Natural mismatch repair mutations mediate phenotypic diversity and drug resistance in Cryptococcus deuterogattii

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Abstract

Pathogenic microbes confront a constant evolutionary conflict between the pressure to maintain genome stability and the need to adapt to mounting external stresses. Prokaryotes often respond with elevated mutation rates, but to date little evidence exists of stable eukaryotic hypermutators in nature. Whole genome resequencing of the human fungal pathogen *Cryptococcus deuterogattii* identified an outbreak lineage characterized by a nonsense mutation in *MSH2*. This defect in mismatch repair results in a moderate mutation rate increase in typical genes, and a larger increase in genes containing homopolymer runs. This allows facile inactivation of genes with coding homopolymer runs including *FRR1*, which encodes the target of the immunosuppressive antifungal drugs FK506 and rapamycin. Our study identifies a eukaryotic hypermutator lineage spread over two continents and suggests that pathogenic eukaryotic microbes may experience similar selection pressures on mutation rate as bacterial pathogens, particularly during long periods of clonal growth or while expanding into new environments.
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Introduction

Mutation is the raw material of evolution. As a result, all organisms must strike a balance between allowing enough random mutations for selection to act upon, and the fact that most of these mutations are likely to be deleterious and must be purged from the population. This is particularly true for pathogenic microbes that take part in Red Queen conflicts with their hosts and require a continuous supply of mutations to maintain competitive fitness. In this case, increased pressure may exist to maintain an elevated mutation rate to increase the rate of adaptation. Increases in mutation rate may also serve to accelerate adaptation when microbes are introduced into new environments or encounter novel stresses, such as antimicrobial therapy.

Adaptive variation in mutation rate is common in bacteria. One example is the Long Term Evolution Experiment (LTEE), where defects in DNA repair emerged and the resulting hypermutator phenotype swept the population in six out of twelve *E. coli* lines (1). Mutator alleles likely emerge frequently but are typically purged from the population because individual mutations are more likely to be deleterious than adaptive; as a result hypermutators generally produce less fit offspring than non-hypermutators. However, occasionally a sufficiently beneficial mutation is potentiated by the presence of the hypermutator and the hypermutator allele is able to hitchhike to higher frequency within the population. This is more likely to occur in populations at local evolutionary minima, where many different large-effect adaptive mutations are possible. For example, hypermutator phenotypes are common in isolates of *Pseudomonas aeruginosa* growing within the lungs of patients with Cystic Fibrosis, where environmental conditions are constantly changing (2). This changing environment causes an ongoing need to adapt, and increases the likelihood of hypermutators emerging by increasing the target size for adaptive variation. However, once beneficial mutations have fixed, and organisms have adapted to their new environment, antimutator suppressor alleles can emerge to reduce the
mutation rate or defective mutator alleles can even be replaced by functional alleles through horizontal
gene transfer (3, 4). This balance is likely critical to evolution in microbes as they face diverse stresses
and environmental changes, and as a result over 1% of natural bacterial isolates display hypermutator
phenotypes (5). Even more extreme cases also exist, like in *Mycoplasma pneumoniae* and *M.
genitalium*, where the mismatch repair machinery has been completely lost (6).

In fact, many mutator phenotypes are the result of defects in mismatch repair, commonly
referred to as MMR. In bacteria, MMR requires the MutHLS system, where MutS binds to mismatches
and recruits MutL, which subsequently activates MutH to excise the mispair (7–10). In eukaryotes,
both the MutS and MutL families have expanded, yielding MutS homolog (MSH) (11) and MutL
homolog (MLH) families (12). Heterodimers of various MSH family members participate in mismatch
repair, but Msh2 in particular is a core component of the majority of mismatch repair pathways in yeast
(13). As a result, defects in MMR, and *MSH2* mutations in particular, result in elevated rates of simple
mismatches, but also dramatically elevated rates of repeat tract instability (12), which are associated
with hereditary colon cancer in humans. MMR also plays a role in rejecting heteroduplexes of
divergent sequences during both mitosis (14) and meiosis (15), meaning that loss of the complex can
lower species boundaries and allow increased recombination between divergent chromosomes (15, 16).

In contrast to bacteria, substantially less is known about hypermutators in natural populations of
eukaryotes. Evolutionary theory predicts that the presence of sex abrogates selection for hypermutator
strains by eliminating genetic linkage between the mutator allele and the associated beneficial
mutations it hitchhikes upon to high frequency (17). Studies of allele incompatibility within the model
fungus *Saccharomyces cerevisiae* have supported this traditional paradigm. Incompatible alleles of
*PMS1* and *MLH1* exist within the population, such that a cross of two strains with wildtype mutation
frequency can give rise to progeny with elevated mutation rates (18, 19). However, the homozygous
incompatible arrangement was not found among any of the original 65 strains examined, and was only identified within one clinical isolate of 1,010 global diverse yeast isolates (20) and four clinical strains out of 93 sequenced in a separate study (21). This frequency is far below that of the individual alleles in the population, and in each case observed suppressors had arisen to restore a wildtype mutation rate despite the incompatibility. In contrast, eukaryotic mutators can adapt and thrive in clonally expanding populations such as in human tumor environments (22). Recent studies of *Candida glabrata* have begun to challenge this dogma in eukaryotic microorganisms as well, with over 50% of clinical *C. glabrata* isolates harboring loss of function mutations in *MSH2*, a critical component of mismatch repair (23). In previous work we also identified a clade of candidate hypermutators within the Pacific Northwest *Cryptococcus deuterogattii* outbreak that contain a coding single base deletion within the critical mismatch repair component *MSH2* (24).

*Cryptococcus deuterogattii*, previously known as *Cryptococcus gattii* VGII (25), is a basidiomycete human fungal pathogen. Unlike the other species of the *Cryptococcus* pathogenic species complex that infect immunocompromised hosts, *C. deuterogattii* is characterized by its ability to cause infection in otherwise healthy hosts (26) and by loss of the RNAi pathway (27, 28). *C. deuterogattii* is responsible for an ongoing outbreak in the Pacific Northwest region of the United States and Canada, which began in the late 1990’s (29). This outbreak was originally analyzed using Multi-locus sequence typing (MLST) and shown to be comprised of three clonal expansions, denoted VGIIa, VGIIb, and VGIIc (30, 31). These three subtypes differ in terms of virulence and total number of cases, with VGIIa, responsible for the majority of infections. More recently, whole genome sequencing studies of these subpopulations showed that the clonal subtypes appeared to have different proximal geographic origins, and most interestingly, that VGIIa has three very closely related isolates, here referred to as VGIIa-like, that were considerably less virulent and differed in isolation location or
date from the outbreak (24). We identified a candidate potential large effect mutation shared by these
three diminished virulence isolates that resulted in a predicted non-functional Msh2. Here we show that
these VGIIa-like isolates exhibit a bona fide hypermutator phenotype. Furthermore, we show that
homopolymer runs are particularly unstable, and that these runs are common within the coding regions
of genes in the Cryptococcus genome. We predict that this results in dramatic phenotypic diversity
from inactivation and possibly also activation or reactivation of genes. Finally, we show that the mutant
msh2 allele is not directly responsible for the decrease in virulence of the VGIIa-like isolates, nor does
it appear to have directly played a role in the evolution of virulence in the VGIIa clonal outbreak.
Rather, it appears to represent a parallel route of adaptation to a new environment in a successful
lineage isolated from two continents over two decades and from both the environment and patient
samples. This work suggests that hypermutator phenotypes are not limited to prokaryotes, but may
represent a frequent avenue to evolutionary change and phenotypic diversity in eukaryotic microbes as
well.

Results
We previously identified a sublineage of strains highly related to the VGIIa outbreak in the
Pacific Northwest denoted VGIIa-like, in part because they differed either in isolation location or time
from the typical outbreak isolates (24). The VGIIa-like sublineage is composed of three isolates:
NIH444, a clinical isolate from Seattle in 1975, well before the start of the outbreak; ICB107, a clinical
isolate from Brazil in 1981; and CBS7750, an environmental isolate from California in 1990. All three
isolates are characterized by diminished virulence relative to the VGIIa clonal outbreak strains (30–
32). The analysis here includes additional genomes for a South American outgroup to both the VGIIa-
like and VGIIa groups (33) (Figure 1A) as well as alignment of these genomes to the improved C.
deuterogattii genome assembly (34). This allowed determination of the ancestral state of the multiple alleles differentiating the VGIa and VGIa-like groups. We also generated a second independent genome sequence for our original NIH444 isolate (NIH444(1) and NIH444(2)), as well as an NIH444 isolate acquired from a different lab (NIH444(v)). We identified a total of 18 mutations that are shared by all of the VGIa-like isolates sequenced: 13 SNPs and 5 INDELs (Table 1). Five SNPs are noncoding while the remaining eight coding SNPs are characterized by a high rate of nonsynonymous mutations (7/8 coding mutations). In addition we confirmed that one of the INDELs in this branch is a single base deletion in a coding exon of MSH2 (24). This deletion is unique to the VGIa-like group.

We confirmed both the presence of this mutation and showed that it results in a hypermutator phenotype. Sanger sequencing verified that this single base deletion is not an Illumina sequencing artifact (Figure 1B). This single base deletion occurs within an early exon of MSH2 and results in a nonsense mutation, with both an out of frame message and an early stop codon present in the predicted transcript (Figure 1C).

**MSH2 defect results in hypermutator phenotype**

Because Msh2 is a critical component of the mismatch repair complex, we next tested whether the VGIa-like strains displayed an increased mutation rate. We utilized a standard fluctuation assay approach to determine the rate of 5-FOA resistance, as previously applied in Cryptococcus (35). The resistance rate was increased in the hypermutator lineage between 4.4-fold at the minimum (CBS7750) and 6.6-fold at the maximum (ICB107) compared to the type strain VGIa R265 (Figure 2A). The majority of 5-FOA resistance in Cryptococcus neoformans is acquired through mutation of URA5 (36); therefore, the URA5 locus was PCR amplified and sequenced in independently derived 5-FOA resistant isolates to determine the genetic basis of resistance. Mutations were characterized as either
substitutions, or indels of either single or multiple bases (Figure 2B). The majority of resistant isolates were the result of substitutions, and the mutation profile was similar between the hypermutator VGIIa-like strains and the non-hypermutator VGIIa strain EJB17 (EJB17 vs NIH444: P=1.00, EJB17 vs CBS7750 P=0.60, EJB17 vs ICB107 p=1.00, Fisher’s Exact Test).

MSH2 mutants accumulate indels in homopolymer runs

Next, the whole genome data was analyzed to ascertain whether the similarity in mutation spectrum observed in the URA5 locus was recapitulated at multiple loci or if the URA5 locus had unique properties. To do this, we focused on two relatively long branches within the C. deuterogattii phylogeny: the branch separating the outgroup VGIIb strain R272 from the VGIIa clonal cluster (58,375 variants) and the private ICB107 variants, which represented the longest available hypermutator branch (763 variants) (Figure 8). The frequency of individual SNPs was examined first. While a high rate of transitions as compared to transversions was maintained in the hypermutator branch, there was a slight reduction in the frequency of A->G and T->C mutations relative to C->T and G->A mutations (Figure 3A). However, much more striking was a dramatic increase in indels within homopolymer runs (Figure 3B). While shifts in homopolymer runs in the R272 branch account for only 0.7% of the variants, in ICB107 they account for 42.9%, exceeding even the proportion of SNPs. By looking further at the context of the homopolymer run mutations within the ICB107 branch, it is apparent that longer base runs appear to be less stable than shorter runs, as approximately 17% of the mutations occur within the context of a nine base homopolymer run (Figure 3C). This peak is likely a function of both the decreased stability of longer runs and the low quantity of longer homopolymer runs in the genome. Many of these longer runs occur within intergenic regions in the genome, but a substantial portion of the coding genes in C. deuterogattii contain at least one longer coding
homopolymer run (Figure 3C). The longest homopolymer run in the URA5 locus contains only four bases, which is predicted to be both relatively stable and among the coding genes with relatively shorter homopolymer runs (Figure 3C). In contrast, the FRR1 gene that encodes the FKBP12 homolog responsible for the antifungal action of both FK506 and rapamycin has a run of seven cytosines (7C) within its coding region. This places it both within the range of commonly mutated homopolymers in the ICB107 branch, and in the upper half of homopolymer containing genes in C. deuterogattii, suggesting that FRR1 is an appropriate locus to test the effect of mismatch repair mutants on homopolymers containing genes (Figure 3C).

**Genes containing homopolymer runs are highly unstable in msh2 mutants**

The FRR1 locus was utilized to further assess whether VGIIa-like strains were hypermutators at more than one locus and to test the hypothesis that homopolymer runs are particularly unstable in these isolates. The protein product of FRR1, FKBP12, binds to either FK506 or rapamycin to form a protein-drug complex that inhibits calcineurin or TOR, respectively. By selecting with both drugs at the same time at the non-permissive temperature of 37°C, loss of function mutations in FRR1 are selected as the only single step mutation conferring resistance to both drugs. Here we utilized a semi-quantitative assay to identify gross differences in mutation rate whereby independent colonies were grown in liquid culture and then swabbed onto quadrants of a selective plate. On FK506/rapamycin media, the wildtype strains produced a small number of resistant colonies (Figure 4A). In contrast, the NIH444 hypermutator strain produced prolific FK506/rapamycin resistant colonies. Surprisingly, the CBS7750 hypermutator was completely resistant to both drugs. Upon examining the FRR1 locus in the whole genome sequence data we discovered that there was already a single base deletion within the 7C homopolymer run in this strain. This may represent an unselected drug resistance phenotype
potentiated by the msh2 defect. We attempted to use this assay for the ICB107 strain as well but were unsuccessful, likely as a result of its inability to grow at the higher growth temperature necessary for this assay that is based upon calcineurin-dependent growth at 37°C (Figure 7B).

Next a standard fluctuation assay approach was employed to quantify the mutation rate difference between the VGIa and VGIa-like hypermutators. There was a greater than 120-fold increase in mutation rate to FK506/rapamycin resistance in NIH444 compared to R265, in contrast with the maximum of 6.6-fold increase observed for 5-FOA resistance (Figure 4C). As before, we selected independent resistant colonies, PCR amplified, and sequenced the FRR1 gene to determine the mechanism of resistance. All resistance was explained by mutations within FRR1 as expected, but while R265 still primarily acquired resistance through substitutions, NIH444 now almost exclusively underwent single base additions or single base deletions, all within the 7C homopolymer run (R265 vs NIH444, p=0.0003, Fisher's Exact Test). These indels resulted in nonsense mutations and resistance to both FK506 and rapamycin.

**Hypermutation phenotype is linked to the msh2 defect**

Once we established a *bona fide* hypermutator phenotype in the VGIa-like strains, we sought to verify that this phenotype was linked to the msh2 del131 allele. We crossed the NIH444 strain to a very closely related R265 congenic strain recently generated by serial backcrossing (37). C. *deuterogattii* matings are less fertile than *C. neoformans*, but we successfully dissected 14 viable spores from this cross. We typed these strains for the MSH2 allele by sequencing and observed 1:1 segregation of the MSH2 and the msh2 del131 alleles. The spores were then tested for rate of FK506/rapamycin resistance using the same semi-quantitative swabbing assay described above (Figure 5A). While five of the msh2 del131 strains displayed the predicted hypermutator phenotype, surprisingly two of the spores
(#4 and 5) appeared wildtype despite inheritance of a defective copy of *MSH2*.

Mating in *Cryptococcus* has previously been demonstrated to produce phenotypic plasticity through generation of aneuploid progeny at a high rate (38). These aneuploids often display defects in growth at high temperature; thus, we tested whether the high growth temperature of 37°C employed in our FK506/rapamycin resistance assay may have masked the ability of these meiotic progeny to develop drug resistance. Congruent with this hypothesis, both spore products #4 and #5 demonstrated growth defects at 37°C and 39°C (Figure 5C). Whole genome sequencing confirmed that both segregants carried an extra copy of one chromosome, suggesting that a 1N+1 aneuploidy was causing temperature sensitivity (Figure 5B). In fact, all of the progeny from this mating had an unusually high rate of aneuploidy with 8 of the 14 progeny aneuploid for at least one scaffold/chromosome (Table 2). By passaging all of the spores on YPD at 37°C we were able to restore the ability to grow at high temperatures in all of the progeny after four to nine passages after which all no longer demonstrated a high temperature growth defect (Figure 5C). As a result the passaged derivatives of isolates #4 and #5 now demonstrated a hypermutator phenotype as expected (Figure 5D) and all 14 progeny produced the results predicted through linkage of the *msh2* defect to the hypermutator phenotype (Supplemental Figure 1). Further, whole genome typing of the 14 meiotic progeny demonstrated that sufficient recombination had occurred to establish linkage. All 14 progeny demonstrated unique SNP profiles and only one SNP demonstrated inheritance congruent with the hypermutator phenotype: a SNP on scaffold 3 that was linked to the *msh2* del131 indel (~175 kb) (Figure 5E).

As a final verification of phenotypic linkage, we constructed two independent deletions of the *MSH2* gene in the R265 VGIIa background via biolistic transformation and homologous recombination replacing the *MSH2* ORF with a neomycin resistance cassette. A complete deletion of *MSH2* resulted in the same hypermutator phenotype as the *msh2* del131 allele, providing further evidence that the
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261 hypermutator phenotype was linked to the loss of function in MSH2 (Figure 5F). A fluctuation assay
262 was used to determine the mutation rate for resistance to FK506 and rapamycin and demonstrated that
263 the null mutants did not have an elevated mutation rate in comparison to the NIH444 strain (Figure
264 5G). This suggests that suppressors have not arisen on the NIH444 background to moderate the effects
265 of the msh2 del131 allele.

266

267 Hypermutation enables phenotypic diversity including reversion of mutations

268 During growth of the msh2Δ::NEO strains a spontaneous ade2 mutant was fortuitously
269 identified based on its classic red pigmentation (Figure 6A). We verified that this was the result of a
270 defect in adenine biosynthesis as this strain grows on YNB supplemented with adenine, but not YNB
271 media (Figure 6B). As expected, sequencing of the ADE2 locus revealed that the ade' phenotype was
272 linked to a single base mutation that results in a His->Arg amino acid change (Figure 6C).
273
274 The red pigment produced by ade2 mutants is a toxic intermediate in adenine synthesis that
275 accumulates in the vacuoles of mutant cells. As a result, these mutants have a growth defect, and
276 suppressor mutations can readily be isolated that eliminate production of this toxic intermediate and
277 now produce white colonies (Figure 6C). We isolated two red and two white derivative colonies and
278 sequenced the ADE2 locus. While the red colonies retained the causative substitution, one of the white
279 colonies retained it and the other had undergone reversion at this site back to the functional nucleotide
280 (Figure 6C). We confirmed that the revertant isolate was no longer an adenine auxotroph and that the
281 second white isolate was still auxotrophic, despite the lack of pigmentation (Figure 6D). This is likely
282 the result of a second mutation upstream in the adenine biosynthetic pathway, resulting in further
283 inactivation and loss of the ability to produce the toxic red intermediate. We tested a number of
284 independent white reverted colonies for the ability to grow on media lacking adenine and observed that
direct reversion to functional adenine biosynthesis was more common than additional inactivating mutations, with 37/42 (88%) white colonies demonstrating adenine prototrophy (Figure 6E). Taken together, these results suggest that hypermutation allows both frequent inactivation of pathways, and also the means to either adapt to the consequences or to simply directly revert the original mutation. In the context of antifungal drug action or activity, this phenotypic switching could be highly important.

**Hypermutation does not directly affect virulence**

We previously demonstrated that the VGIIa-like mutants exhibit diminished virulence in a nasal instillation model of virulence (30, 31). Here, we tested whether loss of Msh2 function was directly responsible for the decrease in virulence but observed no difference in virulence between R265 and two independent *de novo* deletions of *msh2* (Figure 7A). This suggested that the defects in murine virulence in the VGIIa-like isolates could be the result of multiple independent deleterious mutations unique to each lineage. We next tested the ability to grow at higher temperature, a critical virulence factor, in the VGIIa-like isolates. All three isolates showed defects in high temperature growth, varying from minor defects in NIH444, the isolate with the shortest branch to R265, to moderate defects in CBS7750, the isolate with an intermediate length branch, and finally severe defects in ICB107, the isolate with the longest branch affected by the hypermutator (Figure 7B). This suggests that instead of an immediate change in virulence, defects in mismatch repair may instead result in loss of virulence over time as mutations accumulate in critical pathways. Thus, rather than the virulence difference being explained by a single change or set of changes shared by all three VGIIa-like strains, instead they may all be avirulent for their own unique reasons.

**Hypermutation is deleterious in rich conditions, but advantageous under stress**
Our data suggests that over time a high mutation rate comes at a cost to organisms. We next addressed the early stages of hypermutator emergence and the direct effects of Msh2 inactivation to address if temporary benefits may be conferred. To this end, independent msh2 de novo deletions were employed in competition experiments with wildtype R265. Briefly, liquid YPD cultures were mixed in 50:50 ratios of wildtype R265 and a neomycin resistant msh2 mutant or a random insertion of the neomycin resistance cassette as a control. After 48 hours incubation, co-cultures were spread onto YPD plates and then colonies were picked and restruck to media containing neomycin to determine the percentage of colonies derived from each original strain. If mutations are neutral, the expectation is that the 50:50 ratio will be maintained. Instead, growth defects were observed for the msh2 mutants at either 30°C or 37°C, suggesting that the msh2 mutation is deleterious under rich growth conditions (Figure 7C). Notably, in three of the 24 replicates at 30°C or 37°C the msh2 mutant was able to grow moderately better than the wildtype parent strain. This may indicate that in these individual experimental replicates the hypermutator allowed a beneficial mutation and hitchhiked to higher frequency. However, under highly stressful conditions, such as exposure to the drugs FK506/rapamycin at 37°C, the hypermutator was highly advantageous. In multiple replicates the hypermutator rapidly acquired resistance and overtook the entire co-culture. This suggests that mutator alleles can be advantageous when a population faces an evolutionary landscape where large effect beneficial mutations exist. In contrast, mutators are deleterious in landscapes where few large effect mutations can provide advantages. Antifungal treatment and transitions from environment to host are likely to provide opportunities for adaptive mutation and favor mutator alleles.

**Hypermulator lineage is derived and hypermutation did not cause the VGIIa outbreak**

Historically, *C. deuterogattii* has been thought of as a tropical and subtropical pathogen. As a
result, the origin of the VGIIa outbreak in the Pacific Northwest, a non-tropical environment, was surprising. In addition, 1) the South American origin of the ICB107 strain (within the proposed cradle of the *C. deuterogattii* species (32)), 2) the isolation of NIH444 in Seattle near the outbreak origin, and 3) the diminished virulence of the VGIIa-like subclade (30, 31), all combined to suggest that the VGIIa-like group might have been the immediate precursors to the clonal VGIIa outbreak cluster (24, 33). Identification of the hypermutator phenotype further suggested that the defect in Msh2, carried by the older VGIIa-like group, may have played a role in adaptation to the climate of the Pacific Northwest and may have even potentiated the increase in virulence. To test this, a maximum parsimony phylogeny of the VGIIa-like and VGIIa strains was constructed, including additional sequences for a related South American VGII clade not included in our previous analysis (33) and using the VGIIb R272 strain as an outgroup (Figure 8). As described above, the *msh2* mutation can allow restoration of function mutations at the exact mutation site that are indistinguishable from the original sequence (Figure 6). Likewise, mating could also reintroduce a functional *MSH2* allele. For these reasons, we tested for the possible presence and impact of the past *msh2* del131 mutator allele throughout the phylogeny by examining the imputed mutation spectrum.

As discussed above (Figure 3), branches with defects in mismatch repair show an increased frequency of shifts in homopolymer runs. Increases in homopolymer run shifts were observed only on the most proximal branch ancestral to the VGIIa-like mutator lineage. This suggests that the mutator phenotype is congruent with the presence of the allele throughout the VGIIa-like phylogeny, and that the VGIIa group did not experience a transient period of *msh2* mediated hypermutation, followed by repair or mating-mediated replacement. Instead, the VGIIa-like lineage may represent a unique pathway to adaptation distinct from that followed by the VGIIa group that resulted in diminished rather than enhanced virulence.
Discussion

In this study we have identified and characterized a successful lineage of eukaryotic hypermutators. Elevated mutations rates are a common adaptive mechanism in bacteria, but are typically thought of as transient states that allow beneficial mutations in the short term but are selected against in the long term (3). Bacteria solve this problem through horizontal gene transfer of genes from the mismatch repair pathway, allowing the initial beneficial mutation to be separated from the deleterious mutator allele (4). In contrast, few cases have been identified in natural isolates of eukaryotes, suggesting that variation in mutation rate may play a less substantial role than in bacteria. Two recent studies have identified exceptions to this rule. In S. cerevisiae, incompatible alleles of PMS1 and MLH1 result in elevated mutation rates if present in combination (18, 19). However, in the rare cases where the incompatible arrangement is found in nature, additional suppressors have arisen to restore wildtype mutation rate, suggesting that hypermutator phenotypes may not be tolerated over long periods of time (20, 21). In addition, a recent study of Candida glabrata has demonstrated that a substantial proportion of clinical isolates (>50%) carry nonsynonymous mutations in the MSH2 gene, some causing elevated mutation rates similar to the null phenotype and others with intermediate changes in mutation rate (23). The authors correlate the mismatch repair defects with multi-drug resistance. A second study with a different sample cohort confirmed the presence of MSH2 mutations, but concluded that drug resistance was better correlated with drug exposure than with mismatch repair defects (39).

In contrast with the studies in S. cerevisiae and C. glabrata, here we identified a group of viable hypermutator strains resulting from a nonsense mutation in MSH2. These strains were isolated over a period of fifteen years from two different continents and include both clinical and environmental
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In addition, the msh2 del131 allele carried by these strains results in a complete loss of function, rather than simply reducing the efficiency of mismatch repair. This is distinct from the mutators identified in C. glabrata, and suggests that the VGIIa-like lineage is a successful and relatively long-lived hypermutator lineage, capable of both disseminating over a large area and persisting in the environment. Cryptococcus deuterogattii may also represent an intermediate between mutator-rich C. glabrata and mutator-poor S. cerevisiae. C. deuterogattii is not an obligate pathogen and can cycle between an environmental lifestyle and an infectious lifestyle as an “accidental pathogen” (26). Elevated mutation rates may be selected for by Red Queen interactions with a host, meaning that pathogens like C. glabrata that grow only in association with their hosts would experience higher mutation rates than facultative pathogens like C. deuterogattii, or effectively nonpathogenic yeasts like S. cerevisiae. In fact, a putative recurrent infection of Cryptococcus neoformans was recently demonstrated to contain nonsense mutations in MSH2, MSH5, and RAD5, predicted to result in a mutator phenotype (40).

An alternate hypothesis is that ploidy may play a role in the success of mutator strains. Both C. deuterogattii and C. glabrata exist primarily as haploids in nature, while S. cerevisiae is predominantly a diploid. Diploid strains are buffered from the effects of loss of function mutations like those observed in homopolymer runs in this study, which may reduce the supply of large effect beneficial mutations for mutator alleles to hitchhike upon in diploids. However, past work in S. cerevisiae suggests that diploids adapt more quickly in msh2 mutant backgrounds, rather than less, suggesting that ploidy may play the opposite role, at least in S. cerevisiae (41).

Evolutionary theory predicts that the admixture provided by sex in a population nullifies the ability of mutator alleles to hitchhike to high frequency (17). Unlike obligately sexual animals, or asexual bacteria, fungi can reproduce both sexually and asexually, with the frequency of sex varying
substantially between different species. We previously described *C. deuterogattii* as a species characterized by long periods of mitotic clonal expansion and only intermittent sexual crosses (24). A contributing factor is likely the highly biased mating type distribution in *C. deuterogattii*, as the vast majority of *C. deuterogattii* isolates are MATα, with only a handful of MATα isolates described globally (31). In *Cryptococcus deneoformans*, which shares a similar biased mating type distribution, this has resulted in the development of a unisexual α-α sexual cycle, dispensing with the obligate need for a MATα partner (42). This unisexual cycle can result in both de novo variation, but also recombination and admixture at similar levels to that observed in typical bisexual crosses (38, 42, 43). However, no laboratory unisexual cycle has been observed in *C. deuterogattii* to this point. Consequently, as strains are introduced to new locales, they may need to survive and adapt via mitotic growth without sexual crosses for long periods of time. This could elevate linkage between mutator alleles and the beneficial mutations they elicit, but also eliminate the ability to separate those beneficial mutations from additional deleterious alleles.

We also showed that hypermutators can allow both inactivating mutations in genes, and reversion of those mutations to the wildtype. Mismatch repair-defective mutator strains are characterized by particularly high rates of slippage within homopolymer runs, and these occur at run lengths that are common within fungal genomes. This may indicate that eukaryotes may harbor contingency loci featuring homopolymer runs, like those observed in bacteria (44). Fungal contingency loci could be critical for responses to antifungal agents, both in the environment and the clinical setting. In addition, variation could be important in host-pathogen interactions as well. For example, in *S. cerevisiae*, tandem repeats are enriched within cell surface genes, which could enable alteration of antigenic diversity during pathogenic interactions (45). While these loci are unstable even in wildtype populations, defects in the MMR pathway could enhance this instability and result in increased
diversity. In *C. neoformans*, one mutator strain has been previously described in a lab-passaged strain notable for frequent mutations in the RAM pathway, which enables a dimorphic transition between pseudohyphal and yeast phase growth, although the molecular basis of the elevated mutation rate is unknown (35). This transition is highly important for survival in the face of environmental threats such as amoeba, but compromises pathways responsible for high temperature growth, suggesting that oscillation between RAM+ and RAM- states may allow populations to survive as conditions change.

In addition, we provided evidence that the VGIIt-like lineage was not a subvirulent progenitor of the Pacific Northwest outbreak, in contrast with previous hypotheses from two groups (24, 33).

Rather, the VGIIt-like lineage may represent either an alternative pathway of adaptation or perhaps a preview of the evolutionary trajectory of the outbreak. The ability to grow at high temperatures is a critical component of virulence for fungal pathogens. However, many pathogenic fungi are thought to be “accidental” pathogens where virulence factors are selected for by other environmental factors. In this model, thermostolerance could be selected by high ambient temperatures in a non-host environment (46). Moving from a tropical/sub-tropical environment in South America to the more temperate climate of the Pacific Northwest may relieve this high temperature selection. Consequently, the loss of ability to grow under thermal stress observed in the VGIIt-like strains may be adaptive for the majority of their growth conditions in the environment, but also render them relatively avirulent in mammals. This could suggest that as the VGIIt outbreak strains continue to adapt to the Pacific Northwest, they will gradually lose virulence potential, making the outbreak self-limiting over time. Alternatively, the defects could simply reflect a decrease in viability caused by a long period of growth with an elevated mutation rate, resulting in mutational meltdown. The most severe temperature defect was observed in ICB107, a strain isolated from a patient in Brazil and also the strain with the largest number of mutations separating it from the VGIIt group, which would support the second hypothesis. These
strains may have been able to persist in the Pacific Northwest in spite of the high mutation load because of a bottleneck effect during their introduction.

Our original hypothesis was that the common variants that differentiated the VGIIa group from the progenitor of the VGIIa-like mutator group would explain the decline in virulence observed in the VGIIa-like strains (24, 31). However, the msh2 mutation, the only predicted large effect mutation, has no obvious immediate effect on virulence in a murine inhalation model. This is interesting, at least in part, because msh2 mutants of C. neoformans were previously reported to exhibit a modest growth advantage in mouse lungs in pooled signature tagged mutant experiments (47). This suggests that the mutator strains are better able to adapt to growth in the mouse lung, but not in a way that correlates directly with virulence in a dissemination and survival model of virulence. In addition, our results suggest that loss of virulence in the VGIIa-like strains may be a consequence of independent private mutations, i.e. each VGIIa-like strain is avirulent for its own reason. In previous work, CBS7750 and ICB107 have the most substantial virulence defects, while NIH444 has a more modest defect (31). This is congruent with the fact that NIH444 has the shortest divergence from the VGIIa clonal group, while CBS7750 and ICB107 have much longer branches, and is also congruent with the high temperature growth defects we observed.

Finally, we demonstrated that mutation spectrum analysis could be utilized within a phylogeny to infer the presence of a hypermutator allele through an increase in the proportion of indels within homopolymer runs. In bacteria, the hypermutator state is often transient and wildtype mismatch repair is often restored via horizontal gene transfer (4). This paradigm could also operate in fungi, with sexual recombination replacing direct DNA transfer. However, these events are difficult to detect if the hypermutator allele is purged from the population by selection after it is separated from its linked beneficial allele. The widespread availability of whole genome sequencing could now allow changes in
mutation spectrum to be used to detect episodic hypermutation throughout entire populations of microbes. We suspect that these episodes are common throughout the evolution of eukaryotic microbes and may be even more common among pathogenic microbes, reflecting their natural history as well as the result of Red Queen host-pathogen conflicts.
Materials and Methods

Strains and media

Strains used in this study are listed in table S1. Strains were routinely grown on YPD media at 30°C and maintained in permanent glycerol stocks at -80°C. Strains marked with neomycin resistance were grown on YPD supplemented with G418.

Genetic crosses and spore dissection

To conduct crosses and isolate spores, NIH444 was cocultured with the R265a congenic strain (37) on solid V8 pH =5.0 medium in the dark for eight weeks. Basidiospores were isolated using a microdissection microscope equipped with a 25-µm needle (Cora Styles Needles ‘N Blocks, Dissection Needle Kit) as previously described (48).

Virulence assays

Approximately eight week old A/J mice were anesthetized with phenobarbital via intraperitoneal injection and then were infected with 5x10^4 cells from strains R265, RBB17, and RBB18 by intranasal inhalation. Ten animals per group were infected. Mice were monitored daily for signs of cryptococcal infection and sacrificed when exhibiting signs of clinical distress. The animal study was conducted in the Division of Laboratory Animal Resources (DLAR) facilities at Duke University Medical Center (DUMC). All of the animal work was performed according to the guidelines of NIH and Duke University Institutional Animal Care and Use Committee (IACUC) under protocol number A245-13-09.
Gene disruption and strain construction

Deletions of the MSH2 gene were constructed using a standard overlap PCR approach as previously described (49). Briefly, 1 kb flanking regions of genomic DNA were amplified from both the 5' and 3' regions flanking the MSH2 open reading frame in R265 and the selectable marker for NEO resistance was amplified from plasmid pJAF1. An overlap PCR was carried out to generate a full-length deletion cassette and R265 was transformed using biolistic transformation. Gene replacement was confirmed using in-gene, 5’ junction, 3’ junction, and spanning PCRs. Two independent transformations of independent overnight cultures of R265 were carried out to isolate independent mutants.

The R265 NEO resistance marked strain was constructed by transforming XbaI digested pJAF1 plasmid into wild-type R265 by biolistics. Transformants were selected for on YPD containing neomycin. Transformants were checked for tandem insertions using primers JOHE40500/JOHE40501 and a strain with only a single insertion was chosen and employed for the competition assay.

Fluctuation assays

Fluctuation assays were performed on either synthetic medium containing 5-FOA (1 g/L) at 30°C or YPD supplemented with rapamycin (1 μg/mL) and FK506 (1 μg/mL) at 37°C. For each strain tested, ten independent 5 mL YPD liquid cultures were grown overnight at 30°C. Cultures were then split and either spread directly on selective media or diluted and spread on solid YPD to determine colony forming units. Resistant colonies and total colonies were counted and mutation rate was calculated using the Maximum Likelihood method as implemented via the FALCOR calculator (50).
Co-culture competition assays

Competition assays were carried out by growing independent liquid cultures overnight in 5 mL YPD. Cultures were then counted using a hemocytometer and 500,000 cells each of a tester neomycin resistant strain and a wildtype R265 culture were mixed in a 5 mL liquid YPD culture. Co-cultures were grown for 48 hours and then spread onto a YPD plate, such that individual colonies could be isolated. All colonies were picked (up to 60 and at least 5) and restruck to neomycin media to determine the proportion of colony forming units derived from each of the original strains in the competition.

Genome sequencing and assembly

DNA was isolated with the CTAB isolation protocol as previously described (51). Library construction and genome sequencing were carried out at the University of North Carolina Next Generation Sequencing Facility. Paired-end libraries with approximately 300-base inserts were constructed. A number of genome sequences were previously available (24, 33). The remaining were generated here and are deposited on the SRA under project accession no. PRJNA387047. All sequences were mapped to the V2 R265 reference genome (34). Alignment was performed using the short read component of the BWA aligner (52). Further processing was carried out using the Genome Analysis Toolkit (GATK) version 3.1-1 (53), including SAMtools (54) and Picard. SNPs and indels were called with UnifiedGenotyper from GATK, using the haploid setting. GATK’s VariantAnnotator was used to define the homopolymer context of indels. VCFtools was used for filtering and extracting private variants (55). SnpEff was employed to determine the predicted impact of mutations (56).
Inference of phylogeny

Maximum parsimony phylogenies were constructed using MEGA6 (57) to analyze SNP matrices extracted from VCF format using a custom Perl script. SNPs and indels were placed on branches where they were predicted to change states using VCFtools (55) to extract the variants supporting each node.

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**Figure legends**

**Figure 1. VGIiA-like isolates harbor a frameshift resulting in a predicted nonfunctional protein.**

A) The VGIiA-like clade is part of the clonal radiation that includes the Pacific Northwest Outbreak. All three isolates are characterized by a frameshift in the MSH2 gene at position 131. B) Sanger sequencing of the MSH2 gene confirms that NIH444 and CBS7750 have both undergone deletion of a single T within the coding region of MSH2, while the VGIiA R265 strain and the outgroup VGIiB strain have not. C) This deletion results in a frameshift beginning in the first functional domain of MSH2 and an early premature stop. This truncated protein is predicted to be non-functional.

**Figure 2. msh2 mutants are hypermutators.**

A) A fluctuation assay for resistance to 5-FOA was carried out for both wildtype VGIiA strains (R265 and EJB17) and hypermutator VGIiA-like strains (NIH444, EJB17, and ICB107), as well as an outgroup VGIiB (R272) strain. Resistance rates were normalized to the rate observed in R265. The VGIiA-like strains demonstrate an increase in mutation rate. Data shown are the mean of ten replicates with 95% confidence interval. B) The molecular basis of resistance was determined for 5-FOA resistance at the URA5 locus. All isolates tested demonstrated predominantly substitutions as the molecular basis of resistance.

**Figure 3. Hypermutators are characterized by high homopolymer instability.**

A) Percentage of transitions and transversions among SNPs private to R272, an outgroup, and ICB107, a hypermutator, are shown. Transitions are more common in both strains, while one type of transition is modestly reduced in the hypermutator. B) All private mutations within R272 and ICB107 were characterized either as SNPs, indels, or homopolymer indels. The cutoff used to distinguish a
homopolymer indel was longer than 4 bases. ICB107 had a substantial increase in homopolymer run shifts. C) Frequency of indels within all homopolymers longer than 1 base from the private ICB107 mutations are shown in blue. All genes from the *C. deuterogatti* transcriptome were grouped by frequency of the longest homopolymer run within the coding region of that gene and shown in red.

*URA5* only has a run of 4 bases, while *FRR1*, which encodes FKBP12, the target of FK506 and rapamycin, has a run of 7 bases.

**Figure 4. A homopolymer run within the *FRR1* gene allows rapid inactivation and resistance to FK506 and rapamycin.**

A) Swab assays of the VGIIa and VGIIa-like strains were conducted to determine whether VGIIa-like stains develop resistance to FK506 and rapamycin at 37°C at an elevated rate compared to VGIIa. NIH444 demonstrated a large increase in resistance rate, while CBS7750 was completely resistant. B) The resistance in CBS7750 is attributable to a single base deletion within the coding 7C run in the *FRR1* gene that has been fixed in this strain. C) Fluctuation assays for NIH444 and R265 show that the hypermutator conferred a greater than 100-fold increase in mutation rate compared to the wildtype VGIIa strain. Data shown are the mean of ten replicates with 95% confidence intervals and are normalized to the rate in R265. D) Analysis of the molecular basis of resistance shows that substitutions are still the predominant mechanism for resistance in R265, but in the hypermutator strain single base additions and single base deletions within the homopolymer run are responsible for the vast majority of resistance.

**Figure 5. The hypermutator phenotype is linked to the frameshift in *MSH2*.**

A) Progeny from a cross between NIH444 and R265 show co-segregation of the *msh2 del131* allele
with the hypermutator phenotype in all except two cases. B) Whole genome depth of coverage plots show that scaffolds five and six are aneuploid in spores #4 and #5, respectively. C) Temperature sensitivity assay for the incongruent spores before and after serial passaging at 37°C. D) After passage at 37°C, the non-congruent spores now properly demonstrate linkage with the msh2 del131 allele. E) Whole genome sequencing of NIH444 x R265a progeny, with variant regions indicated as reference in blue or alternate in red. The boxed region indicates a SNP in close linkage to the msh2 del131 allele. F) Two independent de novo deletions of MSH2 via biolistic transformation demonstrate elevated mutation rates in an FK506/rapamycin swabbing assay at 37°C. G) Fluctuation assays for R265, NIH444, and two independent deletions of msh2 in R265 show that the hypermutator phenotype is recapitulated in the null mutants. Data shown are the mean of ten replicates with 95% confidence intervals and are normalized to the rate observed in R265.

**Figure 6. Hypermutation allows inactivation and reactivation of adenine biosynthetic pathway.**

A) A spontaneous red colony was isolated from the de novo msh2 deletion (RBB18). B) This colony (RBB22) demonstrated adenine auxotrophy, suggesting that it was an ade2 mutant. C) Sequencing of the ADE2 locus confirmed that the original colony was an ade2 mutant. In addition, two red (RBB25 and RBB26) and two white derivatives (RBB23 and RBB24) were tested. One white derivative had reverted the original mutation (RBB23), while the second had eliminated production of the red intermediate but had not reverted the original ade2 mutation (RBB24). D) One revertant colony (RBB23) demonstrated adenine prototrophy, while the other (RBB24) remained an auxotroph despite losing the red pigmentation. E) An assay to test direct reversion frequency versus secondary mutation to eliminate the red toxic intermediate demonstrated that the most common mutations were direct reversions and restoration of adenine prototrophy.
Figure 7. 

Hypermutteration does not have immediate virulence defects but may potentiate long term deficits.

A) Virulence tested in the murine inhalation model was not strongly affected by deletion of the MSH2 gene. B) All three VGIIa-like strains demonstrated defects in high temperature growth via a spot dilution assay in comparison with the VGIIa R265 strain and the VGIIb R272 strain as an outgroup. Strains with longer branches exhibited larger high temperature growth defects. C) Competition experiments between a tester strain with the neomycin resistance marker and the wildtype R265 strain. (Strain used: SEC501, RBB17, RBB18). Original cultures were mixed in a 1:1 ratio and then grown overnight in liquid YPD. Both hypermutator strains showed a modest growth defect at 30°C and 37°C but a dramatic growth advantage in the high stress FK506/rapamycin 37°C condition. Boxplots show minimum, first quartile, median, third quartile, and maximum values. Points represent the results from six individual replicates summarized by the box plot. The NEO vs WT competition is shown in gray, while the two msh2Δ::NEO competitions are shown in dark and light blue.

Figure 8. The VGIIa-like hypermutator is derived and not ancestral to the Pacific Northwest outbreak.

A maximum parsimony phylogeny of the VGIIa group with the VGIIb R272 genome as an outgroup demonstrates that the VGIIa-like group is a branch parallel to the VGIIa group. To test for the presence of a defect in MMR throughout the tree, the mutation spectrum was examined on each branch. High rates of homopolymer run shifts were observed throughout the VGIIa-like group, but no evidence was apparent at branch A). Instead it appears that the hypermutator first arose on branch B).
Supplemental Figure 1. Passaged spore progeny demonstrate linkage between msh2del131 allele and mutator phenotype.

Four independent overnight cultures of spore progeny passaged at 37°C to select against aneuploidies that confer temperature sensitivity were struck on YPD media containing FK506 and rapamycin at 37°C. All fourteen passaged spore progeny demonstrated behavior predicted by the state of the MSH2 locus for the majority of the four samples tested. Spores #4 and #5 and controls are depicted in Figure 5.

Supplemental Figure 2. Reversion of ade2 mutants primarily occurs through repair of ADE2.

42 white colonies were selected as independent revertants from a red ade2 mutant (RBB22). Colonies were struck to both YPD and YNB to test for auxotrophy indicative of either direct reversion or a second site mutation upstream of ade2 in the adenine biosynthetic pathway.
| Strain  | Genotype                              | Source/Reference |
|---------|---------------------------------------|------------------|
| R265    | VGIla                                 | (29)             |
| EJB17   | VGIla                                 | (31)             |
| R272    | VGIlb                                 | (30)             |
| NIH444  | VGIla-like, msh2del131                | (30)             |
| NIH444(v) | VGIla-like, msh2del131               | (58)             |
| CBS7750 | VGIla-like, msh2del131                | (30)             |
| ICB107  | VGIla-like, msh2del131                | (30)             |
| RBB17   | R265, msh2Δ::NEO                      | This study       |
| RBB18   | R265, msh2Δ::NEO                      | This study       |
| RBB22   | RBB18, msh2Δ::NEO, ade2               | This study       |
| RBB23   | RBB22, msh2Δ::NEO, white, ADE2        | This study       |
| RBB24   | RBB22, msh2Δ::NEO, white, ade2        | This study       |
| RBB25   | RBB22, msh2Δ::NEO, red, ade2          | This study       |
| RBB26   | RBB22, msh2Δ::NEO, red, ade2          | This study       |
| ECT1    | XL280 ade2::NEO                       | This study       |
| R265a   | R265, MATa                            | (37)             |
| SEC016  | NIH444xR265a spore #1, msh2del131     | This study       |
| SEC017  | NIH444xR265a spore #2                 | This study       |
| SEC018  | NIH444xR265a spore #3, MATa, msh2del131 | This study     |
| SEC019  | NIH444xR265a spore #4, MATa, msh2del131 | This study   |
| SEC020  | NIH444xR265a spore #5, msh2del131     | This study       |
| SEC021  | NIH444xR265a spore #6, MATa, msh2del131 | This study   |
| SEC022  | NIH444xR265a spore #7                 | This study       |
| SEC023  | NIH444xR265a spore #8                 | This study       |
| SEC024  | NIH444xR265a spore #9, MATa           | This study       |
| SEC025  | NIH444xR265a spore #10, MATa          | This study       |
| SEC026  | NIH444xR265a spore #11                | This study       |
| SEC027  | NIH444xR265a spore #12, MATa          | This study       |
| SEC028  | NIH444xR265a spore #13, msh2del131    | This study       |
| Research Article | Final 5/23/17 |
|------------------|--------------|
| SEC029           | NIH444xR265a spore #14, MATa, msh2del131 This study |
| SEC501           | R265, NEO This study |
| SEC559           | SEC016, passed 4 times at 37°C This study |
| SEC560           | SEC017, passed 4 times at 37°C This study |
| SEC561           | SEC018, passed 4 times at 37°C This study |
| SEC562           | SEC019, passed 4 times at 37°C This study |
| SEC563           | SEC020, passed 4 times at 37°C This study |
| SEC564           | SEC021, passed 4 times at 37°C This study |
| SEC565           | SEC022, passed 4 times at 37°C This study |
| SEC566           | SEC023, passed 4 times at 37°C This study |
| SEC567           | SEC024, passed 4 times at 37°C This study |
| SEC568           | SEC025, passed 4 times at 37°C This study |
| SEC569           | SEC026, passed 4 times at 37°C This study |
| SEC570           | SEC027, passed 4 times at 37°C This study |
| SEC571           | SEC028, passed 4 times at 37°C This study |
| SEC572           | SEC029, passed 4 times at 37°C This study |
| SEC573           | R265a, passed 4 times at 37°C This study |
| SEC574           | NIH444, passed 4 times at 37°C This study |
| SEC575           | SEC562, passed 5 additional times at 37°C This study |
Supplemental Table 2. Oligonucleotides used in this study.

| Primer                | Sequence (5' to 3')          | Comment                  |
|-----------------------|------------------------------|--------------------------|
| M13F                  | GTAAACGACGGCGCCAG            | universal oligo          |
| M13R                  | CAGGAAACAGCTATGAC            | universal oligo          |
| JOHE41021/BB265       | GAGCGTATTTGGAAGCAGGG         | MSH2 5' F deletion       |
| JOHE41022/BB266       | CATCTGGCCATAGTGACGC          | MSH2 5' F nested deletion|
| JOHE41023/BB267       | CTGGCCGCTGTTTTTACAAAGACGAA  | MSH2 5' R deletion       |
| JOHE41024/BB268       | GTCATAGCTGTGTTCCTGCGTTCGCA   | MSH2 3' F deletion       |
| JOHE41025/BB269       | CGACATTTGATGGAACCTCCACC     | MSH2 3' R deletion       |
| JOHE41026/BB270       | GTCATAATCGCCGGACACTTGCG     | MSH2 3' R nested deletion|
| JOHE41027/BB271       | CCGAGAAAGCAGAAAGTGACC       | MSH2 F sequencing        |
| JOHE41245/BB275       | CCAAGCAGATGCGTATCG          | MSH2 R sequencing        |
| JOHE42336/BB280       | CCTTCAAACAGGCGAAAGTGGA      | ADE2 5' F for sequencing |
| JOHE42337/BB281       | GTGTTTGTGCGCTGACTCGG        | ADE2 5' F nested for sequencing |
| JOHE42338/BB282       | CGGAAGTCAGCGCAAAGTCC        | ADE2 F ~400 for sequencing |
| JOHE42339/BB283       | GGCCTGAGAAAGGCGCTGGG        | ADE2 F ~800 for sequencing |
| JOHE42340/BB284       | GGTATGGATCTCTCTGGGACCTCC    | ADE2 F ~1300 for sequencing |
| JOHE42341/BB285       | CCGACATCAAGCGCAACAGGG       | ADE2 3' R for sequencing |
| JOHE42342/BB286       | CGTGCTGCAGATGCTGGG          | ADE2 3' R nested for sequencing |
| JOHE42343/BB287       | CCAGGCGTCTAGTGACCTG         | ADE2 F ~1600 for sequencing |
| JOHE42344/BB288       | GCAGCACCAGAAACAGTGAGAGG     | ADE2 F ~1100 for sequencing |
| JOHE42345/BB289       | CCAACCTTCAGCATAAAGAGGG      | ADE2 F ~600 for sequencing |
| JOHE26938/BB14        | GTCTTTCCAAGCCTCGAATCCT      | URA5 5' F for sequencing |
| JOHE26939/BB15        | CCGGTAGCCATACCATGCAGC       | URA5 5' F nested for sequencing |
| JOHE26941/BB17        | CCTGTACCTTCCTGCACCTCAG      | URA5 3' R for sequencing |
| JOHE26942/BB18        | CCCACTTTCCGGAGCCTCC         | URA5 3' R nested for sequencing |
| JOHE40363/BB257       | GTGGTTGGACGAGCAGTGGG        | FRR1 3' R for sequencing |
| JOHE40364/BB258       | GCAGCAATGCAATCCCTGG         | FRR1 3' R nested for sequencing |
| JOHE40365/BB259       | GGTACAGGGCGTTGGGACC         | FRR1 5' F for sequencing |
| JOHE40366/BB260       | CGACCTGCAATAGTTCCCGAGC     | FRR1 5' F nested for sequencing |
| Sample ID | Primer Sequence | Confirmation Type |
|-----------|----------------|------------------|
| JOHE40500/SEC129 | GGGATGACAGGAAGATCCTGC | NEO 3' R for insert confirmation |
| JOHE40501/SEC130 | GCAACAATCCATCCGTGCTGG | NEO 5' F for insert confirmation |
Figure 1

A

V Gill-like
msh2del131

V Gilla
MSH2

V Gillb
(outgroup)
MSH2

WM clade
MSH2

B

R265
V Gilla

NIH444
V Gilla-like

CBS7750
V Gilla-like

R272
V Gillb (outgroup)

TT

TT → T

TT → T

TT

C

Msh2

MutS I
MutS II
MutS III
MutS IV

MSH2
ATP-binding cassette

msh2 del 131 stop

del 131 stop
Figure 3

A

B

C

Length of longest coding homopolymer run per gene in *C. deuterogattii*  Homopolymer length context of indels of ICB107
Figure 6

A

YPD

msh2Δ::NEO-2

B

YNB

XL280 ade2

R265

YNB + Adenine

R265 msh2Δ::NEO-2 ade2

Figure 6

C

R265

R265 msh2Δ original red (RBB22)

R265 msh2Δ red #1 (RBB25)

R265 msh2Δ red #2 (RBB26)

R265 msh2Δ white #1 (RBB23)

R265 msh2Δ white #2 (RBB24)

D

YNB

YNB + Adenine

Figure 6

E

Percentage of white revertants

0 25 50 75 100

ade- ADE+

Figure 6
| Mutation | Location   | Predicted Effect          | Gene         | Predicted Function          |
|----------|------------|---------------------------|--------------|-----------------------------|
| SNP      | SC2.1:1493731 | Missense (Ser->Pro)      | CNBG_0538    | t-RNA ligase                |
| SNP      | SC2.2:591810 | Splice site acceptor     | CNBG_0776    | HAD hydrolase               |
| SNP      | SC2.2:965997 | Missense (Gly->Asp)      | CNBG_0907    | Clp protease subunit        |
| SNP      | SC2.4:899106 | Missense (Leu->Val)      | CNBG_1894    | RhoGEF                      |
| SNP      | SC2.6:160627 | Missense (Ala->Val)      | CNBG_2540    | Ubiquitin-like              |
| SNP      | SC2.6:410824 | Intergenic               | -            |                             |
| SNP      | SC2.6:1157157 | Intergenic               | -            |                             |
| SNP      | SC2.7:290975 | Missense (His->Tyr)      | CNBG_3034    | BRCT domain-containing      |
| SNP      | SC2.8:594146 | Synonymous               | CNBG_3547    | snRNP                       |
| SNP      | SC2.10:94373 | Intronic                 | CNBG_3750    | Dcp2                        |
| SNP      | SC2.13:413395 | Missense (Leu->Ser)      | CNBG_4876    | esterase                    |
| SNP      | SC2.15:96883 | Intergenic               | -            |                             |
| SNP      | SC2.16:143098 | Missense (Pro->Leu)     | CNBG_5437    | Protein kinase Nrc2         |
| INDEL    | SC2.1:15401-15402 | Intronic            | CNBG_0005    | sugar metabolism            |
| INDEL    | SC2.3:240763-240764 | Nonsense          | CNBG_1661    | Msh2                        |
| INDEL    | SC2.9:241948-241949 | Intronic            | CNBG_4065    | SAM methyltransferase       |
| INDEL    | SC2.14:162875-162876 | Intergenic         | -            |                             |
| INDEL    | SC2.15:299154-299155 | Intergenic         | -            |                             |
Table 2. Aneuploidy in NIH444 x R265a cross

| Spore # | Aneuploid scaffold(s) | MSH2 allele |
|---------|-----------------------|-------------|
| 1       | -                     | msh2del131  |
| 2       | 19,21,26              | WT          |
| 3       | -                     | msh2del131  |
| 4       | 5                     | msh2del131  |
| 5       | 6                     | msh2del131  |
| 6       | -                     | msh2del131  |
| 7       | 6                     | WT          |
| 8       | -                     | WT          |
| 9       | 6                     | WT          |
| 10      | 6                     | WT          |
| 11      | 6                     | WT          |
| 12      | -                     | WT          |
| 13      | -                     | msh2del131  |
| 14      | 19,21,26              | msh2del131  |
YPD + FK506/rapamycin at 37°C

Passaged spores
Supplemental Figure 2

YNB

Rev 1-6  Rev 7-12  Rev 13-18  Rev 19-24  Rev 25-30  Rev 31-36  Rev 37-42

YPD

ade- WT

Controls

ade- WT

YPD