Calcineurin Is Essential for *Candida albicans* Survival in Serum and Virulence

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Calcineurin is a calcium-activated protein phosphatase that is the target of the immunosuppressants cyclosporin A and FK506. In T cells, calcineurin controls nuclear import of the NF-AT transcription factor and gene activation. In plants and fungi, calcineurin functions in stress responses (e.g., temperature, cations, and pH) and is necessary for the virulence of the fungal pathogen *Cryptococcus neoformans*. Here we show that calcineurin is also required for the virulence of another major fungus that is pathogenic to humans, *Candida albicans*. *C. albicans* calcineurin mutants had significantly reduced virulence in a murine model of systemic infection. In contrast to its role in *C. neoformans*, calcineurin was not required for *C. albicans* survival at 37°C. Moreover, *C. albicans* calcineurin mutant strains exhibited no defects in known *Candida* virulence traits associated with host invasion, including filamentous growth, germ tube formation, and adherence to and injury of mammalian cells. *C. albicans* calcineurin mutant strains failed to colonize and grow in the kidneys of infected animals and were unable to survive when exposed to serum in vitro. Our studies illustrate that calcineurin has evolved to control aspects of the virulence of two divergent fungal pathogens via distinct mechanisms that can be targeted to achieve broad-spectrum antifungal action.

*Candida albicans* is the most common cause of invasive fungal disease. Interestingly, *C. albicans* is also a component of the normal human biota that resides on mucosal surfaces, including the gastrointestinal tract and the oral and vaginal mucosae. Superficial colonization of immunocompetent or immunocompromised patients by the fungus can lead to oral thrush, vulvovaginitis, esophagitis, and cutaneous infections. Patients who have undergone abdominal surgery or organ transplantation, have late-onset diabetes, are taking broad-spectrum antibiotics, or have neutropenic are all at increased risk for serious invasive disease (6, 43, 58). Often, the first stage of invasive disseminated infection is candidemia, a bloodstream infection with *Candida* that can be associated with foreign-body catheters or entry of yeast through a damaged intestinal mucosa (reviewed in reference 7). Candidemia can progress to acute or chronic hematogenously disseminated candidiasis, in which organisms such as the liver, kidney, and spleen are involved. During the progression from a common commensal organism to a pathogen, *C. albicans* faces a number of environments in vivo that require it to sense and respond to the extracellular milieu for survival.

A great deal is known about the events leading to the pathogenic transition of *C. albicans*, especially about those pathways important for tissue invasion: the morphological switch from yeast-like growth to filamentous growth, expression of adherence molecules in the cell wall, and protease production (7, 11, 25, 42, 54, 55). Germ tube production, an early stage of filamentation, plays a multifunctional role in virulence. The germ tube is the site of localization for proteins such as the adherence protein Hwp1 (53) and the secreted aspartyl proteinases Sap4 to Sap6 (28). Also, filament production by the fungus inside macrophages causes lysis of the macrophages and thus allows the fungus to survive this host defense (14, 38). Afilamentous mutants of *C. albicans* have markedly attenuated virulence, although it is not known whether filamentous structures are themselves essential or whether it is the loss of specific proteins associated with the hyphal state that causes the attenuation of virulence (5, 15, 23, 26, 38). Disruption of genes encoding the adhesins and proteases that are expressed by filamentous *C. albicans* is known to reduce the virulence of *C. albicans*. Mutant strains lacking the adhesin Hwp1 have attenuated virulence in an oroesophageal model of candidiasis in immunodeficient mice (56). Mutants lacking the Sap1 to -6 proteases also have attenuated virulence and, in some instances, demonstrate either a reduced capacity to invade host tissues or enhanced clearance from host organs (19, 29, 50). Many of the proteins involved in germ tube formation, adhesion, and host cell damage are dependent on host cues, such as high pH and high temperature, for their expression (16, 19, 47, 48), and thus the ability to sense these environmental cues is essential for the virulence of *C. albicans*.

A candidate for such a sensor is calcineurin, a highly conserved protein that is important for mediating stress responses in both plants and fungi. Calcineurin is composed of two distinct subunits, the catalytic A and the regulatory B subunit. Calcineurin B binds to an α-helical extension of calcineurin A distinct from the active site. In response to calcium influx,
calmodulin binds to calcineurin A and blocks the action of the autoinhibitory C-terminal domain of calcineurin A, resulting in the formation of the active calcineurin complex (reviewed in references 2, 27, 34, and 49). Calcineurin is important for responses to environmental temperature stress, including low (Arabadopsis thaliana and Schizosaccharomyces pombe) and high (Saccharomyces cerevisiae) temperatures (35, 44, 62). Calcineurin also functions in response to cation stress in A. thaliana, C. neoformans, and Saccharomyces cerevisiae (9, 13, 35, 44). Additionally, calcineurin is also important for response to and growth in alkaline pH media in both S. cerevisiae and C. neoformans (44, 51). Thus, we hypothesized that calcineurin might function in C. albicans to sense host cues to enable host tissue invasion and disseminated infection.

Calcineurin has recently been identified in C. albicans and shown to play a key role in enabling cells to survive membrane perturbation by drugs that target ergosterol biosynthesis (12, 45). The resulting model is that calcineurin function becomes essential when cell integrity is compromised. Based on these observations, we hypothesized that calcineurin might also be necessary to maintain the viability of C. albicans during the stresses inherent in infection. Here we show that calcineurin is indeed important for virulence in a murine tail vein injection model of candidiasis. Histological examination of the renal parenchyma of infected mice reveals that calcineurin is required to establish tissue infection. However, analysis of the known virulence traits important for organ invasion did not uncover an explanation for the inability of strains lacking calcineurin to produce renal infection. Instead, we found that calcineurin is important for survival in serum. Thus, we conclude that calcineurin plays a key role early in infection, during candidemia, a critical stage of the C. albicans infectious cycle.

MATERIALS AND METHODS

Strains and media. Strains used in this study are listed in Table 1. Strains were grown on either yeast extract-peptone-dextrose (YPD) rich medium, defined synthetic dextrose medium, prepared as previously described (52), or fetal bovine serum (FBS) (Sigma). FK506 (Fujisawa), niacin (Sigma), or thiamine (Sigma) was added to media at the concentrations described.

To create prototrophic calcineurin mutants for in vivo analysis, plasmid pGEM-HIS1 was linearized by SacI-EcoQ digestion (59). The linearized fragment was transformed into the calcineurin B mutant strains DAY364 and DAY365 (both ura3::ura3 arg4::arg4 his1::his1 cnb1::URA3/cnb1::ARG4) (12) by the lithium acetate procedure (30), and transformants were selected on synthetic dextrose media lacking histidine. The prototrophic strains JRB64 (from DAY364) and JRB71 (from DAY365) were isolated after this transformation and have the genotype ura3::ura3 arg4::arg4 HIS1/cnb1 cnb1::URA3/cnb1::ARG4. His+ isolates were grown on YPD medium for several generations and then tested on synthetic dextrose media to ensure that the HIS1 gene had been stably integrated into the genome.

Virulence in a murine model. The virulence of the cnb1/cnb1 mutant C. albicans strains was tested in a murine tail vein injection model. Two tests were performed, one with 5 animals in each injection group and a second with 10 animals in each group. Strains were grown overnight in Sabouraud’s agar, and a light suspension of each was made in Sabouraud’s broth. One drop of this suspension was added to 50 ml of Sabouraud’s broth, which was then shaken at 250 rpm overnight at 35°C. Cells were washed three times, resuspended in phosphate-buffered saline (PBS), and counted with a hemocytometer. The cells were then diluted to yield 3 × 10^8 CFU per 200 μl and injected intravenously through the tail vein into outbred ICR mice. Inoculum concentrations were verified by culture and ranged from 2.46 × 10^6 to 3.12 × 10^6 cells per inoculum in the 5-mouse experiment (Fig. 1A) and from 1.74 × 10^6 to 4.2 × 10^6 cells per inoculum in the 10-mouse experiment (Fig. 1B). The end point for survival was determined by clinical signs of pain or distress, at which point animals were sacrificed by CO2 asphyxiation.

Adherence to epithelial cells. Adherence of yeast cells was measured using a buccal epithelial cell (BEC) protocol modified from the work of Panagoda et al. and of Kimura and Pearsall (33, 46). Wild-type, calcineurin B mutant, and reconstituted strains were grown overnight in liquid YPD medium, washed twice with PBS, and then resuspended in PBS at a concentration of 10^7 cells per ml. BECs were harvested from healthy human subjects by gently rubbing the inside of the cheek with sterilized cotton-tipped swabs (Fisher), and cells were suspended in PBS. These cells were washed twice in PBS and then resuspended in PBS at a concentration of 10^6 cells per ml. For the adherence tests, 0.5 ml of the epithelial suspension and 0.5 ml of the yeast inoculum were mixed in Eppendorf tubes and incubated at 37°C for 1 h with shaking. As a control, 0.5 ml of the BEC suspension or 0.5 ml of the C. albicans suspensions were mixed with 0.5 ml of PBS and incubated for 1 h at 37°C without shaking. Following incubation, all samples were filtered (Ultra-ware 25-mm microfiltration system; Kontes Glass Company, Vineland, N.J.) through 10-μm-pore-size polycarbonate filters (Milipore, Bedford, Mass.). To remove unattached yeast cells, the filters were washed twice with 10 ml of PBS. Filters were then removed from the filtration apparatus, gently pressed onto glass slides, and then carefully pulled off of the slides to leave the epithelia and attached yeast cells attached to the slides. Slides were air dried, heat fixed, Gram stained (Fisher Protocol) Gram Stain Stain Stat Pack (Fisher Scientific, Pittsburgh, Pa.), and mounted with Permount (Fisher Scientific). The number of C. albicans cells attached to the first 100 BECs was counted.

Endothelial cell injury. The endothelial cells used in this study were harvested from human umbilical cord veins by the method of Jaffe et al. (32). Cells were grown in M-199 medium supplemented with 10% FBS, 10% defined bovine calf serum, and 2 mM L-glutamine with penicillin and streptomycin. Cells were grown in 24-well plates coated with gelatin and incubated at 37°C under 5% CO2. Endothelial cell damage caused by SC5314, JRB64, and MCC85 was measured by a chromium release assay as previously described (20). A total of 10^5 C. albicans cells in RPMI 1640 medium were added to each well in a 24-well tissue
culture plate and incubated for 3 h. All experiments were performed in triplicate and repeated twice.

**Histology.** Healthy BALB/c mice were infected by tail vein injection with ~3 \( \times 10^8 \) cells of strain CAF2 (wild type), JRB64 (cnb1/cnb1), JRB71 (cnb1/cnb1), or MCC85 (cnb1/cnb1 + CNB1) per 200 \( \mu \)l. Two animals from each infection group were sacrificed on day 1 postinfection. Both kidneys were harvested from each sacrificed animal and fixed in zinc formalin fixative (Polysciences, Inc., Warrington, Pa.). Fixed kidneys were processed and embedded in paraffin. Renal sections were subsequently prepared and subjected to staining with methylene silver (Grocott’s modification) to visualize yeast cells or with hematoxylin and eosin to visualize host cells. Processing and staining were performed at the Duke University Medical Center Immunohistology Research Laboratory, Durham, N.C. Renal sections were examined under a Nikon Eclipse E400 microscope at \( \times 200 \), and photographs were taken with a Nikon Coolpix digital camera.

**Survival in serum.** Fungal strains were grown overnight in rich liquid medium at 30°C, washed twice in PBS, and counted with a hemocytometer. Cells were diluted into FBS, porcine serum, or sheep serum (Sigma) at a concentration of 2.5 \( \times 10^3 \) cells/ml and allowed to grow for the indicated time at 30°C with agitation. In certain experiments, the vitamins niacin and/or thiamine (Sigma), yeast extract (Difco), or peptone (Difco) was added to the serum at the specified concentrations, after which the serum solution was filtered through a Millex-GV syringe-driven filter unit with a pore size of 0.22 µm (Millipore) and then incubated with yeast. Following incubation, cells were diluted appropriately and 100 µl was plated onto YPD plates such that there were approximately 100 to 500 CFU per plate, except in the case of the calcineurin mutants of *C. albicans*, which with no dilution had only 1 to 2 CFU after 24 h of incubation in serum.

**Statistical analysis.** Analysis of the virulence experiments was performed with Kruskal-Wallis and Dunn’s multiple comparison tests. Endothelial cell injury was analyzed with a single-factor analysis of variance. Analysis of the data for yeast survival in serum and adherence to BECs was performed by using an unpaired, two-tailed Student t test.

**RESULTS**

*C. albicans* calcineurin mutants have attenuated virulence.

In divergent microorganisms, calcineurin senses and responds to a variety of stresses similar to those present in mammalian hosts. In particular, calcineurin is essential for the virulence of the pathogenic basidiomycete *C. neoformans* (13, 21, 44). This led us to hypothesize that calcineurin might also be important for the virulence of *C. albicans*. To test this hypothesis, we generated an isogenic series of prototrophic calcineurin mutants lacking the regulatory B subunit (cnb1/cnb1) and a strain in which calcineurin B was reconstituted (cnb1/cnb1 + CNB1) (Table 1) (12). Virulence studies were performed in which groups of 5 or 10 ICR outbred mice were infected intravenously with ~3 \( \times 10^6 \) cells of the wild-type strain (CAF2), the cnb1/cnb1 mutant strains (JRB64 and JRB71), or the cnb1/cnb1 + CNB1 reconstituted strain (MCC85).

All of the mice infected with the wild-type *C. albicans* strain succumbed to lethal infection by day 8 postinfection, whereas the survival of mice infected with either of two different independent calcineurin B mutant strains was dramatically prolonged (Fig. 1). Of 30 animals infected with the calcineurin B mutant strains, 27 survived until the experiment was terminated on day 30 postinfection. Three of the 30 mice infected with the calcineurin mutant strains died during the course of the experiment, but the survival time for 2 of the 3 mice was longer than that of any of the animals infected with the wild-type strain. Importantly, reintroduction of the wild-type calcineurin B gene via integration at the *HIS1* locus restored virulence, and all of the mice infected with the reconstituted strain died during the course of both virulence experiments, with a survival rate that approached that of animals infected with the wild-type strain (Fig. 1). In summary, calcineurin plays a critical role in virulence in the murine tail vein injection model of candidiasis.

**Calcineurin is dispensable for high-temperature growth of *C. albicans*.** Calcineurin is essential for optimal growth of *C. neoformans* strains at 37°C, and as a consequence, *C. neoformans* calcineurin mutant strains are avirulent (13, 21, 44). Wild-type and calcineurin mutant strains of *C. albicans* were analyzed for growth at 30, 37, and 42°C on rich and minimal media (Fig. 2 and data not shown). The *C. albicans* cnb1/cnb1 calcineurin B mutant strains did not exhibit a growth defect at any temperature tested (22 to 42°C) (Fig. 2 and data not shown). Thus, in contrast to the requirements of *C. neoformans*, calcineurin is not required for growth at elevated temperatures in *C. albicans*, and its impact on virulence must therefore be attributable to a distinct molecular mechanism.
Calcineurin is not required for germ tube formation. Several virulence attributes, including germ tube formation, adherence to host cells, and host cell invasion, have been shown to play a role in tissue colonization and infection by *C. albicans* (5, 15, 19, 23, 26, 36, 38). Adhesins and proteases important for host cell adherence and invasion are localized to the germ tube (28, 53, 57), and the hyphal structure itself may promote invasion of host cells. High temperature, alkaline pH, and other undefined in vivo cues signal a change in morphology from yeast to filamentous growth in *C. albicans*. Two pathways have been implicated in this dimorphic switch, and additional regulatory inputs likely also participate (reviewed in references 10 and 42). Because calcineurin is important for growth under high-salt, alkaline pH, and elevated-temperature conditions in other fungi (9, 13, 35, 44, 51, 55), we tested whether calcineurin controls the hyphal growth of *C. albicans* and thereby contributes to virulence.

Hyphal differentiation was analyzed in vitro on solid or liquid media. There were no differences between the calcineurin mutants and the wild-type strain in the ability to filament on solid, filamentation-inducing media (Lee's, 10% serum, and spider media) at 30 and 37°C (data not shown). The calcineurin mutants did exhibit a filamentation defect at 25°C, but only on spider medium (data not shown). Calcineurin mutants also exhibited no filamentation defect in liquid media containing 10% FBS either with regard to the number of cells producing germ tubes or with regard to the length of the germ tubes and hyphae formed (Fig. 3A and data not shown). In summary, calcineurin is not required for germ tube formation or filamentous growth under most conditions in vitro.

**Calcineurin is not necessary for host cell adherence.** *C. albicans* adheres to host cells to avoid clearing from the mucosal surface, and adherence is an essential initiating event for establishing infection during invasive candidiasis (54). An in vitro test was performed with human BECs to compare the

![FIG. 2. Calcineurin is not required for growth of *C. albicans* at 37°C. Wild-type and calcineurin B mutant strains of *C. albicans* and *C. neoformans* were grown at 30 and 37°C on rich medium for 72 h and photographed.](image)

**FIG. 3.** Calcineurin is not required for germ tube formation or adherence to epithelial cells. (A) Wild-type (WT) (SC5314), calcineurin mutant (cnb1/cnb1) (JRB64), and calcineurin reconstituted (cnb1/cnb1 + CNB1) (MCC85) strains were grown in rich liquid medium with 10% FBS at 37°C for 2.5 h, and germ tubes produced by all three strains were visualized by Nomarski optics and photographed at a magnification of ×200. At this stage, 89.6% of wild-type, 82.5% of calcineurin mutant, and 87.0% of reconstituted cells had produced germ tubes (only the difference between the data for wild-type and calcineurin mutant cells was statistically significant [P = 0.03]). (B) A total of 5 × 10^6 wild-type (SC5314) or calcineurin mutant (JRB64) cells were incubated with 5 × 10^6 human BECs. Cells were Gram stained, visualized with a Zeiss Axioscop 2 plus microscope, and photographed at a magnification of ×400. Arrows indicate representative *C. albicans* cells attached to epithelial cells. Bars, 25 μm.
abilities of calcineurin B-null mutant and wild-type cells to adhere to epithelial cells. C. albicans strains were incubated with human BECs in PBS, and the number of adherent yeast cells was quantified after 1 h of coincubation. The number of yeast cells per BEC for wild-type versus calcineurin mutant strains did not differ significantly (81.5 yeast cells per 100 BECs for the wild type and 142 yeast cells per 100 BECs for the calcineurin mutant; \( P = 0.21 \)) (Fig. 3B). This assay was also performed in TC199 cell culture medium, and no significant difference was noted between wild-type and mutant adherence, even though adherence was greatly enhanced in TC199 medium compared to that in PBS (data not shown). Calcineurin, therefore, is not essential for C. albicans adherence to host cells.

**Calcineurin is largely dispensable for endothelial cell injury.** The next stage of tissue infection by C. albicans involves invasion of host cells to promote host cell injury, evade the host immune defense, and scavenge nutrients from host cells. Here, endothelial cell injury was monitored with a \(^{51}\text{Cr} \) release assay (20). Wild-type C. albicans, a calcineurin-null mutant, and the reconstituted mutant strain were incubated with umbilical vein endothelial cells preloaded with \(^{51}\text{Cr} \), and release of \(^{51}\text{Cr} \) from endothelial cells was measured after a 3-h incubation with fungal cells. The calcineurin mutant strain exhibited a defect in endothelial cell injury in this assay (20% less than that with the wild type) (Fig. 4). This statistically significant (\( P = 0.001 \) for comparison with the wild type) but modest defect might contribute to the observed virulence defect of the mutant in the murine whole-animal model. However, it is unlikely that the defect in endothelial cell injury displayed by the calcineurin mutant strains is sufficient to explain the avirulent phenotype in the whole-animal model. Supporting this assertion, the extent of endothelial cell injury caused by the reconstituted mutant strain was not statistically significantly different from that caused by the wild type (\( P = 0.9 \)) and was only slightly different from that caused by the calcineurin mutant strain (\( P = 0.029 \)) (Fig. 4), even though this strain is nearly as virulent as the wild type in the murine model. It should also be noted that, though tested in a more susceptible mouse model at a lower inoculum, the rim101/rim101 mutant strain, which exhibits a more severe defect in endothelial cell wounding, led to a median survival time of only 10 days (15), compared to the almost avirulent calcineurin mutant strains, which had a much less severe endothelial cell injury defect.

**Calcineurin mutants fail to colonize and grow in the kidneys.** C. albicans infections frequently start as candidemia that can progress to acute hematogenously disseminated candidiasis and result in the infection of many organs. The murine tail vein injection model mimics aspects of this disease progression in that the fungus is first introduced into the bloodstream and then disseminates to cause disease in organs including the kidney and liver. In this model of disease progression, acute infection of the kidney appears to be the main cause of attributable mortality (1, 3, 4).

To monitor disease progression in animals infected with either wild-type or calcineurin B mutant strains, and to determine whether the mutant strains might have an in vivo defect in filamentation that was not detected in our in vitro assays, kidneys were harvested and subjected to histopathological analysis. Mice infected with the wild-type strain displayed multiple aggregates of filamentous cells, approximately 75 to 250 \( \mu \text{m} \) in diameter, in the kidney (Fig. 5). Significantly fewer fungal cells were present in the kidneys of mice infected with the calcineurin B mutant strain, and these were typically observed as single cells in the renal section rather than as aggregates (Fig. 5). However, the mutant strains did produce filaments in vivo, thus demonstrating that the calcineurin mutant virulence defect is not likely attributable to a filamentation defect in vivo. On the other hand, these observations indicate that calcineurin is required for C. albicans to initiate and maintain infection in the renal parenchyma. This inability to grow within the kidney likely contributes to the marked attenuation in virulence of the mutant strains.
Calcineurin is essential for survival in serum. Calcineurin is important for virulence and growth within the kidney, yet calcineurin mutants appear to have only minor defects in the known steps preceding tissue invasion. We therefore focused on an even earlier stage of the infection cycle, candidemia. In the murine tail vein injection model of candidiasis, one of the first environments that the yeast cells encounter is the blood. Blood is composed of host cells and plasma, which constitutes about 55% of whole blood. We therefore tested whether calcineurin is important for survival in this medium. This was accomplished by testing the survival of wild-type, calcineurin mutant, and reconstituted strains for persistence in serum.

Calcineurin mutants exhibited a defect in survival in 100% FBS relative to survival of the wild-type and reconstituted strains. After 3 h in serum, there was no difference in growth between the wild-type and calcineurin mutants. However, by 6 h of incubation, the wild-type and reconstituted strain populations had increased six- to eightfold whereas the population of the calcineurin mutant strains had decreased, and this trend continued at 9 h (Fig. 6A). Finally, after 24 h of incubation, the mutant cells were essentially inviable while the wild-type and reconstituted strain populations had increased more than 14,000-fold (Fig. 6B). Thus, we conclude that calcineurin is crucial for the survival of C. albicans in serum. This finding was not specific to FBS, because similar observations were made with porcine, sheep and murine sera (data not shown). In addition, a wild-type strain grown in FBS with 1 μg of the calcineurin inhibitor FK506/ml was as sensitive to serum as the calcineurin B mutant strain. This finding reinforces the conclusion that calcineurin is essential for survival in serum.

Why is calcineurin required for growth in serum? Calcineurin is essential for survival in serum, but it is not known whether serum contains an active component that is killing the fungal cells or lacks a component that is required for the growth and survival of calcineurin mutant cells. We tested whether serum lacks components necessary for the survival of calcineurin mutants by nutrient addition experiments. The growth of wild-type, calcineurin mutant, and reconstituted strains of C. albicans was measured in FBS diluted 1:1 with PBS or YPD rich medium; YPD diluted 1:1 with PBS was used as a control. The growth of strains in the 1:1 dilution of serum with PBS was similar to that in undiluted FBS after 24 h of incubation, but replacement of PBS with YPD liquid medium led to an almost complete recovery of calcineurin mutant growth such that the difference in growth between the wild-type and the calcineurin mutant strains was less than 2-fold, compared to an approximately 1,000,000-fold difference in serum alone at 24 h (Fig. 7A).

YPD medium is composed of peptone, yeast extract, glucose, and water. When tested individually, none of these components, used at the concentrations present in YPD, had any obvious effects on the growth of the calcineurin mutants in serum, but a 10-fold increase in the yeast extract concentration partially complemented the growth defect of the calcineurin mutants (~2-fold difference in fold population growth between the wild type and the calcineurin mutant). Peptone restored growth only weakly (data not shown). Peptone and yeast extract are similar in composition, but several differences exist, most notably in the concentrations of certain vitamins. Niacin, para-aminobenzoic acid, pantothenic acid, pyridoxine, riboflavin, and thiamine are ~25- to 5,000-fold more abundant in yeast extract than in peptone. Supplementation of serum with niacin and thiamine restored the growth of calcineurin mutants in serum, but only at supraphysiologic levels, concentrations...
YPD medium and approximately 1.5/10000 (JRB64) (hatched bars), and reconstituted (MCC85) (open bars) strains were grown for 24 h at 30°C in serum diluted 1:1 with PBS or YPD, or in YPD diluted 1:1 with PBS. Statistically significant differences were noted between the wild-type strain and both the calcineurin mutant strain (P < 0.005) and the reconstituted strain (P = 0.006) but not between the calcineurin mutant and reconstituted strains (P > 0.05) in the serum-PBS experiment. Also, no significantly significant difference in growth was found between the samples in the serum-YPD and YPD-PBS experiments (P > 0.05). (B) Wild-type (WT) (SC5314) and calcineurin mutant (cnb1/cnb1) (JRB64) strains were grown for 24 h at 30°C in serum either alone or supplemented with either 5.95 mg of niacin/ml or 5.3 mg of thiamine/ml. Fold changes in population size were measured by comparing the CFU at the conclusion of the experiments with the initial CFU. Differences in growth between the wild-type and calcineurin mutant strains in both the serum-alone (P < 0.001) and thiamine addition (P = 0.004) experiments were statistically significant. Values below the dashed line represent a fold change below 1, indicating cell death.

FIG. 7. Medium supplementation restores growth of calcineurin mutants in serum. (A) Wild-type (SC5314) (shaded bars), calcineurin mutant (JRB64) (hatched bars), and reconstituted (MCC85) (open bars) strains were grown for 24 h at 30°C in serum diluted 1:1 with PBS or YPD, or in YPD diluted 1:1 with PBS. Statistically significant differences were noted between the wild-type strain and both the calcineurin mutant strain (P < 0.005) and the reconstituted strain (P = 0.006) but not between the calcineurin mutant and reconstituted strains (P > 0.05) in the serum-PBS experiment. Also, no significantly significant difference in growth was found between the samples in the serum-YPD and YPD-PBS experiments (P > 0.05). (B) Wild-type (WT) (SC5314) and calcineurin mutant (cnb1/cnb1) (JRB64) strains were grown for 24 h at 30°C in serum either alone or supplemented with either 5.95 mg of niacin/ml or 5.3 mg of thiamine/ml. Fold changes in population size were measured by comparing the CFU at the conclusion of the experiments with the initial CFU. Differences in growth between the wild-type and calcineurin mutant strains in both the serum-alone (P < 0.001) and thiamine addition (P = 0.004) experiments were statistically significant. Values below the dashed line represent a fold change below 1, indicating cell death.

~1,000-fold higher than those at which they are present in rich YPD medium and approximately 1.5 × 10^5-fold (niacin) (31) to 1 × 10^6-fold (thiamine) (17) higher than those in human blood (Fig. 7B). Niacin and thiamine had no appreciable affect on the growth of calcineurin mutants in serum at −1, 10, and 100-fold their concentrations in YPD (data not shown). Therefore, absence of vitamins might contribute to the growth defect of calcineurin mutants in serum, but other factors must contribute as well.

Candidates for an active lethal factor in serum include complement components, antibodies, or antimicrobial peptides. Size fractionation of serum excluded the possibility that complement and antibodies were the active factor, because PBS that was size fractionated to exclude components of >3 kDa retained the activity that kills calcineurin mutants of C. albicans (data not shown). In addition, serum that was boiled for 15 min and pelleted to remove denatured proteins still retained the ability to kill C. albicans calcineurin mutants (data not shown). Hence, an active component in serum, if any, must be heat stable and <3 kDa.

DISCUSSION
We have discovered that calcineurin, a calcium-regulated signaling enzyme conserved throughout evolution, is essential for the virulence of C. albicans. Sanglard and colleagues have independently reached a similar conclusion (personal communication). We find that calcineurin is not required for hyphal development in vitro or in vivo or for adherence to host cells, two well-studied processes important for virulence. Calcineurin mutants have a modest defect in endothelial cell injury, but it is not likely of sufficient magnitude to confer the marked reduction in virulence. Histopathological analysis revealed that calcineurin is required for dissemination and growth in the renal parenchyma. Calcineurin was found to be necessary for survival in serum, and we propose that calcineurin mutants fail to survive long enough in the bloodstream to disseminate efficiently and cause significant disease in target tissues for mortality such as the kidneys. The necessity of calcineurin for survival in serum is interesting in light of a recent study exploring the expression profile of C. albicans exposed to human blood in vitro or mouse blood in vivo (22). A number of classes of genes were discovered to be upregulated in serum or blood, including genes involved in glycolysis, the glyoxylate cycle, and fatty acid metabolism. It will be of interest to determine whether calcineurin plays a role in regulating any of these processes that could be important for growth and survival in blood.

We propose two models to explain the sensitivity of calcineurin mutants of C. albicans to serum. In the first, serum lacks one or more components necessary for the proliferation of calcineurin mutant cells. In the second, an active factor in serum kills calcineurin mutants. Our studies reveal that 1:1 supplementation of serum with YPD rich medium, addition of 100 mg of yeast extract/ml to serum (10-fold the level in YPD), or addition of supraphysiological levels of the vitamins thiamine and niacin rescues the growth of calcineurin mutants, providing support for the first model. Furthermore, our studies reveal that boiling serum and passing it though a 3-kDa filter does not destroy activity against calcineurin mutants, eliminating an involvement of most serum proteins, including complement and antibodies.

In contrast to the nutrient addition studies, the rapid loss of CFU by the calcineurin mutants in serum suggests an active
killing component. In support of this second hypothesis, calcineurin mutant cells incubated overnight at 30°C in PBS, which lacks nutrients, persist in this medium (data not shown). Thus, the phenotype of the calcineurin mutants cannot simply be due to a lack of nutrients in the serum. The addition of supraphysiologic nutrients to serum in the nutrient addition experiments might simply promote growth and mask the effect of active factors, which could include small antimicrobial peptides (AMPs) with molecular weights below 3 kDa. AMPs, such as the platelet-produced antimicrobial peptides, are known to have antifungal capabilities (61). AMPs have little to no tertiary structure, are heat stable, have broad activity against a number of organisms, including bacteria, fungi, and some viruses, and generally act by disrupting the pathogen membrane (reviewed in references 24 and 60). The membrane disruption activity of the AMPs is reminiscent of the membrane disruption caused by fluconazole, which is lethal to strains lacking calcineurin but only fungistatic to strains with active calcineurin (12). Thus, AMPs in serum might be contributing to the growth defect of calcineurin mutants in serum via a mechanism similar to the azole-hypersensitive phenotype observed with C. albicans cnb1/cnb1 calcineurin mutants. It is also possible that both the active factor and nutrient models are correct and that their effects compound to lead to the rapid killing of calcineurin mutants. Further studies will be necessary to elucidate in detail the role of calcineurin in promoting the growth and survival of C. albicans in serum.

Although calcineurin is highly conserved, calcineurin mutants of the distantly related pathogenic basidiomycete C. neoformans and the closely related ascomycete S. cerevisiae exhibit distinct phenotypes. Calcineurin is required for C. albicans to survive in serum, but not for the survival of C. neoformans or S. cerevisiae (J. R. Blankenship and J. Heitman, unpublished data). In contrast, calcineurin is required for survival at 37°C in C. neoformans but not in C. albicans or S. cerevisiae (13, 21, 44). Thus, the calcineurin signaling cascade has evolved to control aspects of virulence of two divergent pathogens via distinct molecular mechanisms. It is also clear from these findings that one cannot extrapolate from model yeasts or even from other human fungal pathogens to establish the cellular functions of these conserved signaling molecules; rather, studies must be conducted with each organism.

Calcineurin is the target of the immunosuppressive and antifungal drugs cyclosporine A and FK506 (tacrolimus) (37). Less-immunosuppressive analogs of these drugs that retain their antifungal activity have been identified, and several groups have explored the potential of these drugs for anti-
Candida therapy (12, 39–41, 45). These drugs have no effect on the growth of C. albicans and S. cerevisiae in rich media but exhibit a potent fungicidal synergism when combined with azoles (12, 18, 39, 41). Edlund and colleagues have shown that the activity of the calcineurin target Crz1 is important for synergy between calcineurin inhibitors and azoles in S. cerevisiae (18). However, other studies suggest that a Crz1-independent role of calcineurin in responding to endoplasmic reticulum stress is responsible for the synergism between azoles and calcineurin inhibitors in S. cerevisiae (8). In our studies, the growth defect of calcineurin mutants in serum was recapitulated by exposing wild-type cells to the calcineurin inhibitor FK506 in serum. Thus, the utility of calcineurin inhibitors in treating Candida infections could be expanded. These studies highlight the potential use of less-immunosuppressive analogs of calcineurin inhibitors for treatment of candidal infections and illustrate that screening drugs for antifungal activity in serum rather than rich medium has the potential to reveal novel classes of antifungal agents for therapeutic intervention.

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