genetic deletion mutants showed increased mutation rates after growth at 37°C (Figure S2). These results suggest that unlike *C. deneoformans*, growth at 37°C reduces mutation rates in wild-type *C. neoformans* strains and does not contribute to or exacerbate hypermutation in Bt65 and Bt81.

Characterization of mutation spectra in *C. neoformans* hypermutator strains

After quantifying the mutation rates of Bt65 and Bt81 strains on media with various antifungal drugs, we investigated the types of mutations conferring resistance to the combination of rapamycin and FK506. PCR amplification of the *FRR1* gene (which encodes FKBP12, the shared target of rapamycin and FK506, and the only gene in which mutations confer resistance to

both FK506 and rapamycin) followed by gel electrophoresis revealed the expected wild-type PCR product size (~1.2 kb) for all but two (35/37) H99 R+F^R colonies; the remaining two produced PCR products smaller than expected, indicative of deletions (Figure S3A). In contrast, large insertions of various sizes were observed in the majority of the Bt65 and Bt81 R+F^R colonies (62/77 and 36/37 independent colonies, respectively) (Figure 1C, S3B, C). Only one resistant colony derived from a non-hypermulator strain, Bt84, (1/10 independent colonies) had an insertion in *FRR1*. No insertions in *FRR1* were observed in any of the other closely related or control strains.

We subsequently sequenced *FRR1* in H99, Bt65, Bt81, and Bt84 R+F^R colonies to determine the genetic changes responsible for the larger PCR products (Figure 1C). In 37 R+F^R colonies of the H99 control strain, SNPs in *FRR1* were largely responsible for resistance (57%, 21/37 colonies), while resistance in the remaining colonies was attributable to small insertions/deletions (microINDELs; 38%, 14/37) or large deletions (5%, 2/37). Conversely, in the hypermutator isolate Bt65, insertions of the non-LTR retrotransposon Cn11 were responsible for the majority of rapamycin + FK506 resistance (77.4%, 24/31). Rapamycin + FK506 resistance in the remaining Bt65 colonies was either due to SNPs (19.4%, 6/31) or microINDELs (3.2%, 1/31). In all sequenced PCR products from R+F^R colonies of Bt81, Cn11 insertions were responsible for resistance (27/27 colonies). Cn11 insertions in Bt65 and Bt81 ranged from 54 bp to ~3500 bp, and this range in transposon sizes is a common characteristic of non-LTR retrotransposons. The single insertion observed in *FRR1* in the R+F^R colony of Bt84 had no homology with any annotated *Cryptococcus* transposons but was identified as a repetitive element by RepeatMasker and shared minor homology with a Copia-58 BG-I transposable element.

The 5-FC- and 5-FOA-resistant colonies of Bt65, Bt81, and H99 were similarly characterized to determine the sources of resistance to antifungal drugs with different mechanisms of action. Resistance to 5-FOA is conferred by mutations in the *URA3* or *URA5* genes of the uracil biosynthesis pathway^{44,45}. Among the subset of H99, Bt65, and Bt81 5-FOA^R colonies sequenced, mutations were only identified in *URA5*. In almost all colonies, resistance was conferred by SNPs or INDELs, and only one Cnl1 insertion event was identified in a Bt81 5-FOA^R colony (Figure S3D). We also PCR amplified genes in which mutations are known to confer resistance to 5-FC, including *FURI* and *UXS1*⁴⁶. Only two of the 5-FC^R isolates analyzed from H99, Bt65, and Bt81 produced a PCR product larger than expected for all assessed genes, and subsequent sequencing revealed a Cnl1 insertion in *FURI* in two independent 5-FC^R Bt81 isolates (Figure S3E).

Analysis of the Cnl1 insertions observed to confer resistance to rapamycin + FK506, 5-FC, and 5-FOA revealed Cnl1 preferentially inserts at guanine- and cytosine-rich regions of target genes, a known property of this element²⁷. Target site duplication sequences flanking Cnl1 insertions were not present in many instances, but when present, ranged from 1 to 12 bp in length. Cnl1 insertions ranged greatly in size, from 25-bp fragments to full-length Cnl1 copies (3,494 bp). The smallest Cnl1 insertion (25 bp) was followed immediately by a 59-bp deletion in *FRR1*. Cnl1 insertions in the *FRR1* gene were observed in both orientations (5' to 3' and 3' to 5' relative to *FRR1* transcription). Of the 51 characterized Cnl1 *FRR1* insertions, 27 were in the 5' UTR, and all but one were in the same orientation as *FRR1* transcription, 23 were in exons (7 oriented 5' to 3', 16 oriented 3' to 5'), and one insertion was in an intron of *FRR1* in the 3' to 5' orientation, potentially disrupting splicing or transcription.

QTL mapping identifies loci that significantly contribute to the hypermutator phenotype

To determine the genetic cause of the hypermutator phenotype and rampant transposition in Bt65 and Bt81 and to determine the genetic consequences of this phenotype, we conducted quantitative trait locus (QTL) mapping. For this purpose, a total of 165 basidiospores were dissected from a genetic cross between Bt65 *MATa* and an H99 *crg1Δ MATa* mutant with an enhanced mating phenotype, and 47 F₁ progeny germinated (28%).

Twenty-eight Bt65a x H99a F₁ progeny were selected for fluctuation analysis and whole-genome sequencing. Aligning the paired-read Illumina sequencing data from the 28 F₁ progeny identified 215,411 bi-allelic SNPs that were utilized for QTL mapping. For 24 of the segregants as well as for the Bt65 and H99 *crg1Δ* parental strains, the mutation rate on rapamycin + FK506 medium served as the phenotype for association tests (Figure 2A). Across the 14 chromosomes and bi-allelic SNP sites, two QTL with large effect (heritability = 64%) were identified at approximately 919-1,120 kb on Chromosome 3 and 987-1,193 kb on Chromosome 11 (Figure 2B and S4). Analysis of these QTLs revealed that the SNPs in each QTL were co-segregating and that they shared the same distributions of phenotype scores (Figure S4 and S5). The borders of the QTL spanning Chromosomes 3 and 11 were determined by calculating 95% confidence intervals and examining recombination break points along each chromosome. Interestingly, these two QTLs span the chromosomal translocation between Chromosomes 3 and 11 that is unique to H99 (Figure S4, S5, S6). We thus treated these QTLs as the same QTL for subsequent analysis.

Within the QTL there are a total of 108 and 85 genes along Chromosome 3 and Chromosome 11, respectively, and for 82 and 77 of these genes (respectively), the published annotation and SNP data was used to characterize differences in predicted protein sequence and expected protein lengths between the H99 and Bt65 parental strains (Figure S4 and Table S2).

Figure 2. QTL analysis of hypermutator phenotype. (A) Quantification of mutation rates on YPD + rapamycin + FK506 medium – sorted smallest to largest, left to right – for F₁ progeny and parental strains, H99 *crg1Δ* and Bt65. Inheritance of the Bt65 *znf3* allele or H99 *crg1Δ* *ZNF3* allele in F₁ progeny is indicated above mutation rates. Colored bar plots and vertical black lines depict the mean mutation rate and associated 95% confidence intervals (CI) per segregant. Mutation rates represent the number of mutations per cell per generation. (B) Manhattan plot showing the strength in association (y-axis) between bi-allelic SNPs and hypermutator phenotype, across the 14 chromosomes (x-axis). Colors separate SNPs across chromosomes. The permutation-based significance threshold ($\alpha = 0.01$) is depicted with a horizontal dashed line. (C) Predicted *ZNF3* gene and *Znf3* protein models in H99 and Bt65. A grey horizontal bar depicts the gene body in the upper panel, and larger grey rectangles represent exons; the gene is depicted 5' to 3' and is 5417 nt in length. The locations of SNPs differing

between Bt65 and H99 are shown by vertical black rungs along the gene model. Amino acids specified by mRNA codons in the indicated region of *ZNF3* exon 1 (nucleotides 364 to 411) are shown for H99 and Bt65 to illustrate the effect of the C to T mutation (nucleotide 409) predicted to cause a nonsense mutation in Bt65. The bottom panel depicts the predicted impact of the nonsense mutation on the Znf3 protein in Bt65. White rectangles along the protein schematic depict the three C2H2-type zinc finger domains of Znf3.

Among these, 71 and 60 genes along Chromosome 3 and Chromosome 11, respectively, have at least one predicted nonsynonymous change in protein sequence, seven of which harbor a predicted nonsense (i.e. stop-gain) or stop-loss mutation. One of these genes is *ZNF3* (CNAG_02700), which encodes a C2H2 type zinc finger protein with three zinc finger domains. Znf3 was previously identified as an RNAi silencing component that localizes to P-bodies and whose mutation results in increased expression of transposable elements^{32,33}. *ZNF3* is located on Chromosome 3 and has a SNP – C to T – within the first exon in the Bt65 genetic background, which is predicted to cause a nonsense mutation, severely truncating Znf3 from 1,561 amino acids to only 96 amino acids (Figure 2C). In addition, this nonsense mutation may also result in nonsense-mediated mRNA decay of the mutant *znf3* mRNA. Based on the publicly available whole-genome sequencing of all isolates in the SDC, the *znf3* nonsense mutation in exon 1 is unique to Bt65 and Bt81 and not present in any other strain. Another gene of known function within the QTL encodes a long-chain acyl-CoA synthetase (CNAG_01836, Chromosome 11) and a SNP – G to A – within the last exon of this gene is predicted to cause an early nonsense mutation in the Bt65 background (Figure S7). Given the dramatic difference in the predicted protein length of *ZNF3* between the H99 and Bt65 parental alleles (relative to other genes in this QTL with predicted stop-loss or nonsense mutations), and previous studies demonstrating the role of Znf3 in RNAi and transposon silencing, we hypothesized *ZNF3* could be the quantitative trait gene (QTG) and the SNP leading to the predicted stop gain in the first exon could be the quantitative trait nucleotide (QTN) underlying the hypermutation phenotype^{32,33}.

Few Bt81 F₁ progeny display a hypermutator phenotype

In addition to generating and analyzing Bt65a x H99a *crgI*Δ F₁ progeny, 42 F₁ progeny were derived from a genetic cross between the other hypermutator strain, Bt81a, and H99a *crgI*Δ. The *ZNF3* alleles of all 42 Bt81 F₁ progeny were sequenced to determine whether they had inherited the non-functional *znf3* allele from Bt81 or the functional *ZNF3* allele from H99. Of the 42 F₁ progeny, only four inherited the mutant *znf3* allele from Bt81, a significantly lower number than would be expected based on Mendelian inheritance patterns (chi-square test, *p*-value < 0.01). The four progeny with non-functional Bt81 *znf3* alleles had the highest mutation rates of 18 F₁ progeny that were analyzed (Figure S8). The mutation rates for three of the four *znf3* progeny, however, were not significantly higher than the Bt81 F₁ progeny with functional the *ZNF3* allele and were not as high as would be expected based on the results from the Bt65a x H99a F₁ progeny (Figure 2A).

Cnl1 elements are organized into subtelomeric arrays in hypermutator genomes

For all strains in the SDC, including Bt65 and Bt81, only short-read WGS data was available¹⁸. Because of the known difficulties in assembling repetitive elements, such as Cnl1, with short-read sequencing data, we conducted long-read WGS with the Oxford Nanopore Technologies MinION sequencing platform to generate more complete assemblies for Bt65, Bt81, and two of the most closely phylogenetically related non-hypermutator strains, Bt89 and Bt133. With the long-read sequencing data, we were able to assemble chromosome-level genomes for all four strains. In the assemblies, we observed the known chromosomal translocation between Chromosomes 3 and 11 unique to H99⁴⁷ and identified a translocation

between Chromosomes 1 and 13 unique to Bt65 and Bt81 (Figure S6). These two gross chromosomal rearrangements explain the relatively low germination frequency (28%) of Bt65a x H99 α F₁ progeny because each translocation should decrease germination by ~50%.

Analysis of the genomes of Bt65 and Bt81 revealed large arrays of the Cnl1 transposon at all but one end of each of the 14 linear chromosomes (27/28 subtelomeric regions in Bt65 and 28/28 in Bt81) (Figure 3A, 3B). The assembled Cnl1 arrays (defined as ≥ 2 Cnl1 copies) in Bt65 and Bt81 range from 5 kb to 80 kb in length. These highly repetitive arrays made it difficult and, in some instances, impossible to confidently assemble telomeric repeat sequences at the ends of each Bt65 and Bt81 chromosome. Using manual telomere extension via read mapping, we were able to identify telomere repeats at only 20 chromosome ends in Bt65 and 13 in Bt81. In contrast to Bt65 and Bt81, genome assemblies for Bt89 and Bt133 were assembled with telomere repeats on all 28 chromosome ends without any manual extension (Figure S6). In these assemblies, some telomeres had no copies of Cnl1 while others had Cnl1 arrays up to 30 kb in length (Figure S9).

Further analysis revealed the Bt65 genome harbors at least 414 fragments of Cnl1, including 105 full-length copies, while the Bt81 genome appears to encode even more Cnl1 elements, with at least 449 fragments, including 147 full-length copies (Table 1). It is important to note that due to incomplete ends for most chromosomes, it is likely Bt81 and Bt65 encode additional copies of Cnl1 that were not assembled. The presence of long Cnl1 arrays in Bt65 and Bt81 was surprising because the *C. neoformans* H99 reference strain encodes only 22 fragments of Cnl1 and no full-length copies, and therefore, *C. neoformans* was not thought to harbor functional Cnl1 elements (Figure 3C) (Table 1).

Apart from the subtelomeres, retrotransposons in *Cryptococcus* are also enriched at centromeres, specifically the LTR retrotransposons Tcn1-Tcn6^{26,28,47}. The changes in Cnl1

Figure 3. Retrotransposon content in the genomes of H99, Bt65, and Bt81. Distributions of the Tcn1 through Tcn6 LTR-retrotransposons and the Cn11 non-LTR retrotransposon in subtelomeric and centromeric regions of (A) Bt65, (B) Bt81, and (C) H99 genomes depicted in Figure S6. In Bt65 and Bt81, 80 kb of subtelomeric regions are displayed, and 50 kb subtelomeric regions are displayed for H99 to show the full distribution of subtelomeric Cn11 elements. Subtelomeric arrays of Cn11 are depicted at the end of each chromosome in Bt65 and Bt81, while only 7 Cn11 elements are localized subtelomerically in H99. Shading corresponds to fragments of the Cn11 elements, and gene arrowheads indicate the direction of transcription for all retrotransposons.

transposon content in Bt65 and Bt81 along with a previous study establishing a link between loss of RNAi and centromere length²⁸ motivated us to characterize the centromeres in Bt65, Bt81, Bt89, and Bt133. Identification of centromeres in these four isolates revealed shorter centromeres on average compared to H99: 40.3 kb in Bt65, 40 kb in Bt81, 35.6 kb in Bt89, and 41.2 kb in

Table 1. Cnl1 burden in H99, hypermutator strains, related non-hypermutator strains, and six Bt65 x H99 *crg1Δa* F₁ progeny based on Nanopore sequencing data.

Strain	Hypermutator Status	Total Cnl1 burden (>50 bp)	Full-length Cnl1 copies (>99% in length)
H99	Non-hypermutator	22	0
Bt65	Hypermutator	414	105
Bt81	Hypermutator	449	147
Bt89	Non-hypermutator	261	24
Bt133	Non-hypermutator	246	23
Progeny 2	Hypermutator	212	30
Progeny 8	Hypermutator	172	40
Progeny 14	Non-hypermutator	296	68
Progeny 18	Non-hypermutator	321	88
Progeny 20	Non-hypermutator	425	136
Progeny 34	Hypermutator	187	41

Bt133, compared to an average length of 47.3 kb in H99. However, this difference did not reach statistical significance (ANOVA, p -value = 0.153, Figure S10 and Table S3). Many centromeres in the assessed VNBII isolates had undergone numerous rearrangements and several inversions relative to one another (Figure S11). Centromeric alterations have also previously been observed in *C. neoformans* genetic deletion mutants lacking the canonical RNAi components Ago1 and Rdp1²⁸. Combined, these analyses suggest that while Cnl1 is more abundant in the hypermutators Bt65 and Bt81 as well as the non-hypermutators Bt89 and Bt133, other retrotransposons are not substantially increased in number compared to H99 (Figure 3 and S9).

Characterization of H99 *crg1Δ* x Bt65 F₁ progeny genomes reveals invasion of Cnl1 elements into naïve telomeres

Expression of transposable elements, including Cnl1, has been shown to be upregulated during sexual reproduction in *C. neoformans*^{30,32,34}. To investigate how increased expression of

Cnl1 during mating impacts the genome, six of the 28 progeny utilized for QTL mapping were also selected for long-read whole-genome sequencing on the nanopore MinION sequencing platform: three hypermutator progeny that inherited the Bt65 *znf3* allele (P2, P8, and P34) and three non-hypermutator progeny that inherited the H99 *ZNF3* allele (P14, P18, and P20). Nanopore sequencing identified recombination points across the genomes of the progeny, providing information on which genomic regions were inherited from either parent and confirming these were F₁ genetic recombinants (Figure 4).

Figure 4. Genetic recombination sites and Cnl1 distribution in Bt65 x H99 F₁ progeny.

Recombination sites along each of the 14 chromosomes for the six Bt65 α x H99 α F₁ progeny for which long-read whole-genome sequencing was conducted. Genomic loci depicted in blue were inherited from the H99 parent, and orange genomic loci were inherited from the Bt65 parent. Cnl1 elements throughout the F₁ progeny and parental genomes are indicated by black arrowheads in the upper panel. Centromeres are indicated by dark blue boxes in only the parental genomes. Hypermutator F₁ progeny are indicated with asterisks, and the *ZNF3* locus is indicated in each strain with a red arrowhead. Regions enlarged below illustrate Cnl1 subtelomeric arrays on several chromosomes and depict examples of Cnl1 array expansion (e.g. Chromosome 4, P18), contraction (e.g. Chromosome 1, P14), and invasion of naïve H99 subtelomeres (e.g. Chromosome 1, P8). Telomeric repeat sequences are indicated by black half circles only in the enlarged panels.

Surprisingly, the genome assemblies for the three progeny with the functional *ZNF3* allele inherited from H99 appear to encode more full-length Cnl1 elements and fragments compared to the three *znf3* progeny. However, of the three *znf3* progeny, telomeric repeat

sequences were only identified at the end of two chromosomes (out of a possible 84 telomeric ends across the three progeny, 2/84 or 2%): on one arm of Chromosome 3 in P8 and one end of Chromosome 12 in P34 (Figure S12). This is in contrast to the 31 telomeres that were accurately assembled across the three *ZNF3* progeny (31/84 or 37%) (Figure S12). The smaller number of telomeres identified in the *znf3* progeny assemblies suggest there might be more Cnl1 elements that were not accurately included, similar to the assemblies for the hypermutators Bt65 and Bt81. Therefore, the Cnl1 burden quantified and presented in Table 1 might not accurately capture the entire Cnl1 burden of these strains.

A previous study found *ZNF3* to be a haploinsufficient gene because no progeny isolated from a *ZNF3* x *znf3*Δ cross showed evidence of sex-induced RNAi-mediated silencing³². This haploinsufficiency allowed us to analyze Cnl1 dynamics both hypermutator and non-hypermutator progeny and identified two additional phenomena. First, nearly all Cnl1 arrays in the progeny show signs of expansion and contraction relative to the Bt65 parental genome, suggesting these elements are either highly mobile during sexual reproduction, undergoing high levels of recombination, or both. Additionally, combined analysis of subtelomeric region inheritance patterns and Cnl1 arrays revealed Cnl1 elements are capable of invading naïve subtelomeric regions inherited from the H99 parent, i.e. regions that previously had no Cnl1 elements/fragments (Figure 4, Figure S12). In the three *znf3* progeny, 65% (28/43) of the naïve telomeric regions inherited from H99 acquired Cnl1 copies and arrays in many cases. In the three *ZNF3* progeny, 81% (35/44) of naïve telomeric regions inherited from H99 now had Cnl1 elements. Overall, both *ZNF3* and *znf3* F₁ progeny inherited roughly equivalent numbers of telomeric regions from either parent and Cnl1 invaded a majority of the naïve H99 telomeres.

Complementation of the nonsense mutation in *ZNF3* significantly lowers the mutation rate and restores production of siRNAs

All of the available evidence thus far suggested the nonsense mutation in *ZNF3* unique to Bt65 and Bt81 is responsible for the hypermutation phenotype, possibly due to compromised RNAi silencing of Cnl1 elements. To test this hypothesis, we used CRISPR-mediated gene editing to restore the functional *ZNF3* allele in Bt65. Gene editing was achieved with the transient CRISPR-Cas9 coupled with electroporation (TRACE) system and the utilization of a functional *ZNF3* allele from a closely related strain, Bt133, such that only the SNP responsible for the nonsense mutation would be changed to the wild-type nucleotide (found in H99 and all SDC isolates except Bt65 and Bt81)⁴⁸. Following transformation and selection, we identified two Bt65 transformants that had successfully integrated a single copy of the Bt133 *ZNF3* allele at the endogenous *znf3* locus, Bt65+*ZNF3*-1 and Bt65+*ZNF3*-2. These two independent Bt65+*ZNF3* transformants were subjected to fluctuation analysis to determine if changing the single SNP responsible for the *znf3* nonsense mutation restored the mutation rate to a wild-type level. On YPD + rapamycin + FK506 medium, both transformants had significantly lower mutation rates than Bt65, similar to those observed in H99 and Bt133 (Figure 5A).

We next sequenced the sRNA repertoires of the Bt65+*ZNF3* isolates as well as Bt65, Bt81, and an H99 *znf3*Δ mutant; H99 and an H99 *rdp1*Δ mutant served as positive and negative controls, respectively. Analysis of the size distribution of sRNAs aligning to the *C. neoformans* genome showed that Bt65, Bt81, and H99 *znf3*Δ displayed profiles similar to that of the canonical RNAi mutant H99 *rdp1*Δ because they lacked the characteristic siRNA peak at 21-24 nt (Figure 5B). sRNA analysis also revealed that *ZNF3* complementation in Bt65 restored the 21-24 nt sRNA peak. We also characterized the 5' nucleotide identity of sRNAs of different lengths

Figure 5. *ZNF3* complementation in Bt65 significantly reduces mutation rates and restores siRNA production. (A) Mutation rates of H99, Bt65, Bt133, and the two independent *ZNF3* complementation mutants, Bt65 + *ZNF3-1* and Bt65 + *ZNF3-2*, on YPD + rapamycin + FK506 medium. Bars represent mutation rate (number of mutations per cell per generation) and error bars represent 95% confidence intervals. (B) Size distributions of sRNA reads from each indicated strain. Dashed vertical lines indicate the 21 to 24 nucleotide size range, the characteristic sizes of siRNAs produced by the RNAi pathway. (C) Proportion of sRNA reads (y-axis) with the indicated 5' nucleotide identity (color of stacked bar) at each sRNA read size (x-axis). siRNAs produced by the RNAi pathway characteristically have a 5' uracil nucleotide. (D) Quantification of sense and antisense sRNAs from Bt65, Bt65 + *ZNF3-1*, and Bt65 + *ZNF3-2* aligning to an array of subtelomeric Cn1 elements on Chromosome 1 of Bt65. Transposable elements along the chromosome are indicated by dark grey boxes, while intergenic regions are light grey.

379 and found only H99 and the two Bt65+*ZNF3* transformants had a peak of 21-24 nt sRNAs with a
380 predominance for a 5' U, another characteristic of siRNAs produced by the RNAi pathway

(Figure 5C).

To determine how complementation of *ZNF3*, and thus restoration of RNAi, specifically impacted silencing of *Cnl1*, we quantified sRNAs aligning to *Cnl1* elements across the Bt65 genome (Table S4). Relative to Bt65, the normalized expression of sRNAs corresponding to *Cnl1* were increased 5.6-fold in H99 (a strain with no full-length *Cnl1* elements and only 22 *Cnl1* fragments), 10.6-fold in Bt65+*ZNF3-1*, and 12.8-fold in Bt65+*ZNF3-2* (Table S4). To demonstrate the marked difference in *Cnl1* sRNAs in the two Bt65+*ZNF3* transformants compared to Bt65, sRNAs were plotted along a telomeric *Cnl1* array on Chromosome 1 in Bt65 (Figure 5D). These results suggest that changing the single nucleotide responsible for the nonsense mutation in *ZNF3* back to the wild-type nucleotide found in closely related strains as well as H99 was able to successfully restore RNAi in Bt65. This result also provides evidence that all other RNAi components have been largely maintained in a functional form despite the loss of RNAi activity in Bt65.

Discussion

Transposable element mobilization can alter gene expression, gene function, and even genomic stability. In this study, we identified two clinical, hypermutator isolates of *C. neoformans*, Bt65 and Bt81, with a massive accumulation of the non-LTR retrotransposon Cn11 at all but one subtelomeric loci of all fourteen chromosomes. Furthermore, the Cn11 element was capable of inserting in non-subtelomeric regions of the genome, resulting in resistance to diverse classes of antifungal drugs. These findings were unprecedented as *C. neoformans* is thought to be an RNAi-proficient species^{13,49}. Additionally, *C. neoformans* isolates were also thought to lack full-length copies of Cn11, and only the sister species *C. deneoformans* has been shown to harbor full-length Cn11 elements capable of mobilization^{13,27}. These findings highlight the importance of intraspecific diversity at both the genotypic and phenotypic levels.

Following isolation, phenotyping, and genotyping of the Bt81a x H99a F₁ progeny, a significantly smaller number of progeny inherited the Bt81 *znf3* allele than expected. These results were surprising giving the approximately 1:1 inheritance pattern of the *ZNF3* allele from either parent in the Bt65a x H99a F₁ progeny. Based on long-read whole-genome sequencing, Bt81 likely has a substantially higher burden of Cn11 than Bt65. Previous studies have shown that expression of transposons, including Cn11, is significantly upregulated during sexual reproduction in RNAi mutants, such as H99 *znf3*Δ^{30,32–34}. The higher Cn11 burden in Bt81 combined with RNAi-deficiency and transposon upregulation during mating could lead to an increased frequency of deleterious Cn11 insertions in progeny lacking *znf3*, and thus a biased inheritance pattern of *ZNF3*. It is also possible that a higher Cn11 burden favored the selection of a suppressor mutation in the Bt81 F₁ progeny. Additionally, *znf3*Δ mutants are the only *C. neoformans* RNAi mutants studied thus far in which progeny from unilateral genetic crosses (i.e.

crosses in which only one parent lacks *ZNF3*) exhibit complete loss of RNAi-mediated silencing³², and thus loss of *ZNF3* results in haploinsufficiency. Overall, the unequal *znf3* inheritance patterns in Bt81 F₁ progeny indicate a sufficiently high burden of Cn11 elements may be deleterious during sexual reproduction.

In Bt65, QTL mapping and genetic complementation demonstrated the hypermutator phenotype was caused by a single SNP in the RNAi gene *ZNF3*. Changing the *znf3* nonsense mutation to the nucleotide found in the laboratory reference strain as well as phylogenetically closely related strains lowered the mutation rate to a wild-type level and restored RNAi function, including expression of siRNAs corresponding to Cn11, thus confirming the role of *Znf3* in Cn11 regulation. Although *ZNF3* complementation restored the mutation rate and siRNA production, the identified QTL which spans Chromosomes 3 and 11 accounted for only 64% of the hypermutator phenotype. This may suggest the existence of additional contributing loci of small effect, such as a mutation in another RNAi component that results in a partial loss of function, although the mapping population used here is under powered to detect such small-effect loci. It is also possible that variation in Cn11 burden across the F₁ progeny accounts for the remainder of the genetic loci that contribute to the hypermutator phenotype. However, because short-read sequencing data was utilized in variant calling, these Cn11 loci may have been difficult to map in the F₁ progeny or failed to pass quality criteria and were subsequently filtered out during preprocessing of the genetic data used in QTL analysis.

Despite having the wild-type nucleotide in the first exon of *ZNF3*, where Bt65 and Bt81 have a nonsense mutation, both Bt89 and Bt133 (two of the most closely related strains) have a substantial accumulation of subtelomeric Cn11 arrays relative to the laboratory reference strain H99, which has none. The considerable Cn11 burden in Bt89 and Bt133 indicates that they may

descend from an ancestral strain that had lost RNAi function, but which was subsequently restored, such as through a genetic cross in which they inherited a wild-type *ZNF3* allele. The bias against *znf3* inheritance in Bt81 F₁ progeny and the lower but still impressive Cnl1 burden in Bt89 and Bt133 potentially illustrate a natural example of how *C. neoformans* genomes have struck a balance in their mutational capability, switching between high mutational capacities during times of RNAi loss, and genomic stability when RNAi is restored, all via one SNP in only one key RNAi gene. The expansion and contraction of Cnl1 arrays as well as the ability of Cnl1 to invade naïve subtelomeric regions inherited from the H99 parent in the Bt65 F₁ progeny genomes was also exceptional. The observed Cnl1 subtelomeric dynamics mirror those observed for the MoTeR transposons of the fungal plant pathogen *Magnaporthe oryzae*, which were also recently shown to localize to highly dynamic subtelomeric arrays⁵⁰. It is possible the Cnl1 subtelomeric arrays identified here could eventually overcome the requirement for telomerase, as in the example of *Drosophila* telomeres, in which the functions of telomerase have been supplanted by a telomeric retrotransposon^{51,52}.

The finding that only a single SNP rendered the RNAi pathway non-functional in Bt65, and that no additional obvious mutations had occurred in other genes that might disrupt RNAi function, suggest Bt65 represents an intermediate evolutionary stage and could illustrate the natural consequences of relatively recent RNAi loss. Further analysis of the consequences of RNAi loss in Bt65 (and potentially Bt81) through experimental evolution or gene regulation analyses could shed light on the short-term consequences of RNAi loss at the genomic and phenotypic levels. Bt65 could thus serve as an interesting intermediate comparator between RNAi-proficient *C. neoformans* isolates and the closely related RNAi-deficient species *C. deuterogattii*^{7,32}. Instances of relatively recent loss of RNAi have also been observed in a natural

Caenorhabditis elegans isolate, which has a large deletion in a RIG-I homolog required for RNAi and was shown to be infected with an RNA virus^{53,54}. Unlike the identified *C. elegans* virus-infected strain and several other RNAi-deficient fungal species, such as *Saccharomyces cerevisiae*, *Ustilago maydis*, and several *Malassezia* species, we were unable to identify a dsRNA virus in either of the two *C. neoformans* hypermutator strains identified here (see Materials and Methods)^{49,55}. It is possible though, that the hypermutators harbor other types of mycoviruses (ssRNA, for example) that we were unable to detect through our approaches or that the mycovirus was cured by common microbiological isolation practices⁵⁵.

The identification of this hypermutator phenotype in natural *C. neoformans* clinical isolates has important implications for antifungal drug resistance and potentially other adaptive consequences. Here, we showed Cnl1 insertion could confer resistance to diverse classes of antifungal drugs, including the clinically relevant antifungal 5-FC. Insertion of Cnl1 into other genes, particularly those involved in the sterol biosynthesis pathway, could confer resistance to amphotericin B and fluconazole, the only other antifungal drugs effective for *C. neoformans* treatment^{23,56}. This mechanism of drug resistance also has interesting implications for a novel antifungal approach that utilizes dsRNA to initiate RNAi silencing in fungal plant pathogens⁵⁷. The effects of Cnl1 insertion at non-coding loci, such as promoters and 3' untranslated regions, could also impact overall genomic stability or alter gene expression to have important phenotypic implications for virulence, similar to the effects of the *Ac/Ds* elements of maize, the first transposable elements discovered⁵⁸. Alterations in gene expression might also confer resistance to drugs for which resistance cannot be gained through loss of function mutations. Even if full resistance isn't acquired, altered gene expression could contribute to antifungal drug

tolerance, like the tolerance observed in *Candida albicans*, which contributes to persistent infections in immunocompetent patients^{59,60}.

At this stage it is difficult to know how Cnl1 transposition and accumulation may be selected for or against over time. The subtelomeric arrays identified in Bt65 and Bt81 may undergo cycles of amplification and recombination-mediated contraction allowing them to exploit Cnl1 mutagenesis when under stress, similar to the retrotransposons replication cycles observed in some plants^{61,62}. Maintaining an RNAi-deficient background could also be adaptive in the context of viral infection, as has been shown in yeast harboring the killer virus, which outcompete neighboring uninfected strains, and in mice harboring latent herpesvirus, which are protected from the bacterial pathogens *Listeria monocytogenes* and *Yersinia pestis*^{63,64}. Conversely, the mutational impact of Cnl1 mobilization could be highly deleterious over the long term and therefore may not represent a massive contributor to the rise of drug resistance. Natural selection could either select for reversion to a functional RNAi-pathway through mutation of the *znf3* nonsense mutation or instead preserve loss of RNAi and eliminate all full-length transposable elements, as has been observed in *C. deuterogattii*²⁸. Future research on the potential for Cnl1 insertion to mediate resistance to amphotericin B and fluconazole, and the impact of hypermutation due to Cnl1 mobilization on *in vivo* drug resistance, adaptive potential, and genomic stability over time will be of great interest.

Materials and Methods

Strains and growth

The *C. neoformans* strains described in this study are listed in Table S5. Strains were stored at -80°C in liquid yeast extract peptone dextrose (YPD) supplemented with 15% glycerol. Strains were inoculated on YPD agar plates, grown for three days at 30°C, and maintained at 4°C. Due to the hypermutator phenotypes associated with several of the strains in this study, strains were not maintained on YPD agar plates for routine use for more than two weeks; fresh cells from frozen glycerol stocks were inoculated to YPD agar plates as needed.

Screening for hypermutator candidates

Assays for the emergence of resistance (papillation assays) were conducted as previously described⁶⁵. In brief, ten independent overnights per strain were grown overnight at standard laboratory conditions in 5mL liquid YPD medium. Cultures were then spun down, washed, and concentrated in 2 mL dH₂O. Each culture was swabbed to a quadrant of either YPD + 100 ng/mL rapamycin + 1 µg/mL FK506 agar medium or YNB + 100 µg/mL 5-fluorocytosine agar medium. YPD + rapamycin + FK506 plates and YNB + 5-fluorocytosine plates were incubated for up to seven days at 37°C and 30°C, respectively. Fisher's exact probability test was used to determine if the associations environmental isolates and the hypermutator phenotype was statistically significant using the VassarStats online software (<http://vassarstats.net>).

Fluctuation assays

Fluctuation assays were conducted as previously described⁶⁵. Briefly, ten independent overnights of each strain were grown overnight in 5 mL liquid YPD medium at 30°C. Cultures

were washed three times and resuspended in dH₂O. Cells were then plated to the appropriate medium (100 μ L 10⁻⁵ cells on YPD, 100 μ L 10⁻² cells on YNB + 5-FC, and 100 μ L undiluted cells on YPD + rapamycin + FK506 and YNB + 5-FOA). Mutation rates and 95% confidence intervals were calculated using the FluCalc program⁶⁶. For the increased temperature fluctuation analysis, strains were grown overnight at either 30°C or 37°C before use in fluctuation assays, as indicated. YPD + rapamycin + FK506 plates were incubated and 37°C; all other media was incubated at 30°C. Mutation rates and confidence intervals for all fluctuation assays in this study are provided in Table S6.

Characterizing mutation spectra

Following selection on antifungal drug media, resistant colonies were streak purified to YPD medium. Genomic DNA was isolated from the purified colonies, and genes in which mutations are known to cause resistance to the corresponding antifungal drug were PCR amplified (*URA5* and *URA3* for 5-FOA-resistant colonies^{44,45}, *FRR1* for rapamycin+FK506-resistant colonies^{41,42}, and *FUR1*, and *UXS1* for 5-FC-resistant colonies⁴⁶). Oligonucleotides used for all PCR reactions in this study are listed in Table S7. PCR products were subjected to gel electrophoresis, products of interest were extracted from agarose gels using a QIAgen gel extraction kit and sequenced through classical Sanger sequencing conducted at Genewiz. Sequenced mutations, including transposon insertions, were characterized with both Sequencher software and the Clustal Omega Multiple Sequence Alignment program⁶⁷. Identified transposon insertion sequences in *FRR1*, *URA5*, and *FUR1* are listed in Table S8.

Illumina sequencing

Single colonies from strains for whole-genome Illumina sequencing were inoculated in 50 mL of liquid YPD medium and grown overnight at 30°C, shaking. Cells were collected and lyophilized as previously described⁶⁵, and high molecular weight DNA was isolated following the CTAB protocol as previously described⁶⁸. Strains were barcoded and sequencing libraries were generated with the Kapa HyperPlus library kit for 300bp inserts, pooled, and sequenced using paired-end, 2 x 150bp reads on an Illumina HiSeq 4000 platform at the Duke University Sequencing and Genomic Technologies Core facility.

Generation of F₁ progeny

Bt65a x H99a *crgI*Δ and Bt81a x H99a *crgI*Δ F₁ progeny were generated by genetically crossing either Bt65 or Bt81 with H99 *crgI*Δ on Murashige Skoog (MS) medium (Sigma) following Basic Protocol 1 as described in Sun et al. 2019⁶⁹. Basidiospores were randomly isolated through microdissection after three weeks of incubation on MS following Basic Protocol 2 as described in Sun et al. 2019⁶⁹.

Nanopore sequencing and genome assemblies

The DNA samples for nanopore sequencing were isolated and purified using the CTAB DNA preparation protocol described previously⁷⁰. The size estimation of the obtained DNA was done using PFGE electrophoresis and quality was determined using NanoDrop. Once the high-quality DNA was obtained, sequencing was performed using the MinION device with the MinKNOW interface. During sequencing, Bt65, Bt89, and Bt133 were multiplexed together whereas six of the Bt65a x H99a progeny were multiplexed for a second sequencing run. For

multiplexing, samples were barcoded using EXP-NBD103/EXP-NBD104 kits and libraries were made using SQK-LSK109 kit as per the manufacturer's instructions. The libraries generated were sequenced on R9.4.1 flow cell and reads were obtained in .fast5 format. These reads were then converted to fastq format using Guppy_basecaller (v 4.2.2_linux64). The reads were de-multiplexed using qcat (<https://github.com/nanoporetech/qcat>) or Guppy_barcode (part of Guppy_basecaller) with barcode trimming option during processing. Bt81 nanopore sequencing was done as a standalone sample using an R9 flow cell (FLO-MN106) and basecalling was performed during the run itself.

The sequences obtained for each sample were then assembled via Canu (v2.0 or v2.1.1) to obtain contig-level genome assemblies. For the assembly, only >2 kb long reads were used for the Bt65a x H99 α F₁ progeny and Bt81, whereas >5 kb were used for Bt65, Bt89, and Bt133 genomes. Contigs were then assigned chromosome numbers based on their synteny with the reference genome, H99. The numbering of chromosomes involved in translocations was assigned based on the respective syntenic centromere. Some of the chromosomes were not fully assembled and were broken into multiple contigs (Chr 1, Chr 2 for Bt65, Chr 2, Chr11, Chr14 for Bt89, and Chr 2, Chr 5 for Bt133). For such cases, the respective contigs were joined artificially and then processed by read-mapping to obtain complete collinear chromosomes. Specifically, the contigs were stitched together in orientation as determined based on their synteny. Corrected reads obtained from Canu were then mapped to the respective genomes and duplicated or missing regions from the junction were identified. The chromosome sequence was then corrected accordingly by inserting/correcting/deleting sequences and a full-length chromosome sequence was obtained. Complete resolution of junctions was obtained for Bt65, Bt89, and Bt133 genomes

by this approach. However, some of the Bt65 F₁ progeny chromosomes could not be resolved, probably due to hybrid origin of sequencing reads, and were left with gaps as such.

Once chromosome level genome assemblies were obtained for the Bt65, Bt81, Bt89, and Bt133 genomes, the genome sequences were further processed to improve telomeric and subtelomeric regions. For this purpose, the corrected reads obtained from Canu were mapped back to the respective chromosome-level genomes using minimap2 v2.14. The obtained bam files were then analyzed manually by IGV and consensus or, in a few cases, individual reads (up to 30 kb) representing extra sequence beyond an assembled chromosome were extracted as sam files. These consensus extra sequences were then added onto the chromosome sequences to obtain longer chromosomes. In some cases, read mapping also resulted in the identification of incorrect sequence assembly at subtelomeric regions, and in those cases, the sequence was trimmed until a consensus sequence was observed at the end of the chromosome. Once these corrections were made, the genome assemblies were polished via one round of nanopolish and five rounds of pilon, except for the Bt81 genome, for which only 5X pilon polishing was performed. As a result of these corrections and polishing, final assemblies were obtained for each of the four isolates and are described in the study. For the Bt65 F₁ progeny genome assemblies, the subtelomeric extension/curation was not performed, but they were polished using both nanopolish and 5X pilon.

Centromere, telomere, and Cn11 mapping

Centromeres in Bt65, Bt81, Bt89, and Bt133 were defined based on their synteny with the reference H99 genome (genome assembly ASM301198v1) The final polished genomes were used and centromere locations were identified by BLASTn analysis using H99 centromere-

flanking genes as query sequences. Once the centromere locations were defined, Tcn1-Tcn6 locations within those regions were mapped by BLASTn analysis. For the representation, only BLAST hits longer than 400 bp were mapped. For the overlapping BLAST hits with multiple Tcn elements, the longest and best BLAST result was used, and the rest of the matches were discarded from further analysis. All the hits were then visualized using Geneious Prime and maps were exported as .svg files, which were then processed using Adobe Illustrator.

For the Cnl1 mapping at the subtelomeres, the longest *CNLI* insertion sequence from the Bt65 genome was used as the query sequence and BLASTn was performed against each genome. BLAST hits longer than 50 bp were mapped to the respective genomes and visualized using Geneious Prime where the hits were color-coded based on their lengths. The zoomed views for these maps were then exported as .svg files, processed using Adobe Illustrator, and combined with centromere Tcn mapping analysis to generate final figures.

RepeatMasker was used to annotate all transposons in the *de novo* genome assembly of Bt65. For this purpose, RepeatMasker (v4.0.7) with Dfam (v3.3) and RepBaseRepeatMaskerEdition-20181026 libraries was used, supplemented with RepBase EMBL database (v26.04)^{71–73}. The “-species fungi” option was used to identify all repeats in the genome and provided additional support for the manual Tcn and Cnl1 mapping.

Syntenic maps

Syntenic comparisons between the genomes were performed using SyMAP v4.2 with the H99 genome as the reference (genome assembly ASM301198v1). The syntenic comparison was conducted using default parameters and syntenic block maps were exported as .svg files. The maps were processed using Adobe Illustrator for visualization. The phylogenetic relationship as

depicted in Figure S6 was drawn based on the earlier representation¹⁸. The telomere and centromere locations were marked manually based on the presence of the telomere repeat sequence and Tcn mapping, respectively.

For the centromere comparisons, all centromere sequences along with Tcn annotations were converted into GenBank format. The files were then used for synteny comparison via EasyFig v2.2.3. The maps were exported as .svg files which were processed in Adobe Illustrator.

Recombination maps for Bt65 x H99 F₁ progeny

Six of the Bt65a x H99α F₁ progeny were sequenced with on the nanopore MinION sequencing platform and their genomes were assembled and polished using the methods described above. Once their genomes were assembled, recombination maps were generated by mapping the Illumina sequence data from the parental strains to each of the progeny genomes. For this purpose, both H99 and Bt65 Illumina reads were used from published datasets (SRR642222 and SRR647805 for H99; SRR836876, SRR836877, SRR836878, SRR836880, SRR836884, and SRR836885 for Bt65). Reads from all runs were merged to obtain a single file for both H99 and Bt65. The reads were then mapped to the progeny genomes using Geneious Prime default mapper with three iterations. Variants with 90X coverage and at least 90% variant frequency were called from these mapped files. These variants along with coverage analysis were then used to identify recombination sites and generate recombination maps. Cn11 mapping for each of progeny genome was performed as described above. The location of *ZNF3* in each genome was identified by BLASTn analysis using H99 *ZNF3* (CNAG_02700) as the query sequence.

Genetic variant calling and segregant filtering

Whole-genome sequencing data of 28 F₁ progeny from the Bt65a x H99a *crg1*Δ cross were aligned via BWA (v0.7.12-r1039)⁷⁴ to an H99 reference genome (downloaded from FungiDB [<http://fungidb.org/fungidb/>] on April 15th, 2020; FungiDB-46_CneoformansH99_Genome.fasta) and genetic variants between Bt65 and H99 were called using SAMtools (v0.1.19-96b5f22941)⁷⁵ and FreeBayes (v1.2.0)⁷⁶. Approximately 300,000 raw genetic variants were identified across the segregants. The genotypic correlation between progeny, the read coverage per genetic variant, and the ratio of reads suggesting the H99 vs. Bt65 allele per variant were monitored across the genome to identify clones, progeny with aneuploid genomes, and heterozygotic diploids (respectively). Two pairs of clones were identified (Supplementary Table S9) and one segregant from each pair was retained for analysis. F₁ progeny 25 was identified as a heterozygotic diploid (Supplementary Figure S13) and removed from initial analysis. Instances of aneuploidy (and partial duplications) are observed along Chromosomes 3, 4, 11, and 13 within six segregants from this cross and for initial filtering and analysis, those with heterozygotic aneuploidy were removed from analysis (Supplementary Table S9).

Genetic variant filtering

After removing clones and samples with aneuploidy or diploidy, raw genetic variants were filtered by limiting sites to bi-allelic SNPs, called across all the progeny (100% call rate), with greater than 10X read coverage (and a maximum of 200X), a minor allele frequency of 5%, and a quality score greater than 4 (and less than 5.4). These filtering criteria were selected after examining the bivariate relationships between allele frequency, read depth, and quality scores per

chromosome (Supplementary Figure S14A and 14B). Genetic variant sites were also removed if within one kb of the centromere along a given chromosome²⁸. After filtering, a total of 215,411 bi-allelic SNPs were retained for further analysis. The median distance between contiguous SNP sites is 45 bp, and less than 0.01% of neighboring sites had a distance larger than two kb. The allele frequencies across the genome ranged between 25 and 75% of segregant with the Bt65 allele, except for a large portion of Chromosome 13, between 0 to 500 kb, where over 88% of segregants inherited the Bt65 allele (Figure S14C). With these data, a Poisson regression (methods described in Roth *et al.* 2018⁷⁷) was used to relate the average number of crossovers across F₁ progeny as a function of chromosome size. With this model, the estimated genome-wide, physical-to-genetic distance in this cross is 8.14 kb/cM.

QTL mapping

For use in association tests, across 24 F₁ progeny and the two parental strains, the 215,411 bi-allelic SNPs were collapsed into 1,237 unique haplogroups made up of genetic variants sites that co-segregated within the segregant genomes, such that, between any two haplogroups, at least one segregant contains a change in allele (i.e. a recombination event between the Bt65a and H99a *crg*/Δ genomes). Collapsing genetic variants into haplogroups reduces the number of repeated tests in association mapping and computational costs⁷⁸. Across the 1,237 haplogroups, the mutation rate on rapamycin + FK506 medium was used as the phenotype and tested for association using a Kruskal-Wallis H-test. The $-\log_{10}(p\text{-value})$ from this test across haplogroups was monitored to identify QTL. Significance thresholds were established via 10,000 permutations with an $\alpha = 0.01$ as described in Churchill and Doerge (1994)⁷⁹, and 95% confidence intervals for the QTL locations were generated as described in

Visscher *et al.* (1996)⁸⁰. The heritability at the peak of identified QTL was estimated using linear regression with the model: $M = \mu + \beta I + e$, where M is the mutation rate $\times 10^7$, e is an error term, μ is the mean mutation rate $\times 10^7$, I is an indicator variable for the allele at the QTL peak – coded as 0 if from H99 α *crgI* Δ or 1 if from Bt65 α – and β is the effect of having the H99 α *crgI* Δ vs. the Bt65 α allele at the QTL. The variation explained (R^2) from this model was used as an estimate of the heritability.

Gene annotation and SNP effect prediction

For genes within the identified QTL spanning Chromosomes 3 and 11, the alleles between H99 and Bt65 were imputed using filtered SNP data (described above). The published H99 reference strain annotation (downloaded from FungiDB [<http://fungidb.org/fungidb/>] on April 15th, 2020; FungiDB-46_CneoformansH99.gff) was used to predict changes in protein sequence between the H99 and Bt65 parental backgrounds.

CRISPR-mediated genetic complementation of *ZNF3* in Bt65

To change the single nucleotide responsible for the nonsense mutation in the first exon of *ZNF3*, a thymine (base 976004 of H99 Chromosome 3 (CNA3 assembly, accession GCA_000149245.3)), to the wild-type cytosine found in H99 and other phylogenetically closely related strains, the Transient CRISPR-Cas9 Coupled with Electroporation (TRACE) system was used⁴⁸. Briefly, the gene encoding Cas9 was PCR amplified from plasmid pXL1-CAS9-HYG. The *SH1-NEO* construct encoding *NEO* (G418 resistance) targeted to a safe haven locus (SH1) was amplified from plasmid pSDMA57⁸¹. For the sgRNA expression construct, the U6 promoter was amplified from XL280 α gDNA, and the sgRNA scaffold was amplified from plasmid

pYF515⁸². Overlap PCR was used to generate the sgRNA construct with the U6 promoter and sequences targeting either SH1 or the nonsense mutation in *ZNF3*. A 2,171bp region was amplified from Bt133, containing the wild-type C nucleotide in *ZNF3* exon 1 and no other mutations relative to Bt65 for integration at the *ZNF3* endogenous locus in Bt65 (1,197bp upstream of the *ZNF3* start codon to 971bp after the start codon). The *SHI-NEO* construct was linearized with the *AscI* restriction enzyme (NEB). 2μg of the Bt133 *ZNF3* recombination template, 2μg of the *SHI-NEO* linearized construct, 250ng of the *ZNF3* gRNA, 250ng of the SH1 gRNA, and 1.5μg of Cas9 DNA were transformed simultaneously into Bt65 via electroporation using a BIO-RAD Gene Pulser.

Transformants were selected on YPD + G418 agar plates. Successful transformants were identified through restriction enzyme digest with *BtsI*-v2 (NEB), which cleaves the first exon of Bt65 *znf3* at the nonsense mutation but does not cleave the first exon of *ZNF3* in Bt133 (or H99) (Figure S15A). PCR was also used to ensure that no transformants had integrated copies of the gene encoding Cas9 or the gRNA constructs and that only a single Bt133 *ZNF3* allele had integrated correctly at the endogenous *znf3* locus (Figure S15B-F). Sanger sequencing was used to further confirm correct replacement of the Bt65 *znf3* allele including the nonsense mutation with the Bt133 *ZNF3* allele. No identified Bt65+*ZNF3* transformants also had a stably integrated copy of the *NEO* gene at the safe haven locus.

sRNA isolation and sequencing

C. neoformans cells were grown overnight in 50 mL YPD medium at standard laboratory conditions. Following culture, cells were spun down, supernatant was removed, and cells were frozen at -80°C for one hour. Cells were then freeze dried with a Labconco Freezone 4.5

lyophilizer overnight. 70 mg of lyophilized material was used for sRNA isolation following the mirVana miRNA Isolation Kit manufacturer's instructions. sRNA was quantified with a Qubit 3 Fluorometer and quality was verified with an Agilent Bioanalyzer using an Agilent Small RNA Kit. sRNA libraries were prepared with a QiaPrep miRNA Library Prep Kit and 1 x 75 bp reads were sequenced on the Illumina NextSeq 500 System at the Duke University Sequencing and Genomic Technologies Core facility.

sRNA data processing

Initial quality control of the small RNA libraries was performed with FastQC 0.11.9⁸³ followed by the removal of QIAseq library adapters (5': GTTCAGAGTTCTACAGTCCGACGATC; 3': AACTGTAGGCACCATCAAT) with cutadapt 2.8⁸⁴. All untrimmed reads or reads smaller than 14 nt were discarded. The surviving trimmed reads were mapped with bowtie v1.2.3⁸⁵ to the *C. neoformans* Bt65 genome, allowing multiple alignments but no mismatches. The resulting SAM files were converted into BAM file format with SAMtools 1.9⁷⁵ and feature read counts of transposable elements were calculated with BEDTools⁸⁶ using the 'intersect -wa' option and the annotations of transposable elements, which were identified with RepeatMasker using the repbase database for *C. neoformans*^{71,72}. Normalization of the read counts to reads per million (RPM) was performed, allowing the comparison of the libraries. Furthermore, the read depth on both DNA strands was analyzed with SAMtools and custom made perl scripts were used to calculate the read size distribution and 5'-nucleotide preference of the small RNA reads as previously described^{87,88}.

Double-stranded RNA enrichment

For dsRNA enrichment, *Cryptococcus* cells were grown overnight in 50mL liquid YPD medium at 30°C. RNA was extracted, and dsRNA was enriched as previously described⁵⁵. dsRNA enrichment in H99, Bt65, and Bt81 did not reveal the presence of any large dsRNA segments (Figure S16).

Data and software availability

Genetic variant filtering, QTL mapping, and SNP effect prediction was conducted in python (anaconda 3.7.3) via custom scripts; hosted at: https://github.com/magwenelab/Hypermutator_QTL . All sequencing data is available under BioProject PRJNA749953.

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Supplementary Figures

Supplementary Figure S1. Bt65 and Bt81 do not display a hypermutator phenotype on 5-FC or 5-FOA. Mutation rates of closely related VNBII strains and controls on **(A)** YNB + 5-FC and **(B)** YNB + 5-FOA media. Bars represent the mutation rate and error bars represent 95% confidence intervals; mutation rates represent the number of mutations per cell per generation. Schematic depicts the phylogenetic relationships of all strains included in fluctuation analyses based on Desjardins et al. 2017¹⁸.

Supplementary Figure S2. Growth at elevated temperature does not result in increased mutation rates in *C. neoformans* strains. Fluctuation assays were used to quantify the mutation rates of strains grown overnight at 30°C or 37°C and plated on YPD + rapamycin + FK506 medium. Bars indicate mean mutation rate and error bars indicate 95% confidence intervals. Mutation rates represent the number of mutations per cell per generation.

Supplementary Figure S3. Gel electrophoresis *FRR1*, *URA5*, and *FUR1* PCR products

from resistant colonies. Gel electrophoresis of *FRR1* PCR products from (A) all H99 rapamycin + FK506-resistant colonies and a subset of (B) Bt65 and (C) Bt81 rapamycin + FK506-resistant colonies sequenced in Figure 1D. PCR amplification of wild-type *FRR1* in *C. neoformans* produces a 1,165 bp electrophoretic species (primers ZC7/8). Gel electrophoresis of a subset of (D) *URA5* PCR products from H99, Bt65, and Bt81 5-FOA-resistant colonies and (E) *FUR1* PCR products from 5-FC-resistant colonies of Bt65 and Bt81.

Supplementary Figure S4. Protein length differences of genes within QTL. In the upper panels, points mark the strength of association (y-axis) between bi-allelic SNP sites and hypermutation for Chromosome 3 and Chromosome 11 (top left and right, respectively). Grey dashed lines depict the 95% confidence intervals (CI) of the two QTL. Lower panels show the predicted differences in lengths of proteins (y-axis) encoded by annotated genes in Bt65 compared to H99 within each 95% CI of the QTL (x-axis) on Chromosome 3 and Chromosome 11 (bottom left and right, respectively). The name of each gene with a predicted nonsense mutation is annotated. Blue and red colors denote the gene orientation.

Supplementary Figure S5. Haplotype maps of Bt65 x H99 F₁ progeny utilized for QTL mapping. For the QTLs on Chromosome 3 and Chromosome 11 (left and right, respectively) the haplotypes (x-axis) are inferred by SNP data per segregant (y-axis) and colored blue or orange if inherited from H99 *crg1Δ* or Bt65, respectively. Segregants are sorted along the y-axis by the quantification of their mutation rate; largest to smallest, top to bottom. Vertical red lines display the boundaries of the QTL(s). Vertical black lines depict the approximate location of the translocation between H99 and Bt65. The boundaries of the QTG, *ZNF3*, are depicted by vertical green lines. Vertical white spaces indicate the approximate locations of the centromeres.

Supplementary Figure S6. Large-scale genomic rearrangements between the H99, Bt65, Bt81, Bt89, and Bt133 genomes. Nanopore whole-genome sequencing followed by synteny analysis was used to identify all indicated genomic rearrangements with respect to the reference strain H99. There is a chromosomal translocation between Chromosomes 3 and 11 that is unique to H99, and a translocation between H99 Chromosomes 1 and 13 that is unique to Bt65 and Bt81. The phylogenetic relationships of these strains are depicted in the top schematic, telomeric repeat sequences accurately identified in the genomic assemblies are indicated by black half circles, and centromeres are indicated by white circles.

Supplementary Figure S7. CNAG_01836 gene model. A grey horizontal bar depicts the entire gene body in the upper panel, and larger grey rectangles show locations of exons. The gene is depicted 5' to 3', left to right, and is 1896 nt in length. The locations of SNPs differing between Bt65 and H99 are shown by vertical black rungs along the gene model. Amino acids specified by mRNA codons in the indicated region of CNAG_01836 Exon 7 (nucleotide 1831 to 1848) are shown to illustrate the G to A mutation (nucleotide 1842) predicted to cause an early nonsense mutation in Bt65. The bottom panel depicts the predicted outcome of the nonsense mutation on the protein encoded by CNAG_01836 in Bt65.

Supplementary Figure S8. Mutation rates of Bt81 x H99 F₁ progeny. Fluctuation analysis was used to quantify the mutation rates of the indicated strains on YPD + rapamycin + FK506 medium (y-axis) – sorted smallest to largest, left to right – for F₁ progeny and the parental strains, H99α *crg1Δ* and Bt81 (x-axis). Bars indicate the mean mutation rate and error bars represent 95% confidence intervals. Mutation rates represent the number of mutations per cell per generation. Inheritance of the Bt81 *znf3* allele or H99 *crg1Δ* *ZNF3* allele in the F₁ progeny is indicated above mutation rates.

Supplementary Figure S9. Subtelomeric and centromeric retrotransposons in *Bt89* and *Bt133*. Distributions of the Tcn1-Tcn6 LTR-retrotransposons and the Cn1 non-LTR retrotransposon in the genomes of (A) *Bt89* and (B) *Bt133*. 50 kb of subtelomeric regions as well as centromeric regions are displayed for both strains. Shading corresponds to the lengths of the Cn1 elements, and gene arrowheads indicate the direction of transcription for all retrotransposons.

Supplementary Figure S10. Centromere lengths do not significantly differ among H99, Bt65, Bt81, Bt89, and Bt133. The length of each centromere (y-axis) is plotted for each strain (x-axis). The thin horizontal black line indicates average centromere length and the thicker black error bars indicate the standard error of the mean. No significant difference was found between the average centromere length of each strain (ANOVA, p -value = 0.153).

Supplementary Figure S11. Centromeric rearrangements among hypermutators and two closely related strains. Shared homology, genomic rearrangements, inversions, insertions and deletions among the centromeres of each of the 14 chromosomes are depicted for Bt65, Bt81, Bt89, and Bt133. The organization of LTR-retrotransposons Tcn1 through Tcn6 are also indicated for each centromere of each strain.

Supplementary Figure S12. Distribution of Cn11 among Bt65 x H99 F₁ progeny and parental strains. The Cn11 non-LTR elements identified in the nanopore-based whole-genome assemblies are depicted for H99, Bt65, three hypermutator F₁ progeny (P02, P08, and P34, all on the left), and three non-hypermutator F₁ progeny (P14, P18, and P20, all on the right). Blue and orange bars under the subtelomeric region of each chromosome indicate which parental strain the region was inherited from (orange for Bt65, blue for H99). Red asterisks indicate invasion of Cn11 into an H99 subtelomeric region that previously had zero Cn11 copies/fragments. Accurate assembly of telomeric repeat sequences at the end of each chromosome is indicated by a black half circle. Cn11 length is also indicated by the shade of black for each element.

Supplementary Figure S13. Genomic diagnostic plot of Bt65 x H99 F₁ progeny 25. For the 14 chromosomes (columns), the log₂ (read depth) (top row) and allelic read depth ratio (bottom row) per genetic variant are shown for the progeny 25. Red and blue colors indicate the prediction of the allele inherited (Bt65 vs. H99, respectively) at each genetic variant. Allelic read depth ratios nearing 0.5 suggest both alleles are present for a given genetic variant. Marked ranges on Chromosomes 3 and 11 log₂ (read depth) plots indicate the significant hypermutator QTL. Black vertical lines depict the boundaries of the centromeres. The boundaries of the *MAT* locus on Chromosome 5 are shown by vertical green lines.

Supplementary Figure S14. Visualization of genetic variant filtering criteria. (A) For the 14 chromosomes (columns), the quality scores of the genetic variants (x-axis) vs. the minimum read depth across the 28 Bt65 x H99 F₁ segregants (y-axis). (B) the quality scores of genetic variants (x-axis) vs. the portion of progeny with the Bt65 allele per genetic variant (y-axis) per chromosome (columns). (C) The portion of progeny with the Bt65 allele per genetic variant (y-axis) across each chromosome (x-axis). The raw genetic variants are shown in black and the filtered SNPs, used in analysis, are shown in red.

Supplementary Figure S15. Identification and confirmation of correct Bt65+ZNF3

transformants. (A) BtsI restriction enzyme digestion of *ZNF3* PCR products from nourseothricin-resistant transformants and controls (primers SJP186/187). **(B)** PCR amplification of the *ZNF3* allele using primers outside of the Bt133 *ZNF3* allele used for homologous recombination (primers SJP208/209). PCR amplification to ensure correct integration of the **(C)** 5' and **(D)** 3' ends of the Bt133 *ZNF3* allele at the endogenous *ZNF3* locus (primers SJP208/187, and SJP186/209, respectively). PCR to ensure neither **(E)** *CAS9* nor **(F)** the gRNA constructs were integrated into the transformants (primers JOHE41657/45812 and JOHE50451/50452, respectively).

Supplementary Figure S16. Enrichment for dsRNA does not identify any fragments likely to be dsRNA mycoviruses. Pictured on the left are RNA samples following LiCl enrichment for dsRNA run on a 1% agarose gel. Total RNA prior to dsRNA enrichment is pictured on the right on a 1% agarose gel. *Ms+* is a *Malassezia sympodialis* strain that harbors a dsRNA virus, and *Ms-* is a congenic virus-cleared strain⁵⁵. Two biological replicates for all samples are shown and labeled (1) and (2). The TriDye 1 kb DNA ladder (NEB) was used to estimate RNA fragment sizes.

1200 **Supplementary Table Legends**

1201 **Supplementary Table S1. Strains included in preliminary screen of SDC isolates for**
 1202 **hypermutation phenotype.**

1203

1204 **Supplementary Table S2. Genetic variants and predicted changes in genes within QTL**
 1205 **between H99 and Bt65.**

1206

1207 **Supplementary Table S3. (A) Centromere lengths in H99, Bt65, Bt81, Bt89, and Bt133, and**
 1208 **(B) one-way ANOVA and Tukey's HSD post hoc statistical tests for differences in mean**
 1209 **centromere length.**

1210

1211 **Supplementary Table S4. sRNA analysis**

1212

1213 **Supplementary Table S5. Strains used in this study.**

1214

1215 **Supplementary Table S6. Mutation rates and 95% confidence intervals for all fluctuation**
 1216 **assays.**

1217

1218 **Supplementary Table S7. Oligonucleotides used in this study.**

1219

1220 **Supplementary Table S8. Cnl1 insertion sequences in PCR products.**

1221

1222 **Supplementary Table S9. Aneuploid, diploid, and clonal Bt65 x H99 F₁ progeny.**