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Genetic Diversity and Genomic Plasticity of Cryptococcus neoformans AD Hybrid Strains

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ABSTRACT Natural hybridization between two strains, varieties, or species is a common phenomenon in both plants and animals. Although hybridization may skew established gene pools, it generates population diversity efficiently and sometimes results in the emergence of newly adapted genotypes. Cryptococcus neoformans, which causes the most frequent opportunistic fungal infection in immunocompromised hosts, has three serotypes: A, D, and AD. Serotype-specific multilocus sequence typing and serotype-specific comparative genome hybridization were applied to investigate the genetic variability and genomic organization of C. neoformans serotype AD isolates. We confirm that C. neoformans serotype AD isolates are hybrids of serotype A and D strains. Compared with haploid strains, most AD hybrid isolates exhibit unique multilocus sequence typing genotypes, suggesting that multiple independent hybridization events punctuated the origin and evolutionary trajectory of AD hybrids. The MATa alleles from both haploid and AD hybrid isolates group closely to form a cluster or subcluster in both the serotype A and D populations. The rare and unique distribution of MATa alleles may restrict sexual reproduction between isolates of opposite mating types. The genetic diversity of the serotype D population, including haploid strains and serotype D genomes of the AD hybrid, is significantly greater than that of serotype A, and there are signatures of recombination within the serotype D population. Given that MATa isolates are relatively rare, both opposite-sex and same-sex mating may contribute to genetic recombination of serotype D in nature. Extensive chromosome loss was observed in AD hybrid isolates, which results in loss of heterozygosity in the otherwise-heterozygous AD hybrid genome. Most AD hybrid isolates exhibit hybrid vigor and are resistant to the antifungal drug FK506. In addition, the C. neoformans AD hybrid genome is highly dynamic, with continuous chromosome loss, which may be a facile route for pathogen evolution through which genotypic and phenotypic variation is generated.

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DNA sequences have been submitted to the GenBank database at NCBI as series JQ014210-JQ014283
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between a male lion (Panthera leo) and a tigress (Panthera tigris), exhibits prominent hybrid vigor in the animal kingdom. Reports of natural hybridization in fungi were rare until recently (Olson and Stenlid 2002). Interspecific hybrids allow the colonization of novel habitats and expanded host range in several plant fungal pathogens (Brasier et al. 1999; Hamilton et al. 2009; Inderbitzin et al. 2011; Moon et al. 2004) and contribute significantly to the generation of novel species and stress adaptability in the Saccharomyces sensu stricto complex (de Barros Lopes et al. 2002; Querol et al. 2003; Sipiczki 2008).

Cryptococcus neoformans is a ubiquitous fungal pathogen that causes meningocencephalitis and pneumonia in immunocompromised hosts. It is one of the most common causes of death in HIV-infected patients (Mitchell and Perfect 1995; Perfect and Casadevall 2002). On the basis of antibody-promoted slide agglutination assays of polysaccharide capsules, C. neoformans has been classified into three serotypes: serotype A, D, and AD (Dromer et al. 1993; Ikeda et al. 1996; Kabasawa et al. 1991). Serotype A is distributed globally and causes >95% of human infections (Mitchell and Perfect 1995), whereas serotype D is mostly found in Europe but has a sporadic global distribution (Dromer et al. 2007). Serotype AD isolates that react with both serotype A and serotype D antisera to capsular polysaccharides are hybrids of serotype A and D strains. Although AD hybrid isolates are generally regarded as less frequent causes of human infections (Kabasawa et al. 1991; Mitchell and Perfect 1995), a recent survey in Europe revealed that 19% of human infections are caused by AD hybrid strains (Viviani et al. 2006). The finding of an unexpectedly high prevalence of serotype AD infections may be explained through the application of polymerase chain reaction (PCR) assays in molecular detection of C. neoformans and is likely caused by the impact of hybridization on virulence potential. Interspecific hybrids of C. neoformans and Cryptococcus gatti have recently been identified in human infections and may be much more common than previously appreciated (Bowers et al. 2006, 2008; Viviani et al. 2006).

Most AD hybrid strains are diploid or aneuploid (Cogliati et al. 2006; Lengeler et al. 2001; Xu et al. 2002; Xu and Mitchell 2003). Molecular analyses provide evidence that AD hybrid strains are the result of the hybridization of serotype A and D strains, a phenomenon in which mating plays a central role (Xu et al. 2000, 2002; Xu and Mitchell 2003). C. neoformans has a bipolar mating system that is controlled by the mating type locus (MAT) (Fraser et al. 2004; Lengeler et al. 2002). Two opposite mating types, a and α, are known in C. neoformans, but MATα predominates in clinical and environmental populations (Kwon-Chung and Bennett 1978). Mating has been observed between strains of opposite mating types in the laboratory. A filamentous dikaryon is produced after cell—cell fusion, and nuclear fusion occurs later in the fruiting body (the basidium), resulting in a transient a/α diploid state that immediately undergoes meiosis and sporulation to produce α and a haploid progeny (Kwon-Chung 1975, 1976). Although most serotype A isolates are exclusively MATα, serotype A MATα strains have been recently identified in Botswana (Litvintseva et al. 2003, 2006), and there is evidence of sexual reproduction and genetic recombination in this population (Litvintseva et al. 2003). C. neoformans can also undergo unisexual mating, especially between α cells, to produce stable α/α diploids as well as haploid progeny (Lin et al. 2005, 2007).

It has been estimated that serotype A and D of C. neoformans diverged ~18 million years ago and have ~10% to 15% nucleotide polymorphism at the whole genome level (Kavanaugh et al. 2006; Sun and Xu 2009; Xu et al. 2002). In addition, genomic rearrangements make the genetic distance between these two serotypes even greater (supporting information, Figure S1) (Kavanaugh et al. 2006; Sun and Xu 2009). When serotype A and D strains are mixed under appropriate conditions in the laboratory, cell—cell fusion proceeds normally but meiosis is impaired because of their highly divergent genetic backgrounds (Lengeler et al. 2001; Sun and Xu 2007; Xu et al. 2000, 2002), thus resulting in few viable haploid progeny (Lengeler et al. 2001; Sun and Xu 2007). Despite impaired meiosis, AD hybrid cells resulting from these intravarietal matings can propagate mitotically as uninucleate diploid yeast cells after nuclear fusion. Thus, many natural AD hybrids remain in the diploid (or aneuploid) state (Lengeler et al. 2001). The authors of a previous study reported approximately 5% viability of the meiotic progeny of AD hybrids (Lengeler et al. 2001). The hybrids may have recombinant chromosomes rather than a simple combination of genomes from the two serotypes. The resulting recombinant genome may generate novel phenotypic traits. The laboratory-constructed AD hybrid strain exhibits greater virulence than parental serotype A and D strains in vivo and in vitro; they were resistant to ultraviolet light and have larger capsules (Lin et al. 2007). In some mouse infection experiments, C. neoformans AD hybrid strains exhibited intermediate or greater virulence than haploid serotype A and D (Barchiesi et al. 2005; Chaturvedi et al. 2002).

Genetic diversity and genomic organization may ultimately explain phenotypic variability of the pathogen, including, for example, host specificity, pathogenicity, antimicrobial resistance, and virulence. The genome organization of C. neoformans AD hybrids is largely unknown; however, the authors of a previous study found that some AD hybrid isolates are aneuploid on the basis of fluorescence-activated cell sorting (FACS) analysis, whereas others exhibit roughly the same DNA content as diploid control strains (Lengeler et al. 2001). Comparative gene genealogical analyses have provided evidence that the AD hybrid results from hybridization of serotype A and D strains (Xu et al. 2000, 2002, 2003). However, these studies were based on only two genes, LAC1 and URAS, and a limited number of AD hybrid isolates were studied. The limited genetic information from these two genes may not be sufficient to investigate the origin and evolutionary trajectory of the AD hybrid. The limited number of available AD hybrid isolates may exclude some important evolutionary lineages that have yet to be isolated. In addition, whether hybridization between the two C. neoformans serotypes contributes to speciation is debatable (Solits and Solits 2009). Whole-genome duplication has been described in the Saccharomyces complex and it contributed greatly to speciation of this group by generating a number of novel species (Querol et al. 2003; Sipiczki 2008).

In this study, we explored the origin, population dynamics, evolutionary trajectory, and genomic plasticity of the C. neoformans AD hybrid by characterizing a large number of isolates in terms of genetic diversity and genomic organization using mating type- and serotype-specific PCR, multilocus sequence typing (MLST), and comparative genome hybridization (CGH). Our results reveal that serotype AD strains of C. neoformans result from multiple, independent hybridization events that yielded a highly dynamic hybrid genome. Thus, the AD hybrid is an important source of genotypic and phenotypic variation in C. neoformans.

**MATERIALS AND METHODS**

**Strains and media**

In total, 64 C. neoformans isolates, including 31 serotype AD and 33 serotype D isolates (32 haploid and one diploid), are included in this study (Table 1). All strains were cultured on yeast extract-peptone-dextrose (YPD) agar medium at 30°. To conduct comparative
Table 1 The mating type, serotype, and origin of *C. neoformans* isolates included in this study

| Isolate   | Mating type | Serotype | Origin     |
|-----------|-------------|----------|------------|
| 713       | αAD         | AD       | Italy      |
| 42-10     | αAD         | AD       | NC, USA    |
| 5-19      | αAD         | AD       | NC, USA    |
| 6-20      | αAD         | AD       | NC, USA    |
| ATCC48184 | αAD         | AD       | Japan      |
| CDC228    | αAD         | AD       | USA        |
| CDC304    | αAD         | AD       | USA        |
| CDC92-74  | -ADx        | AD       | USA        |
| CDC94-383 | -ADx        | AD       | Unknown    |
| CBS132    | αAD         | AD       | Italy      |
| IT752     | αAD         | AD       | Italy      |
| IT756     | αAD         | AD       | Italy      |
| IUM92-4686| αAD-        | AD       | Italy      |
| IUM92-6198| αADx        | AD       | Italy      |
| KW5       | αADa        | AD       | Kuwait     |
| MMRL752   | αADx        | AD       | Italy      |
| MMRL774   | αADx        | AD       | Italy      |
| MMRL1351  | -ADx        | AD       | USA        |
| MMRL1365  | αADx        | AD       | USA        |
| NC34-21   | αADx        | AD       | NC, USA    |
| ZG287     | αADa        | AD       | China      |
| ZG290     | αADa        | AD       | China      |
| AD6-93    | αAD         | AD       | Sub-Saharan Africa |
| AD7-97    | αAD         | AD       | North Africa |
| AD6-38    | αAD         | AD       | Europe/North Africa |
| AD7-91    | -ADx        | AD       | Sub-Saharan Africa |
| AD7-85    | -ADx        | AD       | Sub-Saharan Africa |
| AD7-69    | αAD-        | AD       | Europe     |
| AD7-75    | αADx        | AD       | Europe     |
| AD2-71    | αADx        | AD       | Europe     |
| AD7-95    | αADx        | AD       | Europe     |
| CDC92-27  | D           | D        | USA        |
| VANC.R461 | D           | D        | Canada     |
| 431       | D           | D        | Denmark    |
| AD7-71    | D           | D        | Europe     |
| AD3-14    | D           | D        | Europe/Africa |
| AD3-13    | D           | D        | Europe/Africa |
| AD2-62    | D           | D        | Europe     |
| AD2-95    | D           | D        | Europe     |
| AD2-96    | D           | D        | Europe     |
| MMRL1076  | D           | D        | USA        |
| 2-14      | D           | D        | NC, USA    |
| 3-15      | D           | D        | NC, USA    |
| 3-28      | D           | D        | NC, USA    |
| NIH12     | D           | D        | Denmark    |
| MMRL751   | D           | D        | Italy      |
| 528       | D           | D        | Italy      |
| 529       | D           | D        | Italy      |
| 709       | D           | D        | Italy      |
| MMRL760   | D           | D        | Denmark    |
| 434       | D           | D        | Denmark    |
| MMRL757   | D           | D        | Italy      |
| 3311      | D           | D        | NC, USA    |
| B3179     | D           | D        | USA        |
| CDC92-18  | D           | D        | USA        |
| Y290-90   | D           | D        | Canada     |
| 2-22      | D           | D        | NC, USA    |
| CAP672    | D           | D        | Unknown    |
| NIH264    | D           | D        | USA        |
| NIH276    | D           | D        | USA        |

Table 1 Continued

| Isolate | Mating type | Serotype | Origin |
|---------|-------------|----------|--------|
| NIH430  | Da          | D        | USA    |
| NIH433  | Da          | D        | USA    |
| AD8-62  | αDAD        | D        | Europe |

NC, North Carolina

a The mating type and molecular serotype were determined by mating-type and serotype-specific PCR amplification of the STE20 gene (see Figure S2).

Phylogenetic analyses, DNA sequences of five MLST markers (IGS, URE1, GPD1, LAC1, and MPDI) in eight serotype A strains (2 VNI, 2 VNNI, and 4 VNB) from previous studies were downloaded from GenBank.

**Ploidy analysis by fluorescence flow cytometry (FACS)**

FACS analysis was performed as described previously (Tanaka et al. 1996). In brief, yeast cells were harvested after growth on YPD medium for 48 hr, washed once in phosphate-buffered saline buffer, and fixed in 1 ml of 70% ethanol overnight at 4°C. Fixed cells were then washed with 1 ml of NS buffer (10 mM Tris-HCl [pH 7.6]; 0.25 M sucrose; 1 mM EDTA [pH 8.0]; 1 mM MgCl2; 0.1 mM CaCl2; and 0.1 mM ZnCl2) and then stained with propidium iodide (12 mg/ml) in 0.2 ml of NS buffer containing RNaseA (1 mg/ml) at 4°C for 8 hr in the dark. Then, 0.05 ml of the stained cell preparation was diluted into 1.95 ml of 50 mM Tris-HCl (pH 8.0) and sonicated for 1 min. FACS was performed on 10,000 cells and analyzed on the FL1 channel of a Becton-Dickinson FACS-can. Haploid isolate H99 and diploid isolate XL143 were included as controls.

**Genomic DNA isolation, PCR amplification, and DNA sequencing**

After 2 days of growth on YPD agar, *C. neoformans* strains were collected directly for genomic DNA isolation. Genomic DNA isolation was performed using the MasterPure Yeast DNA Purification Kit (Epicentre Biotechnologies) with minor modifications. In brief, 500 μl of glass beads (425–600 nm) and 300 μl of cell lysis solution were added to the cells to assist cell wall disruption. Others steps followed the protocol provided by the manufacturer.

All PCR assays were conducted in a PTC-200 automated thermal cycler (MJ Research, Waltham, MA); 300 ng of DNA was amplified in a 25-μl reaction mixture containing 10 μM of each primer, 2 mM of each nucleotide (dATP, dCTP, dGTP, and dTTP), 2.5 μl of 10x Ex Taq buffer, 0.125 μl of Ex Taq polymerase (Takara, Shiga, Japan), and an appropriate volume of distilled water. The following conditions were used for the amplification: an initial 2 min of denaturation at 98°C, followed by 35 cycles of denaturation for 10 sec at 98°C, an annealing time of 15 sec at 54°C, and an extension cycle for 1 min at 72°C. The amplification was completed with an extension period of 5 min at 72°C. Sterile water served as a negative control in each assay. PCR products were analyzed on 1% agarose gels (MLST and serotype/mating type markers). In brief, yeast cells were harvested after growth on YPD medium for 48 hr, washed once in phosphate-buffered saline buffer, and fixed in 1 ml of 70% ethanol overnight at 4°C. Fixed cells were then washed with 1 ml of NS buffer (10 mM Tris-HCl [pH 7.6]; 0.25 M sucrose; 1 mM EDTA [pH 8.0]; 1 mM MgCl2; 0.1 mM CaCl2; and 0.1 mM ZnCl2) and then stained with propidium iodide (12 mg/ml) in 0.2 ml of NS buffer containing RNaseA (1 mg/ml) at 4°C for 8 hr in the dark. Then, 0.05 ml of the stained cell preparation was diluted into 1.95 ml of 50 mM Tris-HCl (pH 8.0) and sonicated for 1 min. FACS was performed on 10,000 cells and analyzed on the FL1 channel of a Becton-Dickinson FACS-can. Haploid isolate H99 and diploid isolate XL143 were included as controls.

Amplicons were purified using the QIAquick PCR Purification Kit (QIAGEN) as recommended by the manufacturer. PCR products were sequenced in both directions using BigDye Terminator version 3.1 cycle sequencing ready reaction mix (Applied Biosystems). Sequencing products were resolved using an ABI 3130 automated sequencer (Applied Biosystems). Sequences were assembled using Sequencher 4.8 (Gene Codes).
Mating type and serotype identification

We initially determined the mating type and serotypes by PCR analysis of the STE20, SXI1a, SXI2a, PAK1, GPA1, and CNA1 genes as described previously (Lengeler et al. 2001). However, some AD hybrid strains are untypable because of PCR failure, which may be attributable to DNA sequence variation in the primer complementary regions. STE20 is not only serotype-specific but also mating-type specific (Frasier et al. 2004), which makes it an appropriate marker for the determination of serotype and mating type of C. neoformans. New STE20 primers were designed on the basis of DNA sequence alignment of STE20 gene from strains JEC21 (D), JEC20 (A), H99 (Ax), and 125.91 (A; Table 2). Genomic DNA from strains JEC21, JEC20, H99, and KN99a served as positive controls, and sterile water was used as negative control. The mating type of most AD hybrid isolates was confirmed through mating assays on V8 and MS media (see the section Mating assays). Even with newly designed primers, the mating types of some AD hybrid isolates were still undeterminable because of negative PCR results. CGH analysis (see RESULTS) demonstrates that loss of the chromosome on which the MAT locus is located in AD hybrids results in negative PCR amplification of STE20. Thus, the use of a single marker, such as STE20, can in some cases misclassify AD hybrid isolates as haploid strains (without FACS data) or diploid isolates of a single serotype (combined with FACS analysis).

To resolve this problem, we also designed primers from other chromosomes for serotype determination. These primers were chosen from regions that are conserved between the serotype A and D genomes and following PCR can generate bands of distinct size from serotype A and D strains. Thus, this PCR assay generated two distinct bands from the serotype A- and D-derived genome of the AD hybrid isolates. To design these primers, we compared the serotype A and D chromosomes by using BLASTn (Altschul et al. 1997) and aligned them by using the ACT tool (Carver et al. 2005).

### MLST and phylogenetic analysis

Previously, 12 housekeeping genes (MPD1, TOP1, MP88, CAP59, URE1, PLB1, CAP10, GPD1, TEF1, SOD1, LAC1, and IGS1) were included in MLST for population genetics studies of C. neoformans serotype A strains (Litvintseva et al. 2003, 2006, 2007). For C.

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### Table 2 Primers used in this study

| Primer sequence, 5'-3' | Target gene | Chr, (D)/(A) | Amplicon size, bp | Purpose |
|-----------------------|-------------|--------------|-------------------|---------|
| GGAAGGGTCTAGTGAAGTCTCCTTC | IGS (A) | 2 (A) | 1092 | PCR and MLST sequencing |
| ATCCATGACCCCCAGGTTCACAGAT | IGS (D) | 2 (D) | 1076 | PCR and MLST sequencing |
| TGGAAGGGTCTAGTGAAGTCTCCTTC | URE1 (A) | 14 (A) | 1093 | PCR and MLST sequencing |
| TCTAGCTGCGAGTAGTTATTATCAC | URE1 (D) | 8 (D) | 1094 | PCR and MLST sequencing |
| CAGACGAGATTTGAAGGAGACTAA | GPD1 (A) | 7 (A) | 889 | PCR and MLST sequencing |
| GTCATAGAATTGAGGGTACCAT | GPD1 (D) | 6 (D) | 897 | PCR and MLST sequencing |
| ATATCGACTGATAGTTATTATCAC | LAC1 (A) | 8 (A) | 779 | PCR and MLST sequencing |
| GGCATGTCAGATGTTATTATCAC | LAC1 (D) | 7 (D) | 901 | PCR and MLST sequencing |
| AATAACCCAGATCCCCAGACTAAG | MPD1 (A) | 8 (A) | 1162 | PCR and MLST sequencing |
| CCCCAATATCCTTGGAGTCTTAA | MPD1 (D) | 7 (D) | 1190 | PCR and MLST sequencing |
| GCCTACTATCATGCGGGTACCTTAC | STE20 (Aa) | 5 (A) | 457 | Mating type/serotype determination |
| CTCATTCTATCCACCGGTTACCTTAC | STE20 (Ax) | 5 (A) | 330 | Mating type/serotype determination |
| CACATCTCAGTGCAGCTTATTTAAC | STE20 (Dd) | 4 (D) | 728 | Mating type/serotype determination |
| AGTCTCAGTCTCGATGTTATATAC | STE20 (Dx) | 4 (D) | 592 | Mating type/serotype determination |
| Intergenic region | 7 (D)/8 (A) | 526 (D)/472 (A) | Serotype determination |
| Intergenic region | 12 (D)/4 (A) | 840 (D)/1128 (A) | Serotype determination |
| Intergenic region | 1 (D)/1 (A) | 227 (D)/227 (A) | Quantitative real-time PCR |
| Intergenic region | 4 (D)/1 (A) | 277 (D)/277 (A) | Quantitative real-time PCR |
| Intergenic region | 9 (D)/1 (A) | 343 (D)/343 (A) | Quantitative real-time PCR |

The amplicon size is based on the DNA sequences of strains JEC21 (serotype D) and H99 (serotype A) (A), serotype A; (Aa), serotype A, MATa; Ax, serotype A, MATa; (D), serotype D; (Da), serotype D, MATa; Dx, serotype D, MATa; MLST, multilocus sequence typing; PCR, polymerase chain reaction.
neuromorphic serotype D, no MLST typing system has been developed, although the LAC1 and URA5 genes have been used to study the population structure of serotype AD strains with a TA cloning strategy (Xu et al. 2000, 2002, 2003). Because AD hybrid isolates have both the serotype A and D genomes in the yeast cells, the previous MLST primers for serotype A amplify both serotype A and D alleles; therefore, the mixed amplicons should be separated by TA cloning. To test a large number of AD hybrid isolates, we designed a serotype-specific MLST system to amplify a specific serotype A or D allele from the AD hybrid isolates.

Taking advantage of 10% to 15% sequence diversity between serotype A and D, we chose primers in serotype-specific regions to amplify the MLST loci on the basis of DNA sequence alignments between strains H99 (Broad Institute) and JEC21 (GenBank accession no. NC_006670-NC_006683). The serotype-specific primers for the five genetic markers are listed in Table 2.

The sequence type is defined as a unique DNA sequence produced by the concatenation of five genetic markers. Here, each AD hybrid isolate is regarded as two composite genomes (serotype A and serotype D). DNA sequences were aligned by ClustalW (Thompson et al. 1994), and alignment was confirmed by MUSCLE (Edgar 2004) and manual inspection. The aligned and verified DNA sequences were then imported into Mega 3.0 (Kumar et al. 2004) and PhyML (Dereeper et al. 2008) to determine the phylogenetic organization of the aligned DNA sequences. The phylogenetic organization was edited in Figure-Tre (http://tree.bio.ed.ac.uk/software/figtree).

Mating assays
For mating assays, strains were pregrown on YPD medium for 2 days. The cells were then mixed on V8 or MS agar medium and incubated at 25° for several days. Hyphae and basidiospore formation was assessed by light microscopy every other day.

Comparative genome hybridization
A C. neoformans array designed at Washington University in Saint Louis, Missouri, is widely used for gene expression analysis. However, given that the serotype A and D genomes share approximately 85% to 90% sequence identity, most array probes could hybridxize with both genomes in the AD hybrid isolates. In addition, probes were designed on the basis of cDNA sequences for expression analysis. To select serotype-specific probes to study the genome organization of the AD hybrid isolates, we compared the probes from the serotype A and D genomes by using BLASTn (e value = 0.0001). Only the probes (70-mers) that exactly matched one serotype but diverged from the other serotype more than 35% were defined as serotype-specific probes. A total of 2140 and 1312 probes distributed over the 14 Cryptococcus chromosomes were selected as serotype A- and D-specific probes for CGH analysis, respectively.

Genomic DNA hybridization was performed as previously described (Alonso-Monge et al. 1999). Basically, genomic DNA was ultrasonicated to generate ~500-bp fragments and purified with a DNA Clean and Concentrator kit (Zymo Research, CA). Then, 5 μg of DNA was used for Cy-3 DUTP or Cy-5 DUTP labeling reactions with the Random Primer/Reaction Buffer mix (BioPrime Array CGH Genomic Labeling System; Invitrogen, Carlsbad, CA). Labeled DNA from the sample (AD hybrid isolate) and the control (equal volume of H99 and JEC21 genomic DNA) was competitively hybridized to microarray slides containing 70 mer oligonucleotides for the C. neoformans JEC21 and H99 genomes (Lin et al. 2006). After hybridization, arrays were scanned with a GenePix 4000B scanner (Axon Instruments) and normalized using GenePix Pro V.4.0 and Gene-spring tools. The log2 ratio of R635/R532 value was plotted according to the H99 and JEC21 genome in Microsoft Excel.

Pulsed-field gel electrophoresis of C. neoformans
Yeast cells from a purified single colony were grown overnight in 5 ml of YPD medium at 30°. Then, 100 ml of the culture were added to 50 ml of yeast nitrogen base minimal medium and grown to OD600 0.5 at 30° in a shaking incubator. To suppress capsule formation, the medium was supplemented with 1 M NaCl. Yeast cells were pelleted, washed three times in 0.5 M NaCl/50 mM EDTA (pH = 8.0), resuspended in 9.5 ml of water with 0.5 ml of β-mercaptoethanol, and incubated for 1 hr at 37° with gentle mixing. Cells were pelleted again, resuspended in 4 ml of spheroplasting solution (1 M sorbitol/10 mM EDTA/100 mM sodium citrate), 100 µl of lysing enzyme solution (20 mg of Sigma-Aldrich [St. Louis, MO] lysing enzyme from Trichoderma harzianum in 100 µl of spheroplasting solution) was added, and cells were incubated for 2 hr at 37° with gentle mixing.

Complete spheroplasting was checked by mixing ~10 ml of cell suspension with the same volume of 10% (vol/vol) SDS and gently pipetting to monitor the increase in viscosity that results from the release of high molecular weight DNA. Spheroplasts were pelleted at 1800 rpm for 10 min at 4°, washed twice with ice-cold spheroplasting solution, resuspended in the remaining spheroplasting liquid, and distilled to a concentration of 10^9-10^10 cells/ml. The spheroplasts were then mixed with three volumes of low-melting agarose (1% low melting agarose [Bio-Rad] in 0.125 M EDTA [pH 8.0]; 50°) and poured into molds. Agarose was solidified for 30 min at room temperature and for an additional hour at 4°.

Agarose plugs containing spheroplasts were removed from the molds and spheroplasts were lysed for 24 hr at 55° in lysis solution (0.5 M EDTA/10 mM Tris HCl [pH 10]/1% Sarcosyl). Plugs were washed three times for 15 min each with 0.5 M EDTA (pH 8.0) and stored in 0.5 M EDTA at 4°. A standard 1% gel (1% pulse field-certified agarose in 0.5 × 90 mM Tris HCl/64.6 mM boric acid/2.5 mM EDTA [pH 8.3]) was run in a Bio-Rad CHEF DRII apparatus. Plugs were placed into the wells of a standard 13 × 14 cm gel, sealed with low-melting agarose, and solidified for 30 min at 4°. The gel was run in 0.5 × 90 mM Tris HCl/64.6 mM boric acid/2.5 mM EDTA (pH 8.3) buffer at 12° with the following settings: initial A-time 75 s, final A-time 150 s, start ratio 1.0, run time 30 hr, mode to 10. Initial B-time 150 s, final B-time 300 s, start ratio 1.0, run time 54 hr, mode to 11. The voltage was set to 125 V.

Spot assay to determine sensitivity to antifungal drugs
Sensitivity to the antifungal drugs FK506 and fluconazole was assessed using a spot growth dilution assay. In brief, after growing in liquid YPD medium overnight at 30°, yeast cells were counted at OD600 and diluted with liquid YPD medium to the cell density OD600 = 0. Yeast cells were serially diluted and spotted on YPD plates with 1 µg/ml FK506 or 0.5 µg/ml fluconazole, respectively. After growth at 30° and 37° for 48 hr, the YPD plates with FK506 were examined for cell growth. The YPD plates with fluconazole were examined for cell growth after growth at 30° for 48 hr.

RESULTS
Most C. neoformans AD hybrid isolates harbor the rare a mating type
Of the 64 C. neoformans isolates included in this study, 32 exhibit a diploid profile, and 32 isolates were determined to be haploid based...
on FACS analysis (Figure 1). The genes in the C. neoformans MAT locus are classified into three categories: serotype-specific, mating type-specific, and serotype/mating type-specific genes (Frasier et al. 2004). The STE20 gene that governs cell polarity at different stages of mating of C. neoformans is serotype/mating type-specific (Frasier et al. 2004; Nichols et al. 2004). Previously, STE20 has been used for mating type and serotype determination (Lengeler et al. 2001; Lin et al. 2007); however, we were unable to PCR amplify the STE20 gene from some AD hybrid isolates with existing primers. This may be partially explained by the sequence specificity of the STE20 primers. We redesigned the STE20 primers for serotype/mating type determination (Table 2). The mating type/serotype of 64 C. neoformans strains were identified by serotype- and mating type-specific PCR amplification of the STE20 gene using these new primers (Table 1).

Of the 32 diploid isolates (on the basis of FACS analysis), 24 are AD hybrids as determined by evidence of amplification of the STE20 genes from both serotypes (Figure 1, Table 1, and Figure S2). Among them, 12 are aADx (39%), seven are oADx (23%), and five are oADA (16%; Table 1). No aADA isolates were identified. The MATa allele was identified in approximately 71% (17/24) of the AD hybrid isolates. Another eight isolates exhibited similar DNA content as diploid control strains in FACS but yielded only one serotype/mating type PCR product of the STE20 gene, which suggests that these isolates are diploids of a single serotype due to genome duplication, or AD hybrids with either loss or divergence of the STE20 gene. We also designed primers on other chromosomes to amplify PCR products of distinct sizes from each serotype. The primers chosen from conserved regions between serotype A and D chromosomes can amplify two fragments of distinct size from the AD hybrid isolates but only a single fragment from haploid or diploid mono serotype isolates. The PCR analysis allows the discrimination of AD hybrid isolates from both haploid and diploid isolates of single serotype. Together with the CGH analysis, we found that one of the eight isolates having similar DNA content as diploid isolates was a serotype DD diploid, whereas the other seven isolates were AD hybrid isolates that lost one copy of the chromosome on which the MAT locus is located (Chr 5 and Chr 4, respectively; see the section Extensive chromosome loss in C. neoformans AD hybrid isolates). Among the seven AD hybrid isolates, five lost the serotype A Chr 5 (4 “-ADx” and 1 “-ADα”), and two have lost the serotype D Chr 4 (one “aAD-” and one “oAD-”). The loss of one copy of the sex-related chromosome results in a failure to amplify the corresponding STE20 gene.

Given that 17 of the 24 (71%) AD hybrid strains of known serotype/mating type in our study harbor the rare MATa allele (Table 1), our results suggest that AD hybrid isolates are a reservoir preserving the rare MATa mating type. Previously, other investigators found that serotype A MATa isolates are extant in Botswana (Litvintseva et al. 2007), but the MATa allele in the AD hybrid isolates has a broader distribution, including isolates from the United States and Europe (Table 1). Of note, at least three clinical isolates (aADx) were recovered from patients born in or having traveled to Africa.

**Serotype A MATa genome of AD hybrid isolates originates from Botswana**

To analyze genetic variation of serotype A and D genomes of the AD hybrid isolates, we designed serotype-specific primers for MLST analysis by using DNA sequence comparisons between the JEC21 and H99 genomes (Table 2). To enable comparison with haploid serotype A and D strains, we also sequenced the five markers from 32 haploid serotype D strains and included five markers of serotype A strains from a previous study (Litvintseva et al. 2006). MLST identified two groups corresponding to the serotype A and D populations.
Each group contained both haploid strains as well as AD hybrid strains, which suggests that AD hybrid isolates result from hybridization of serotype A and D strains. MATa predominates in C. neoformans natural populations, and Litvintseva et al. identified three serotype A lineages, VNI, VNII, and VNB (Litvintseva et al. 2006), with MATa isolates being strictly grouped in the VNB lineage (Litvintseva et al. 2003, 2006). Our MLST analysis based on five genetic markers grouped the serotype A MATa and MATa-containing genomes into the VNB and VNI lineages, respectively (Figure 2). This strict distribution of the serotype A genome in the AD hybrid was not associated with the mating type of the serotype D genomic contribution. Our study further supports that the serotype A MATa genome of AD hybrid isolates originates from Botswana and that the serotype A MATa genome possibly has a global distribution.

**Serotype D population has high recombination**

In the MLST analysis based on five markers, we classified 62 serotype D genomes (including 29 AD hybrid isolates, 32 serotype D haploid isolates, and 1 serotype DD diploid isolate) into 31 STs (Table S1). In the MLST analysis we identified 40 STs among 123 serotype A genomes, including 100 from haploid strains in the previous study by Litvintseva et al. 2006 and 23 serotype A genomes from AD hybrid isolates in our study. The genetic diversity of the serotype D population is significantly greater than the serotype A population (t-test; \( P = 0.0139 \)). The greater genetic diversity of serotype D is supported by low bootstrap values in the phylogenetic organization (Figure 2). The phylogenetic organization based on five markers of serotype D is distinct from each other (Figure S3). In addition, recombination is observed in the serotype D population based on the informative paired allele graphs of three MLST markers (LAC1, GPDI, and MPD1), showing that all four possible combinations are observed for alleles of any two genes (AB, ab, Ab, and aB; Figure S4). No recombination was observed in the serotype A population analyzed here.

To further characterize the possible roles recombination may play in the population structure of the serotype D population, we used

![Figure 2](image-url)

**Figure 2** Phylogenetic organization of C. neoformans serotype A (left) and serotype D (right). This analysis was based on the concatenation of 5 MLST markers: IGS, URE1, GPDI, LAC1, and MPD1. The five loci were specifically amplified and sequenced from the serotype A and D genomes of the AD hybrid isolate by the use of serotype-specific primers. For comparison with haploid strains, 32 serotype D haploid strains, 1 serotype DD diploid strains, and 7 serotype A strains (VNI, VNII, and VNB lineages) from a previous study (Litvintseva et al. 2006) were included in the phylogenetic analysis. The concatenated DNA sequences were aligned by the use of ClustalW and MUSCLE. The phylogenetic organization was constructed with Mega 3.1 via the neighbor-joining method. The MATa strains are shown in red. The serotype and mating type of each isolate are indicated in brackets.
MultiLocus software to examine the percentages of compatible loci and indices of association ($I_\lambda$). The analysis of both the percentage of compatible loci and $I_\lambda$ produce $P$-values that, if significant, reject the null hypothesis of recombination. If the $P$-values are not significant, the null hypothesis cannot be rejected and recombination can be inferred. When we performed the recombination test by using the MultiLocus software, the results suggested that random recombination of the serotype D population could not be rejected on the basis of the genotyping profile of three loci ($\text{LAC1, GPD1, and MPD1}$; $I_\lambda = 0.0531401$ and $P = 0.3$). These results suggest that the serotype D population has a greater level of recombination than serotype A. Both opposite- and same-sex mating, which are observed in serotype D strains in the laboratory, may contribute to the population dynamics of serotype D isolates.

Although the serotype D population is highly diverse based on MLST, most serotype D $\text{MAT}^a$ strains, including haploid strains and the serotype D genome from AD hybrid isolates, are grouped closely together to form a subcluster containing strains of both mating types (Figure 2). Recombination tests via the use of the MultiLocus software suggested that random recombination in this subgroup could not be rejected on the basis of the genotyping profile of three loci ($\text{LAC1, GPD1, and MPD1}$; $I_\lambda = 0.00114586$ and $P = 0.398$). This phylogenetic organization is similar to that of the serotype A $\text{MAT}^a$ population and suggests that the serotype D $\text{MAT}^a$ genome in AD hybrid isolates originate from serotype D haploid strains. However, only four serotype D $\text{MAT}^a$ isolates were included in the phylogenetic analysis, and the geographic origins of these isolates are unknown. This does not allow us to speculate about whether the serotype D $\text{MAT}^a$ isolates are also limited to some regions. However, the serotype D genome of the AD hybrid isolates has a wide distribution including China, Italy, and Kuwait. In addition, two serotype D genomes from two AD hybrid isolates (AD6-93 and AD7-97: $\text{aADx}$) distinctly form a unique cluster that was supported by a high bootstrap value (Figure 2). This unique lineage in the serotype D population may be a sign of an ancient hybridization event followed by several independent evolutionary paths of the AD hybrid. Alternatively, haploid isolates corresponding to this lineage may remain to be found or have become extinct since hybridization.

Although serotype A and D genomes of the AD hybrid isolates are phylogenetically grouped with serotype A and D haploid isolates, respectively, and exhibited comparable phylogenetic organization, most AD hybrid isolates have unique sequence types for both serotype A and D genomes in comparison to the haploid isolates. Only two AD hybrid isolates (AD7-75 and AD2-71) share the same serotype D ST (ST24) with two serotype D haploid isolates (AD2-95 and AD2-96; Table S1). This finding suggests that after the initial hybridization event, the AD hybrid isolates underwent independent evolution, which drives the AD hybrid isolates to become genetically diverse from their parental haploid isolates while still being phylogenetically related.

**Extensive chromosome loss in C. neoformans AD hybrid isolates**

Previous investigators found that AD hybrid isolates are morphomorphic or homoygous at some loci but are heterozygous at many loci, including the $\text{MAT}$ locus (Lengeler et al. 2001). However, information on the genomic organization of AD hybrids is limited. C. neoformans AD hybrid isolates are known to result from hybridization of serotype A and serotype D genomes; however, the genome organization of serotype AD strains is mainly unclear. Hu et al. (2008) used tiling array-based CGH to identify chromosome loss events in three AD hybrid isolates (two $\text{aADx}$ and one $\text{oADa}$) that have opposite mating types of serotype A and D genomes. In our study, whole-genome hybridization analysis of the AD hybrid isolates demonstrated that C. neoformans AD hybrid isolates ($\text{aADx}$, $\text{oADa}$, and $\text{nADx}$) have extensive chromosome loss of either the serotype A or serotype D chromosomes (Figure 3). Both whole and partial chromosome loss is observed, which may generate novel chromosomes (Figure 3). Chromosome loss based on CGH analysis was confirmed by PCR (Figure S2). The PCR primers located in regions conserved between the serotype A and D genomes were used to amplify a polymorphic region with size variation between serotype A and D, resulting in two different products of distinct sizes. Chromosome loss in AD hybrid isolates results in a single product in this PCR assay.

Our FACS results demonstrated that all of the AD hybrids have about the same DNA content as diploid isolates (Figure 1), although in CGH analysis we detected extensive chromosome loss (Figure 3). To explore this, CGH analysis found an approximately twofold log2 ratio (corresponding to copy number) of the chromosomes with loss of heterozygosity in comparison with the chromosomes that have both the serotype A and D parental chromosomal homologs (Figure 3), and thus the monomorphic chromosome has undergone duplication after the loss of one parental chromosome (Figure 3).

To test whether the remaining chromosome was duplicated, we used quantitative real-time PCR to investigate the copy number of genes located on chromosomes that were lost. Our real-time PCR primers were selected from conserved regions of the serotype A and serotype D genomes and can amplify both serotype A and D alleles (~250 bp). We compared the copy number of genes that are homozygous (one serotype chromosome was lost) and those that are heterozygous. An identical gene copy number was obtained from heterozygous vs. homozygous genes (Figure 4), suggesting that duplication of the remaining chromosome follows chromosome loss from one serotype in the AD hybrid isolates. This further explains the greater log2 ratio of the remaining chromosome and the similar DNA content of the AD hybrid isolates as diploid strains based on FACS analysis.

**The genome of C. neoformans AD hybrid isolates is highly dynamic**

A previous study found that AD hybrid strain KW5 lost serotype D Chr 1, Chr 6, and Chr 7 and serotype A Chr 14 (Hu et al. 2008). However, an MLST marker from serotype A Chr 14 ($\text{URE1}$) was positive in isolate KW5 (Figure 5). We first considered that this might be a false-positive PCR because of DNA contamination. We repurified strain KW5 and isolated eight single colony subclones. Genomic DNA was isolated from these subclones for PCR, sequencing, and CGH analysis. PCR amplification of the serotype A $\text{URE1}$ gene failed for clones C1, C4, C5, C6, and C8, whereas clones C2, C3, and C7 yielded a PCR product (Figure 5). Serotype/mating type specific PCR demonstrated that all eight clones shared the same mating type and serotype ($\text{oADa}$; Figure 5). This led us to speculate that the KW5 strain may be a mixture of two groups of isolates, one group that has lost and another that has retained the serotype A Chr 14. However, this does not exclude possible contamination by another isolate. We therefore sequenced in all eight clones the most variable MLST marker, $\text{IGS}$, which is unique in strain KW5 compared with the other isolates in our strain collection. The $\text{IGS}$ sequences of the eight clones are identical, unique, and the same as the original KW5 strain. Furthermore, CGH demonstrated that clone C5 lost the serotype A Chr 14 compared to clone C7 (Figure 5). They have highly similar...
chromosomal profiles for all other chromosomes. Based on these analyses, we conclude that the populations derived from isolate KW5 are a mixture of two groups of isolates, one with serotype A Chr 14 and another group that has lost serotype A Chr 14.

We further explored the genome dynamics of another AD hybrid isolate, CDC228 (aADα), which is self-fertile on mating media (Lengeler et al. 2001). In a previous study, 21 CDC228 progeny obtained on V8 mating medium were dissected (Lengeler et al. 2001). Four CDC228
progeny were investigated here by CGH. Each progeny has a distinct genome organization (Figure 6). The parental strain CDC228 lost only serotype D Chr 1, but the progeny isolate P5 additionally lost serotype D Chr 3 and Chr 12 and serotype A Chr 5, Chr 8 (partial), Chr 10 (partial), and Chr 13 (Figure 6). The CGH results of strain CDC228 and its progeny, and KW5 and its derivatives, show that the AD hybrid genome is not stable and that genomic organization is dynamic and may contribute to phenotypic variation. We further tested the progeny of CDC228 and observed that some progeny lost the resistance to FK506 characteristic of CDC228 (Figure 7; as mentioned in the section Many C. neoformans AD hybrid isolates are resistant to the antifungal drug FK506).

Many C. neoformans AD hybrid isolates are resistant to the antifungal drug FK506

We tested the susceptibility of C. neoformans AD hybrids to two antifungal drugs, fluconazole and FK506. Our results demonstrate that most AD hybrids are resistant to FK506 (Figure 7), although they remain sensitive to fluconazole (data not shown). The self-fertile AD hybrid isolate CDC228 is also resistant to FK506. Given that the CDC228 progeny have extensive chromosome loss and reorganization (Figure 6), the question is whether the genome reorganization contributes to phenotypes such as resistance to antifungal drugs. Spot dilution growth assays on FK506 medium demonstrated that some CDC228 progeny maintain whereas others lost the FK506 resistance phenotype (Figure 7). We further studied the chromosome organization of some progeny of distinct phenotype by separating chromosomes on a CHEF gel and observed that the CDC228 progeny P5, which lost the FK506 resistance phenotype, is missing some chromosomes compared with the parental strain and other progeny (Figure 6). This novel genomic organization may result in the loss of resistance to FK506.
Hybridization and its role in population dynamics and speciation have been widely investigated in plants and animals (Dowling and Secor 1997; Soltis and Soltis 2009). It has also been shown that intraspecific and interspecific hybridization of plant fungal pathogens leads to host expansion and adaptation to novel environmental niches (Hamilton et al. 2009; Moon et al. 2004; Olson and Stenlid 2002); in addition to sexual reproduction, this may be an important biological force driving evolution of plant fungal pathogens. In comparison, hybrids of human fungal pathogens are less characterized and understood. Both intervarietal (AD hybrid) and interspecies hybrids (C. neoformans x C. gattii hybrid) of C. neoformans have been either isolated from nature or patients or constructed in the laboratory (Bovers et al. 2006, 2008; Cogliati et al. 2006; Lengeler et al. 2001; Lin et al. 2007; Tanaka et al. 1996; Viviani et al. 2006; Whelan and Kwon-Chung 1986). A recent study in Europe revealed that approximately 19% of cryptococcosis cases are caused by AD hybrid isolates (Viviani et al. 2006). Given the high prevalence of C. neoformans hybrids in human infections, hybridization may play a fundamental role in the evolution of this fungal pathogen. Indeed, mixed infection of serotype A and D isolates as well as isolates of the same serotype but with different molecular types or mating types have been documented (Desnos-Ollivier et al. 2010; Mandal et al. 2005; Sullivan et al. 1996). The high rate of mixed infection may provide an opportunity for hybridization in the host in addition to routes to hybridization in nature.

Comparative gene genealogical analyses suggest that AD hybrids result from hybridization of serotype A and D strains (Xu et al. 2000, 2002, 2003). In our study, the serotype A and D genomes of the AD hybrid isolates group with haploid serotype A and D isolates, respectively, on the basis of phylogenetic analysis of five MLST markers (Figure 2). The consensus phylogenetic organization of haploid isolates and AD hybrid isolates confirms a recent hybridization origin of the AD hybrid. However, the serotype A and D genomes of most AD hybrid isolates are genotypically distinct from the haploid isolates, except for two AD hybrid isolates (AD7-75 and AD2-71) that share the same sequence type (ST24) with two serotype D haploid isolates (AD7-91 and AD7-85; Table S1). The consistent phylogenetic organization but genotypic divergence between the AD hybrid and haploid isolates supports the role of hybridization in the origin of the AD hybrid but also suggests ongoing independent evolutionary
trajectories. In addition, the serotype D genomes of two AD hybrid isolates (AD6-93 and AD7-97; aADx) exclusively form a basal lineage in the serotype D population (Figure 2). Although haploid serotype D isolates belonging to this lineage may remain to be found or be extinct, the AD hybrid isolates in this lineage may result from an ancient hybridization event followed by independent evolution to form a distinct lineage. However, the serotype A genomes of these two AD hybrids are classified into the VNB lineage and are closely related to isolate Bt63 (MATa) from Botswana.

On the basis of MLST, three lineages (VNI, VNII, and VNB) have been previously identified in the serotype A population (Litvintseva et al. 2006). The serotype A MATa and MATa genomes of AD hybrid strains group into the VNB and VNI lineages, respectively (Figure 2). Given that the serotype A MATa VNB isolates are mainly identified in Botswana (Litvintseva et al. 2006), it is hypothesized that the serotype A MATa genome of the AD hybrid strains originates from Botswana, whereas the MATa genome of the AD hybrid has a global distribution. Our result is in accord with a previous analysis of the serotype A genome of the AD hybrid isolates, in which three MLST markers (CAP10, URE1, and GPD1) were used (Litvintseva et al. 2007). Similarly, the MATa genomes from both the serotype D haploid isolates and AD hybrid isolates group closely to form a subcluster (Figure 2). The limited number of serotype D MATa isolates included in the phylogenetic analysis and the unknown geographic origin of the serotype D MATa haploid isolates (which are very likely from Europe where most serotype D isolates were identified) do not allow us to conclude that serotype D MATa genomes of the AD hybrid originate from a restricted geographic region as is the case with the serotype A MATa genome. Identification and analysis of more serotype D MATa isolates will be necessary to address this question in the future.

Given the strict distribution of serotype D MATa strains in the phylogenetic organization and the limited number of MATa strains in nature, the serotype D population is assumed to be relatively clonal except the subcluster containing both MATa and MATa isolates. Indeed, random recombination in the subcluster containing both MATa and MATa isolates could not be rejected on the basis of genotyping with three MLST markers (LAC1, GPD1, and MPD1; \( I_X = 0.00114586 \), and \( P = 0.398 \)). This finding reveals ongoing recombination in this subcluster that may be attributable to sexual reproduction of strains of opposite mating types. However, our recombination test using MultiLocus software also suggests ongoing recombination in the haploid serotype D population on the basis of three MLST markers (LAC1, GPD1, and MPD1; \( I_X = 0.0531401 \), \( P = 0.3 \)). In addition, the serotype D population (31 STs among 62 genomes) has a significantly greater genetic diversity than the serotype A population (40 STs among 123 genomes; \( t \)-test, \( P = 0.0139 \)). High genetic diversity within a population is often associated with sexual reproduction in a sexually compatible microorganism (Campbell and Carter 2006). Although the serotype D population is dominated by MATa isolates, sexual reproduction between MATa isolates is observed in the laboratory (Lin et al. 2005). Same-sex mating may contribute to increase genetic diversity of the serotype D population in nature. Same-sex mating of serotype A strains has not been observed in the laboratory thus far, but population genetic evidence supports that it occurs in nature, but may be less frequent. If so, this could explain in part why the serotype D population is more diverse than serotype A.

Of the 31 AD hybrid isolates included in this study, 24 have an identified serotype/mating type. The serotype/mating type of the remaining seven AD hybrid isolates could not be determined because of the loss of the mating type chromosome of one serotype (Table 1). Of the 24 AD hybrid isolates with designated serotype/mating type, seven had only one mating type (MATa; aADx) and 17 had both mating types (12 aADx and 5 aADA) present in their diploid genomes. aADx hybrid isolates originate from unisexual mating and hybridization of MATa strains of serotype A and D strains (Lin et al. 2007). On the contrary, aADA and aADA hybrid isolates are products of opposite-sex mating and hybridization events. Thus, there have been at least three independent AD hybridization events inferred based on only the analysis of MAT configuration. No aADA hybrid isolates have been identified in nature thus far, although they can be constructed in the laboratory (Lin et al. 2008). This may be explained by the rarity of MATa isolates in the population or that same-sex mating is less common between MATa isolates (Lin et al. 2008).

Previous authors found that AD hybrids are heterozygous at some loci but homozygous at others (Cogliati et al. 2006; Lengeler et al. 2001). On the basis of serotype-specific CGH, we provide strong evidence that the homozygous chromosomes in AD hybrid isolates

Figure 7 AD hybrid isolates exhibit phenotypic plasticity. Most AD hybrid isolates are resistant to the antifungal drug FK506 (left) and the progeny of the self-fertile strain CDC228 (aADx), which is resistant to the antifungal drug FK506, have altered FK506 resistance (right). After growing for 48 hr on YPD + FK506 (1 \( \mu \)g/ml) medium at 37°C, five AD hybrid isolates (SG290, NC34-21, MMRL1365, MMRL774, and MMRL752) exhibited resistance compared with the H99 and JEC21 haploid strains (left). After growing for 48 hr on YPD + FK506 (1 \( \mu \)g/ml) medium at 37°C, progeny P5 was sensitive to FK506 (right).
result from the loss of the chromosome from the genome of one serotype (Figure 3). Most chromosomes, including the mating type chromosome, have the potential to be lost. However, chromosomes 2, 6, 7, and 8, on which the five MLST markers are located, seem to be more stable than others on the basis of our analyses. On the basis of the chromosomal synteny between the serotype A and D genomes (Figure S1), chromosomes 3 and 11 are less collinear because of a translocation. Chromosome 9 has a significant inversion between the two serotypes. In addition, Chr 1 seems to be more frequently lost from the serotype D genome, which results in homoyzogosity for the serotype A Chr 1 typically by reduplication (Figure 3). Disomy for Chr 1 in haploid strains (1N+1) has been recently associated with heteroresistance to fluconazole in C. neoformans serotype A (Sionov et al. 2010). However, we found that AD hybrid isolates are generally not more resistant to fluconazole than haploid isolates (data not shown).

Our results demonstrate that the genomic organization of the AD hybrid isolates is highly complex and diverse rather than a simple combination of genomes from the two parents, and this highly diverse genomic organization significantly increases the population diversity of C. neoformans. Moreover, we observed that the genomic organization of AD hybrid isolates is highly dynamic with great plasticity. Mitotic propagation of AD hybrids can generate population diversity, including the loss or duplication of whole and partial chromosomes as well as reorganization of the chromosomes. The AD hybrid isolate KW5 is an example demonstrating the complexity of genome dynamics in C. neoformans AD hybrids. Our study demonstrated that populations derived from the KW5 isolate are a mixture of at least two groups, one with and the other without the serotype A copy of Chr 14 (Figure 5). When and how the serotype A Chr 14 was lost from the subpopulation is not known. Isolate CDC228 (aADo) is another example of genomic diversity in an AD hybrid. This strain is self-fertile, and no mating partner is required in this process. Although the viability of the meiotic progeny is low, the surviving progeny are highly diverse in genomic organization (Figure 6). This is an efficient route for C. neoformans to generate diversity, which may contribute to the ability to adapt to changing environments. The parental AD hybrid isolate CDC228 is resistant to the antifungal drug FK506, but some of its progeny have lost this phenotype. We speculate that this may also generate progeny with increased fitness in response to other selective pressures not analyzed. Our study suggests that caution should be exercised when studying AD hybrids, and especially that the analysis of one single colony should be avoided.

Hybridization is an important mechanism for speciation in angiosperms by generating novel gene combinations and distinct phenotypes (Paun et al. 2009; Rieseberg et al. 2003; Soltis and Soltis 2009). In many cases, hybridization between different species or subspecies is prevented by prezygotic barriers, or the hybrid is sterile or exhibits reduced fertility because of genetic divergence between the parental strains. In some cases, however, the hybrid can also be stably fertile via either hybridization followed with genome doubling (allopolyploidy) or introgressive hybridization without a change in chromosome number (homoploidy) (Gompert et al. 2006; Rieseberg et al. 2003; Salazar et al. 2010). Then, the stable fertile hybrid can be productively isolated from its parental species by postzygotic barriers, and is therefore regarded as a novel species (Gutierrez-Marcos et al. 2003). In C. neoformans, there is no evidence that the AD hybrid has undergone genome doubling after hybridization between serotype A and D strains. Our FACS and CGH analyses found that the AD hybrid isolates have similar DNA content as the diploid, but not tetraploid, isolates. Thus, allopolyploid speciation does not likely occur in AD hybrids. Although introgression between serotype A and D genomes has been detected, it is on a smaller scale and occurred historically (Kavanaugh et al. 2006). The authors of a previous study identified an approximately 40-kb region (~0.2% of the genome) that is nearly identical (98.5%) between the serotype A and D genomes (Kavanaugh et al. 2006). This identity island may result from a non-reciprocal transfer event from serotype A to D approximately 2 MYA (Kavanaugh et al. 2006). This introgression event likely occurred during the hybridization process of serotype A and D strains and was facilitated by transposon-mediated DNA transfer. This is the only introgression region identified in C. neoformans thus far. No AD hybrid genomic sequence is currently available, mostly because of the difficulties of assembling the sequence reads into two genomes that share 85% to 90% sequence identity. Thus, the potential recombination events in the AD hybrid isolates are largely unknown. However, given the high percentage of transposons in the C. neoformans genome (Loftus et al. 2005), extensive introgression and recombination events may occur in AD hybrids during the long evolutionary period, which may be able to generate homoploidy to result in speciation. We observed that AD hybrid isolates have unique sequence types compared with their parental isolates, although they are highly related phylogenetically. This finding suggests limited gene flow between AD hybrids and the haploid population. Although the causes of genetic divergence in AD hybrid isolates are not known, this may be an early sign of speciation of the AD hybrid. In addition, most AD hybrid isolates cannot mate with haploid parental isolates in the laboratory, which may result in reproductive isolation and ultimately speciation of the AD hybrid.

We observed that most AD hybrid isolates are resistant to the antifungal drug FK506 (Figure 7). Together with the previous observation that AD hybrids are resistant to ultraviolet light (Lin et al. 2007), we hypothesize that the AD hybrid has hybrid vigor that may be attributable to the genome combination and genomic dynamics. Regulation of the genomic organization is a rapid and efficient route for human fungal pathogens to generate favorable phenotypes. For example, C. neoformans becomes resistant to fluconazole by chromosomal duplication (Sionov et al. 2010), and aneuploidy and the formation of novel chromosomes contributes to antifungal drug resistance in C. albicans and Candida glabrata (Polakova et al. 2009; Selmecki et al. 2006; Sionov et al. 2010).

The AD hybrid genome is highly dynamic. We detected both partial and complete chromosome loss and duplication in some AD hybrid isolates (Figure 3). Progeny P5 of strain CDC228 has a distinct genomic organization resulting from the partial loss of the serotype A copy of Chrs 14 and 10 (corresponding to serotype D Chr 8 and Chr 10, respectively) and the complete loss of some other chromosomes (Figure 6). Partial chromosome loss may result in genome arrangement or the formation of novel chromosomes. Indeed, the loss of resistance to the antifungal drug FK506 in P5 may be attributable to this distinct genomic organization.

The AD hybrid generates population diversity efficiently. Although most progeny of AD hybrids are sterile and have low viability, the surviving progeny may have considerable hybrid vigor. Hybrid vigor may promote the global spread of AD hybrids, which may contribute to the preservation and exportation of the rare MatA genome. This is common in some plant species. The Senecia species are native in France, but the interspecific hybrid resulting from hybridization and allopolyploidy has spread to England (Abbott and Lowe 2004). Tragopogon is native to Euroasia, but the hybrid species invaded into North America (Soltis and Soltis 1999).

In summary, we characterized in this study the genetic diversity and genomic organization of the C. neoformans AD hybrid. Our
results demonstrate that the AD hybrid originated from hybridization between serotype A and D strains and arose multiple times via independent hybridization events. The AD hybrid genome is highly dynamic, providing an efficient way for the AD hybrid to generate population dynamics and hybrid vigor. The hybrid vigor of the AD hybrid may facilitate it to spread globally, which may preserve the rare $MATa$ allele. The potential contribution of the AD hybrid to speculation is an interesting topic for future investigation.

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LITERATURE CITED

Abbott, R.A., and A. J. Lowe, 2004 Origins, establishment and evolution of new polyploid species: Senecio cambrensis and S. eboracensis in the British Isles. Biol. J. Linn. Soc. Lond. 82: 8.

Alonso-Monge, R., F. Navarro-Garcia, G. Molero, R. Diez-Orejas, M. Gustin et al., 1999 Role of the mitogen-activated protein kinase Hog1p in morphogenesis and virulence of Candida albicans. J. Bacteriol. 181: 3058–3068.

Altenschul, S. F., T. L. Madden, A. A. Schaffer, J. Zhang, Z. Zhang et al., 1997 Gapped BLAST and PSI-BLAST: a new generation of protein database search programs. Nucleic Acids Res. 25: 3389–3402.

Arnold, M. L., 1997 Natural Hybridization and Evolution, Oxford University Press, New York.

Barchiesi, F., M. Cogliati, W. C. Esposto, E. Spreghini, A. M. Schimizzi et al., 2005 Comparative analysis of pathogenicity of Cryptococcus neoformans serotypes A, D and AD in murine cryptococcosis. J. Infect. 51: 10–16.

Bowers, M., M. Hagen, E. E. Kuramae, M. R. Diaz, L. Spanjaard et al., 2006 Unique hybrids between the fungal pathogens Cryptococcus neoformans and Cryptococcus gattii. FEMS. Yeast Res. 6: 599–607.

Boyes, M., F. Hagen, E. E. Kuramae, H. L. Hoogveld, F. Dromer et al., 2008 AIDS patient death caused by novel Cryptococcus neoformans x C. gattii hybrid. Emerg. Infect. Dis. 14: 1105–1108.

Brasier, C. M., D. E. Cooke, and J. M. Duncan, 1999 Origin of a new Phytophthora pathogen through interspecific hybridization. Proc. Natl. Acad. Sci. U S A 96: 5878–5883.

Campbell, L. T., and D. A. Carter, 2006 Looking for sex in the fungal pathogens Cryptococcus neoformans and Cryptococcus gattii. FEMS. Yeast Res. 6: 588–598.

Carver, T. J., M. R. Mutherman, M. Berberan, M. A. Rajandream, B. G. Barrett et al., 2005 ACT: the Artemis Comparison Tool. Bioinformatics 21: 3422–3423.

Chaturvedi, V., J. Fan, B. Stein, M. J. Behr, W. A. Samsonoff et al., 2002 Molecular genetic analyses of mating pheromones reveal intervarietal mating or hybridization in Cryptococcus neoformans. Infect. Immun. 70: 5225–5235.

Cigliati, M., M. C. Esposto, A. M. Tortorano, and M. A. Viviani, 2006 Cryptococcus neoformans population includes hybrid strains homozygous at mating-type locus. FEMS. Yeast Res. 6: 608–613.

de Barros Lopes, M., J. R. Bellon, N. J. Shirley, and P. F. Ganter, 2002 Evidence for multiple interspecific hybridization in Saccharomyces sussu stricto species. FEMS. Yeast Res. 1: 323–331.

Dereeper, A., V. Guignon, G. Blanc, S. Audic, S. Buffet et al., 2008 Phylogeny.fr: robust phylogenetic analysis for the non-specialist. Nucleic Acids Res. 36: W465–W469.

Desnos-Ollivier, M., S. Patel, A. R. Spaulding, C. Charlier, D. Garcia-Hermoso et al., 2010 Mixed infections and in vivo evolution in the human fungal pathogen Cryptococcus neoformans. mBio 1: 10.

Dowling, T.E., and C. Secor, 1997 The role of hybridization and introgression in the diversification of animals. Annu. Rev. Ecol. Syst. 28: 27.

Dromer, F., E. Gueho, O. Ronin, and B. Dupont, 1993 Serotyping of Cryptococcus neoformans by using a monoclonal antibody specific for capsular polysaccharide. J. Clin. Microbiol. 31: 359–363.

Dromer, F., S. Mathoulin-Pelissier, O. Launay, and O. Lortholary, 2007 Determinants of disease presentation and outcome during cryptococcosis: the CryptoA/D study. PLoS Med. 4: e21.

Edgar, R. C., 2004 MUSCLE: multiple sequence alignment with high accuracy and high throughput. Nucleic Acids Res. 32: 1792–1797.

Fraser, J. A., S. Diezmann, R. L. Subaran, A. Allen, K. B. Lengeler et al., 2004 Convergent evolution of chromosomal sex-determining regions in the animal and fungal kingdoms. PLoS Biol. 2: e384.

Gompert, Z., J. A. Fordyce, M. L. Forister, A. M. Shapiro, and C. C. Nice, 2006 Homoploid hybrid speciation in an extreme habitat. Science 314: 1923–1925.

Gutierrez-Marcos, J. F., P. D. Pennington, L. M. Costa, and H. G. Dickinson, 2003 Imprinting in the endosperm: a possible role in preventing wide hybridization. Philos. Trans. R. Soc. Lond. B Biol. Sci. 358: 1105–1111.

Hamilton, C. E., S. H. Faeth, and T. E. Dowling, 2009 Distribution of hybrid fungal symbionts and environmental stress. Microb. Ecol. 58: 408–413.

Hu, G., I. Liu, A. Sham, J. E. Stajich, F. S. Dietrich et al., 2008 Comparative hybridization reveals extensive genome variation in the AIDS-associated pathogen Cryptococcus neoformans. Genome Biol. 9: R41.

Ikeda, R., S. Nishimura, A. Nishikawa, and T. Shinoda, 1996 Production of agglutinating monoclonal antibody against antigen 8 specific for Cryptococcus neoformans serotype D. Clin. Diagn. Lab. Immunol. 3: 89–92.

Inderbitzin, P., R. M. Davis, R. M. Bostock, and K. V. Subbarao, 2011 The ascomycete Verticillium longisporum is a hybrid and a plant pathogen with an expanded host range. PLoS ONE 6: e18260.

Kabasawa, K., H. Itagaki, R. Ikeda, T. Shinoda, K. Kagaya et al., 1991 Evaluation of a new method for identification of Cryptococcus neoformans which uses serologic tests aided by selected biological tests. J. Clin. Microbiol. 29: 2873–2876.

Kavanaugh, L. A., J. A. Fraser, and F. S. Dietrich, 2006 Recent evolution of the human pathogen Cryptococcus neoformans by intervarietal transfer of a 14-gene fragment. Mol. Biol. Evol. 23: 1879–1890.

Kumar, S., K. Tamura, and M. Nei, 2004 MEGA3: Integrated software for Molecular Evolutionary Genetics Analysis and sequence alignment. Brief. Bioinform. 5: 150–163.

Kwon-Chung, K. J., 1975 A new genus, Filobasidiella, the perfect state of Cryptococcus neoformans. Mycologia 67: 1197–1200.

Kwon-Chung, K. J., 1976 A new species of Filobasidiella, the sexual state of Cryptococcus neoformans B and C serotypes. Mycologia 68: 943–946.

Kwon-Chung, K. J., and J. E. Bennett, 1978 Distribution of a and alpha mating types of Cryptococcus neoformans among natural and clinical isolates. Am. J. Epidemiol. 108: 337–340.

Lengeler, K. B., G. M. Cox, and J. Heitman, 2001 Serotype AD strains of Cryptococcus neoformans are diploid or aneuploid and are heterozygous at the mating-type locus. Infect. Immun. 69: 115–122.

Lengeler, K. B., D. S. Fox, J. A. Fraser, A. Allen, K. Forrester et al., 2002 Mating-type locus of Cryptococcus neoformans: a step in the evolution of sex chromosomes. Eukaryot. Cell 1: 704–718.

Lewontin, R.C., and C. Birch, 1966 Hybridization as a source of variation for adaptation to new environments. Evolution 20: 22.

Lin, X., C. M. Hull, and J. Heitman, 2005 Sexual reproduction between partners of the same mating type in Cryptococcus neoformans. Nature 434: 1017–1021.

Lin, X., J. C. Huang, T. G. Mitchell, and J. Heitman, 2006 Virulence attributes and hyphal growth of C. neoformans are quantitative traits and the MAta allele enhances filamentation. PLoS Genet. 2: e187.

Lin, X., A. P. Litvinetsve, K. Nielsen, S. Patel, A. Floyd et al., 2007 alpha AD alpha hybrids of Cryptococcus neoformans: evidence of same-sex mating in nature and hybrid fitness. PLoS Genet. 3: 1975–1990.

Lin, X., K. Nielsen, S. Patel, and J. Heitman, 2008 Impact of mating type, serotype, and ploidy on the virulence of Cryptococcus neoformans. Infect. Immun. 76: 2923–2938.
