Candida albicans VMA3 Is Necessary for V-ATPase Assembly and Function and Contributes to Secretion and Filamentation

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The vacuolar membrane ATPase (V-ATPase) is a protein complex that utilizes ATP hydrolysis to drive protons from the cytosol into the vacuolar lumen, acidifying the vacuole and modulating several key cellular response systems in Saccharomyces cerevisiae. To study the contribution of V-ATPase to the biology and virulence attributes of the opportunistic fungal pathogen Candida albicans, we created a conditional mutant in which VMA3 was placed under the control of a tetracycline-regulated promoter (tetR-VMA3 strain). Repression of VMA3 in the tetR-VMA3 strain prevents V-ATPase assembly at the vacuolar membrane and reduces concanamycin A-sensitive ATPase-specific activity and proton transport by more than 90%. Loss of C. albicans V-ATPase activity alkalinizes the vacuolar lumen and has pleiotropic effects, including pH-dependent growth, calcium sensitivity, and cold sensitivity. The tetR-VMA3 strain also displays abnormal vacuolar morphology, indicative of defective vacuolar membrane fission. The tetR-VMA3 strain has impaired aspartyl protease and lipase secretion, as well as attenuated virulence in an in vitro macrophage killing model. Repression of VMA3 suppresses filamentation, and V-ATPase-dependent filamentation defects are not rescued by overexpression of Rim8, Md53, Efg1, Cst20, or Ume6, which encode positive regulators of filamentation. Specific chemical inhibition of Vma3p function also results in defective filamentation. These findings suggest either that V-ATPase functions downstream of these transcriptional regulators or that V-ATPase function during filamentation involves independent mechanisms and alternative signaling pathways. Taken together, these data indicate that V-ATPase activity is a fundamental requirement for several key virulence-associated traits in C. albicans.

Candida albicans is a major opportunistic human fungal pathogen and is responsible for 6.8% of hospital-acquired infections in the United States (1). Despite the availability of several classes of antifungal drugs, attributable mortality, cost of care, and length of stay due to invasive candidiasis remain unacceptably high (2, 3). In addition, resistance to currently available antifungal drugs is emerging (see reference 4 for a review). Therefore, development of new antifungal drug targets remains a critical need. A diverse set of factors contributing to C. albicans virulence have been identified, including the secretion of aspartyl proteases and lipases, filamentation, and biofilm formation (5–8). Understanding the biology and regulation of these processes and pathways may illuminate new candidates for antifungal therapy.

The vacuole is a dynamic acidic organelle found in yeast and plants that is analogous to the mammalian lysosome. It functions in an array of cellular homeostatic processes and thereby plays an important role in stress response, adaptation to novel environments, and cell differentiation (9–13). Furthermore, in C. albicans, intact vacuolar function is important for filamentation and virulence (12–15). Vacular function depends on the maintenance of acidic pH by the vacuolar H-ATPase (V-ATPase), an enzyme complex that functions in organelle acidification across eukaryotes (16, 17). The V-ATPase utilizes hydrolysis of ATP to transport protons from the cytosol into a variety of organelles. V-ATPase-mediated acidification and membrane energization are necessary for important vacuolar functions, including calcium and metal homeostasis (18), cargo sorting and membrane trafficking in endocytic and secretory pathways (19), and drug resistance (20). In Saccharomyces cerevisiae, the V-ATPase is expressed at the vacuolar membrane and the membrane of prevacuolar compartments and the Golgi compartment.
regarding the specific functions of the V-ATPase complex in C. albicans (17).

In S. cerevisiae, the VMA3 gene encodes the c subunit of the V_{o} subcomplex. The c subunit forms a hexameric ring with the c' and c'' subunits of the V-ATPase, which are encoded by VMA11 and VMA16, respectively. This hexameric ring is the main site of proton transport from the cytosol to negatively charged glutamic acid residues in the c ring involves V_{o} subunit a (V_{o}a) (17, 23). The V_{o}a subunit is the only fungal V-ATPase subunit encoded by two isoforms, VP1 and STV1. We have recently shown that V-ATPase pumps containing Vph1p versus Stv1p contribute differently to C. albicans cell biology and virulence-related traits (24). Whereas C. albicans VMA3 has not been previously studied, loss-of-function mutations in the VMA3 gene have been investigated extensively in S. cerevisiae (25–28) and result in pleiotropic effects, including vacuolar alkalization, impaired disassembly of the V_{o} and V_{1}V_{o} complexes (28), and the vma growth phenotype (26).

In this study, we generated a C. albicans tetracycline-regulable VMA3 mutant (tetR-VMA3 strain) in order to analyze the contribution of VMA3 to V-ATPase function, vacuolar morphology, and virulence-related phenotypes. Importantly, we demonstrate that the V-ATPase plays a central role in the induction of C. albicans filamentation; Vma3p-dependent filamentation defects are dominant and independent of several well-characterized filamentation and pH-responsive signaling pathways.

**MATERIALS AND METHODS**

**Identification of Vma3p.** A single potential ortholog of S. cerevisiae Vma3p (http://www.yeastgenome.org/) was identified by a BLASTp search of the Candida Genome Database (http://www.candidagename.org/). The resulting sequence (orf19.5886) was aligned with that of S. cerevisiae Vma3p using the software program MAFFT (29). The alignment was used as a query in the program PRALINE to analyze protein conservation, hydrophobicity, and transmembrane structure (30).

**Strains and media.** Strains used in this study are listed in Table 1. Throughout, unbuffered medium is used to refer to any growth medium where no buffering agents were added to maintain the pH of the medium. For testing of specific pH-dependent phenotypes, medium was buffered to pH 4.0 or 5.0 using 50 mM succinic acid–50 mM Na_{2}PO_{4} or to pH 7.5 where no buffering agents were added to maintain the pH of the medium. Throughout, unbuffered medium is used to refer to any growth medium without buffering agents.

**Growth at various pHs.** The ability of strains to grow on media without a pH buffer was tested on unbuffered YPD with or without doxycycline was assayed using reverse transcriptase PCR (RT-PCR). RT-PCR was performed using the Access RT-PCR system (Promega) according to the manufacturer’s instructions and using the primers RT-VMA3-5Det and RT-VMA3-3Det. The tetracycline-regulable system described by Nakayama et al. (34), with modifications allowing PCR-directed targeting as described by Bates et al. (35), was used to place the remaining VMA3 allele under a tetracycline-regulable promoter. The primers tetVMA3-5DR and tetVMA3-3DR were used to amplify plasmid pDDB57. Strain THE1 was transformed with the resulting PCR amplicon to generate strain VMA3Δ+/tetR (Table 1). Correct genomic integration of the gene disruption cassette was confirmed via PCR using the primers VMA3+/tetR-VMA3-5Det and VMA3+/tetR-VMA3-3Det. Expression of VMA3 in the THE1-CIP10 control strain and the tetR-VMA3 strain after 24 h of growth in unbuffered YPD with or without doxycycline was assayed using reverse transcriptase PCR (RT-PCR). RT-PCR was performed using the Access RT-PCR system (Promega) according to the manufacturer’s instructions and using the primers RT-VMA3-5Det and RT-VMA3-3Det and 5 μg total mRNA as the template. Absence of contaminating DNA was tested in parallel PCR-based reactions. The correct genotype of these strains was confirmed by Southern blotting following standard protocols (36). In brief, genomic DNA prepared from candidate strains was digested with XhoI (New England BioLabs) and run on a 0.6% (wt/vol) agarose gel. A digoxigenin-labeled probe (nucleotide [nt] 800 to 500 of orf19.5886) was prepared from genomic DNA isolated from strain THE1 with the primers VMA3-3Sblt and VMA3-3Sblt (Table 2) and reagents supplied in the PCR DIG probe synthesis kit (Roche).

The ability of strains to grow on medium without a pH buffer was tested on unbuffered YPD and unbuffered complete synthetic medium (CSM) (0.67% yeast nitrogen base without amino acids [YNB], 0.079% complete synthetic mixture, 2% glucose, and 2% agar) with or without doxycycline added. The ability of strains to grow over a pH range was tested on CSM buffered to pH 4.0 to 8.5 with or without doxycycline added. Cells from overnight cultures were washed and counted as previously described (37). Phosphate-buffered saline (PBS) was inoculated with cells from overnight cultures to a starting density of 10^6 cells/ml. Then, a total of five fold dilutions were completed in 96-well plates, and cells were stamped onto agar plates using a multiblot replicator (VP 408H; VP Scientific) and incubated at 30°C for 48 h. Growth at pH 4.0 to 8.5 was also tested in liquid medium: cells from overnight cultures were diluted to an optical density of 600 nm (OD_{600}) of 0.05 in CSM buffered to pH 4.0 to 8.5 and with or without doxycycline added. Then, cells were grown at 30°C using a Biotek Synergy H1M instrument with double orbital shaking at fast speed and 2-mm frequency, with OD_{600} readings taken at 15-min intervals.

**Stress response.** The ability of the tetR-VMA3 strain to grow on media containing various stressors was tested on agar plates with and without
doxycycline. Plates tested for calcium sensitivity were unbuffered YPD with 200 mM CaCl$_2$ and YPD with 200 mM CaCl$_2$ buffered to pH 7.5. Plates used to test the ability of strains to grow on medium containing doxycycline. Plates tested for calcium sensitivity were unbuffered YPD buffered to pH 7.5. Haemagglutinin (HA), hexamethylenetetramine (HMTA). TetR-ura3 strains were CSM (pH 4.0) plates containing either 1 M NaCl, 200 mM caspofungin, 5 mM taurine (YEP)-2% ethanol–3% glycerol. The ability of strains to respond to glycerol as the sole carbon source were unbuffered yeast extract and peptone (pH 7.6) for 5 min. Cells were washed twice with 100 mM phosphate (pH 7.6) for 5 min. Cells were washed twice with 100 mM HEPES–50 mM sodium phosphate–2% glucose (pH 7.6) and resuspended in the same solution. Cells were visualized via differential interference contrast (DIC) and fluorescence microscopy. In order to quantify the vacuolar pH, cells were stained with 2’-7’-bis-(2-carboxyethyl)-5-(and-6)-carboxyfluorescein (BCECF)-acetoxymethyl ester (AM) (from Invitrogen), as described previously (24).

**TABLE 1 C. albicans strains used in this study**

| Strain name or description | Parent strain or description | Relevant genotype$^a$ | Source or reference |
|---------------------------|----------------------------|-----------------------|--------------------|
| BWP17                     | SC5314                     | ura3Δ::ura3Δ arg4Δ::arg4Δ hisΔ::hisΔ VMA3/VMA3 | Wilson et al. 1999 (33) |
| BWP17-VMA3Δ/+             |                           | VMA3::mya3Δ::dpl200-0-URA3-dpl200 | This study |
| THE1                      | CA18                       | ade2Δ::hisGade2Δ::hisG ura3Δ::imm434::ura3Δ::imm434 | Nakayama et al. 2000 (34) |
| THE1-Clp10                | THE1                       | ade2Δ::hisGade2Δ::hisG ura3Δ::imm434::ura3Δ::imm434 | Bernardo et al. 2008 (14) |
| vma3Δ/+ strain            |                            | VMA3::mya3Δ::dpl200-0-URA3-dpl200 ade2Δ::hisGade2Δ::hisG ura3Δ::imm434::ura3Δ::imm434 | This study |
| vma3Δ/+ FOA strain        | VMA3Δ/+ strain             | VMA3::mya3Δ::dpl200 ade2Δ::hisGade2Δ::hisG ura3Δ::imm434::ura3Δ::imm434 | This study |
| tetR-VMA3 strain          |                            | VMA3::mya3Δ::dpl200 ade2Δ::hisGade2Δ::hisG ura3Δ::imm434::ura3Δ::imm434 | This study |
| tetR-VMA3+ NAT1-PENO1-RIM8 strain | tetR-VMA3 strain         | VMA3::mya3Δ::dpl200 ade2Δ::hisGade2Δ::hisG ura3Δ::imm434::ura3Δ::imm434 | This study |
| tetR-VMA3+ NAT1-PENO1-MDS3 strain | tetR-VMA3 strain         | VMA3::mya3Δ::dpl200 ade2Δ::hisGade2Δ::hisG ura3Δ::imm434::ura3Δ::imm434 | This study |
| tetR-VMA3+ NAT1-PENO1-UME6 strain | tetR-VMA3 strain         | VMA3::mya3Δ::dpl200 ade2Δ::hisGade2Δ::hisG ura3Δ::imm434::ura3Δ::imm434 | This study |
| tetR-VMA3+ NAT1-PENO1-EFG1 strain | tetR-VMA3 strain         | VMA3::mya3Δ::dpl200 ade2Δ::hisGade2Δ::hisG ura3Δ::imm434::ura3Δ::imm434 | This study |
| tetR-VMA3+ NAT1-PENO1-CST20 strain | tetR-VMA3 strain         | VMA3::mya3Δ::dpl200 ade2Δ::hisGade2Δ::hisG ura3Δ::imm434::ura3Δ::imm434 | This study |

$^a$ HA, hemagglutinin.

**Vacuolar acidification assays.** Quinacrine staining was performed to visualize acidified vacuoles as described previously (38), with some modifications. First, cells were grown in unbuffered YPD with or without doxycycline for 24 h to ensure complete turnover of vacuolar Vma3p. Then, cells were resuspended in fresh unbuffered YPD, with or without doxycycline, and grown to early log phase. Cells were cooled on ice for 1 min and resuspended in 200 μM quinacrine in YPD buffered with 50 mM sodium phosphate (pH 7.6) for 5 min. Cells were washed twice with 100 mM HEPES–50 mM sodium phosphate–2% glucose (pH 7.6) and resuspended in the same solution. Cells were visualized via differential interference contrast (DIC) and fluorescence microscopy. To quantify the vacuolar pH, cells were stained with 2’-7’-bis-(2-carboxyethyl)-5-(and-6)-carboxyfluorescein (BCECF)-acetoxymethyl ester (AM) (from Invitrogen), as described previously (24).

**V-ATPase assembly and activity assays.** Starter cultures were grown for 6 to 8 h in unbuffered YPD with and without doxycycline. Vacuoles were prepared by resuspending cells in YPD (pH 4.0) with and without doxycycline and growing to an OD$_{590}$ of 1.0 to 1.5 (approximately 18 h). Vacuolar membranes were purified by Ficoll density gradient centrifugation (39). For Western blots, 80 μg of vacuolar protein was separated by SDS-PAGE and transferred to nitrocellulose overnight at 150 mA. The V$_A$, V$_A$ subunit was visualized with a 1:1,000 dilution of anti-human V$_A$ rabbit polyclonal antibody (40); the human V$_A$ antibody cross-reacts with the C. albicans V$_A$ protein. To quantify ATP hydrolysis in purified vacuolar vesicles, vacuolar vesicles (15 μg) were added to an enzymatic assay in which the rate of ATP hydrolysis is coupled to the oxidation of NADH, measured as a loss of absorbance at 340 nm over time (41). Proton transport of purified vacuolar vesicles (30 μg) was measured via quenching of 1 μM 9-amino-6-chloro-2-methoxyxacinidine (ACMA) upon the addition of 0.5 mM ATP–1 mM MgSO$_4$ (MgATP) as described previously (42, 43). Fluorescence at 410-nm excitation/490-nm emission was monitored for 1 min prior to MgATP addition and for an
additional 40 s after. Proton transport was calculated as the change in fluorescence for the first 15 s following MgATP addition. For both assays, the V-ATPase inhibitor concanamycin A (100 nM) was used to assess V-ATPase-specific activity.

**Vacuolar morphology.** For all vacuolar staining, doxycycline was added to the appropriate treatments upon each medium change. Cells were visualized via DIC and fluorescence microscopy. To simultaneously stain vacuoles with FM4-64 \(N\)-(3-triethylammoniumpropyl)-4-(6-(4-(diethylamino)phenyl)hexatrienyl) pyridinium dibromide\) and CMAC (7-amino-4-chloromethyl coumarin), cells were grown in unbuffered YPD for 24 h in the presence or absence of doxycycline. Then, cells were resuspended in fresh unbuffered YPD with or without doxycycline and grown to

### TABLE 2 Primer sequences used in this study

| Primer     | Primer sequence (5’–3’) | Source          |
|------------|-------------------------|-----------------|
| VMA3-5DRb  | TATAATATATAATTAGCTATGAACTGATTGATTACATATAATGGAAGCCCAACTTGG | This study |
| VMA3-3DRb  | GAAACAGGGTTTCCCCAGTCAGGATTG | This study |
| VMA3-5Det  | AATCCGCTAGCTTCTG | This study |
| VMA3-3Det  | GACTGCGCAAATACTG | This study |
| VMA3-35B   | CCAATCTTTCAGCAGTC | This study |
| VMA3-3Sb   | TGACGTTACCAGTC | This study |
| tetVMA3-5DR | TAAGTTGGGATTGTTGACGAAGATTTGGAAGCTGCAGGAATATGC | This study |
| tetVMA3-3DR | GTTGGAGGAGAGATTTGGAAGCTGCAGGAATATGC | This study |
| tetVMA3-5Det | ACGAGGAATTTTCCAAGGAAATTTGGAG | This study |
| tetVMA3-3Det | GTTGGAGGAGAGATTTGGAAGCTGCAGGAATATGC | This study |
| Vacuolar morphology. For all vacuolar staining, doxycycline was added to the appropriate treatments upon each medium change. Cells were visualized via DIC and fluorescence microscopy. To simultaneously stain vacuoles with FM4-64 \(N\)-(3-triethylammoniumpropyl)-4-(6-(4-(diethylamino)phenyl)hexatrienyl) pyridinium dibromide\) and CMAC (7-amino-4-chloromethyl coumarin), cells were grown in unbuffered YPD for 24 h in the presence or absence of doxycycline. Then, cells were resuspended in fresh unbuffered YPD with or without doxycycline and grown to
early log phase. Cells were resuspended to an OD$_{600}$ of 2 to 4 in unbuffered YPD with 40 μM FM4-64, incubated for 15 min at 30°C, and then resuspended in fresh unbuffered YPD and incubated for 45 min at 30°C. Next, cells were resuspended to an OD$_{600}$ of 0.1 in 10 mM HEPES–5% glucose (pH 7.4). CMAC was added to a concentration of 100 μM, and cells were incubated at room temperature for 15 min and examined via microscopy using Texas Red (FM4-64) and 4,6-diamidino-2-phenylindole (DAPI) (CMAC) filters. To create a three-dimensional (3D) image of the vacuole, FM4-64 staining was completed as previously described (44), except that prior to staining, cells were grown for 24 h in unbuffered YPD with or without doxycycline to ensure complete turnover of Vma3p. A Zeiss ApoTome system was used for capturing Z-stack images. 3D image assembly was completed using AxioVision 4.7 software (Zeiss).

**Secretion and filamentation assays.** Secretion was assessed on solid media: extracellular protease secretion was assayed on unbuffered bovine serum albumin (BSA) plates (45), and lipase secretion was assayed on unbuffered YNB plus 2.5% Tween 80 plates (46). All plates were prepared with and without 20 μg/ml doxycycline. First, cells were grown in YPD with or without doxycycline for 24 h. Then, 3 μL cells were spotted onto plates. BSA plates were incubated at 30°C for 48 h, and Tween 80 plates were incubated at 37°C for 5 days.

Filamentation was assessed on solid and in liquid media. Solid media tested were YPD with 10% fetal calf serum (FCS), medium 199 supplemented with t-glutamine, Spider medium as previously described (47), and RPMI–l-glutamine. All but Spider medium were prepared with 2% (wt/vol) agar. Spider medium was prepared with 1.35% (wt/vol) agar. Filamentation assays were completed (i) with all media buffered to pH 4, and (ii) on standard filamentation media, unbuffered YPD plus FCS, unbuffered M199 (pH 7.5), and unbuffered Spider (pH 7.2) agar, and on RPMI agar buffered to pH 7.0 with 165 mM MOPS. All plates were prepared with and without 20 μg/ml doxycycline. Three microliters cells from overnight cultures were spotted to agar plates, and plates were incubated at 37°C for 5 days. Filamentation in liquid medium was tested in RPMI–l-glutamine (pH 4.0) in the presence or absence of doxycycline. Medium was inoculated with cells from overnight cultures to a starting density of 5 × 10$^5$ cells/ml. Cells were grown at 37°C with shaking at 200 rpm for 2 to 24 h. Cells were visualized via light microscopy at selected time points. We were unable to assess filamentation in liquid fetal calf serum at pH 4.0 due to denaturing of serum proteins at low pH.

We also assessed the effect of chemical inhibition of VMA3 on filamentation in the wild-type strain SC5314. Filamentation was tested in RPMI–l-glutamine buffered to pH 7.0 with 165 mM MOPS. Medium was inoculated with cells from overnight cultures to a starting density of 5 × 10$^5$ cells/ml, and either 10 μM bafilomycin A1 in dimethyl sulfoxide (DMSO) or 5 μM concanamycin A in DMSO was added. The final concentration of DMSO was 3% for bafilomycin A1 and 1% for concanamycin A; therefore, to eliminate the possibility of DMSO effects on filamentation, a 1% DMSO-only control was used. Cells were grown at 37°C with shaking at 200 rpm for 2 to 24 h and were visualized via light microscopy at selected time points.

**Overexpression of positive regulators of filamentation.** A PCR-based transformation method using nourseothricin as a positive selection marker was used to overexpress RIM8, MD3S, UME6, EFG1, or CST20 by inserting the ENO1 promoter directly upstream of each gene (48) in the tetR-VMA3 strain. The genotype of the five resulting strains, listed in Table 1, was tetR-VMA3 + P$_{ENO1}$-RIM8, tetR-VMA3 + P$_{ENO1}$-MD3S, tetR-VMA3 + P$_{ENO1}$-UME6, tetR-VMA3 + P$_{ENO1}$-EFG1, and tetR-VMA3 + P$_{ENO1}$-CST20. Amplons for transformation were generated via PCR using the primers shown in Table 2 and the plasmid pNAT1-ENO1 (from S. Bates, University of Exeter). C. albicans tetR-VMA3 cells were transformed using the lithium acetate method, with a 4-h growth step in YPD added after heat shocking the cells in order to allow integration and translation of the NAT1 gene before exposing the cells to nourseothricin, as described previously (48). Correct integration of the PCR amplons was confirmed by allele-specific PCR using one primer inside the amplon (ENO-SF) (Table 2) and one primer within the open reading frame of the gene targeted for overexpression (Table 2). Integration was further confirmed using an alternative allele-specific PCR with up- and downstream primers flanking the region targeted for insertion (Table 2). Then, tetR-VMA3 + P$_{ENO1}$-RIM8, -MD3S, -UME6, -EFG1, or -CST20 cells were spotted to filamentation-inducing media as described above to determine whether overexpression of these positive regulators of filamentation would rescue the filamentation defect observed in the tetR-VMA3 strain in the presence of doxycycline. The tetR-VMA3 + P$_{ENO1}$-UME6 strain was further investigated; first, tetR-VMA3 and tetR-VMA3 + P$_{ENO1}$-UME6 cells were grown in unbuffered YPD with or without doxycycline for 24 h. Then, cells were washed twice in 1× PBS and resuspended in PBS. Cells in PBS were visualized via light microscopy at 0 h. Finally, cells were seeded in unbuffered YPD plus 10% FCS to a concentration of 5 × 10$^5$ cells per ml and incubated at 37°C, 200 rpm, for 24 h, with visualization via light microscopy at selected time points.

**Biofilm formation.** Biofilm formation was tested using the XTT-reduction assay as previously described (37). Biofilms were formed in RPMI–l-glutamine buffered to pH 4.0 to 8.5. Each treatment was performed in quadruplicate, and each experiment was repeated twice.

**Macrophage killing assays.** The *in vitro* model of macrophage infection was performed as previously described (12). The J774A.1 murine macrophage cell line was purchased from ATCC. Macrophage cells were grown in unbuffered high-glucose Dulbecco’s modified Eagle medium (DMEM) supplemented with 10% FCS at 37°C with 5% CO$_2$ for 72 h. Next, fresh unbuffered DMEM plus 10% FCS was seeded with 2 × 10$^5$ macrophage cells/ml, and 0.75 ml of this solution was used to seed Lab-Tek chambered slides (Nalge-Nunc). The slides were incubated at 37°C, 5% CO$_2$ overnight. Spent medium was removed, and adherent macrophage cells were washed twice with PBS. Overnight cultures of *C. albicans* strains were washed three times in PBS, and *C. albicans* cells were added to unbuffered DMEM plus 10% FCS, with or without doxycycline, to a multiplicity of infection (MOI) of 2. *C. albicans* cells were coincubated with adherent macrophage cells overnight at 37°C with 5% CO$_2$. Then, cells were washed twice with PBS, and macrophage viability was assessed using the Invitrogen Live/Dead viability/cytotoxicity kit, following the manufacturer’s instructions. Live macrophages from 12 separate fields of each chamber were counted, and the results were analyzed for statistical differences using one-way analysis of variance (ANOVA), followed by the Tukey’s multiple-comparison test (GraphPad Prism 5.01). The experiment was performed independently three times, and a representative experiment is presented.

**RESULTS**

**Genetic analysis and disruption of VMA3.** Vma3p is a highly conserved protein (49) largely composed of hydrophobic residues. A BLASTp search of the *Candida* genome database using the *S. cerevisiae* Vma3p sequence as a query revealed a single 161-amino-acid predicted protein with 87.5% identity and 94.4% similarity to *S. cerevisiae* Vma3p. Hydrophobic residues such as isoleucine, phenylalanine, valine, and leucine comprise 39% of *C. albicans* and 40% of *S. cerevisiae* Vma3p. Hydrophobic residues such as isoleucine, phenylalanine, valine, and leucine comprise 39% of *C. albicans* and 40% of *S. cerevisiae* Vma3p. Transmembrane structure prediction of Vma3p using the software program Phobius (50) revealed four transmembrane domains in the open reading frame which are present at identical locations in *C. albicans* and *S. cerevisiae* Vma3p. Given its high degree of structural conservation, we anticipated that genetic deletion of VMA3 in *C. albicans* would mimic findings for *S. cerevisiae* in that it should prevent V$_s$ assembly, eliminate all V-ATPase function, and allow us to assess the contribution of VMA3 and V-ATPase function to *C. albicans* physiology and virulence-related phenotypes.

Therefore, we attempted to construct a vma3Δ null mutant in *C. albicans*. Because vma mutants in *S. cerevisiae* and *C. albicans* grow poorly on alkaline media, we buffered all selective medium.
plates to either pH 4.0 or 5.0 to assist in the selection for positive transformants. One allele of VMA3 was readily deleted in the BWP17 background using the PCR-based “mini-UraBlaster” cassette (31) to generate a VMA3/vma3Δ::dlp200-URA3-dlp200 strain. However, in our hands, we were unable to recover second-allele deletion strains at pH 4.0 to pH 5.0 or on unbuffered medium. Therefore, we constructed a conditional VMA3 mutant, the tetR-VMA3 strain, using a tetracycline-repressible system (34, 35) in which VMA3 expression is suppressed in the presence of doxycycline. We used THE1-C1p10, a strain from the THE1 background in which the URA3 gene has been integrated into the genome, as an additional control (14). Strain construction was confirmed via Southern blot analysis (see Fig. S1 in the supplemental material) and RT-PCR (Fig. 1). RT-PCR analysis indicated that in the presence of doxycycline, the VMA3 transcript was absent in the tetR-VMA3 strain after 24 h (Fig. 1). The VMA3 transcript was present in the tetR-VMA3 strain in the absence of doxycycline and in the THE1-C1p10 strain both with and without doxycycline (Fig. 1). After 6 and 18 h of treatment with doxycycline, the VMA3 transcript was still present in the tetR-VMA3 strain. In all subsequent experiments, all strains were grown for 24 h in the presence or absence of doxycycline prior to the start of the experiment to ensure complete disruption of VMA3.

The tetR-VMA3 strain exhibits the vma phenotype. Cells carrying genetic disruptions of V-ATPase subunits develop the vma phenotype in S. cerevisiae, characterized by pH-dependent lethality (16). To test for pH-dependent growth, tetR-VMA3 cells were spotted on medium adjusted to a broad pH range (pH 4.0 to 8.5). tetR-VMA3 strain growth was comparable to that of the THE1-C1p10 control strain under derepressing conditions (Fig. 2). After addition of doxycycline to repress VMA3 expression, tetR-VMA3 strain growth was decreased at an alkaline pH (pH 7.5 and pH 8.5) (Fig. 2) but not at an acidic pH (pH 4.0 and pH 5.0) (Fig. 2). The vma phenotype was also observed in liquid CSM buffered to pH 4 to 8.5 (data not shown). The tetR-VMA3 cells grew at nearly wild-type levels on nonbuffered YPD and at wild-type levels on CSM under derepressing and repressing conditions (Fig. 2), likely due to acidification of the surrounding medium or of key cellular components. This phenotype has previously been observed in S. cerevisiae vmaΔ mutants (26).

VMA3 is involved in stress responses. The S. cerevisiae vma3Δ mutant grows poorly on media containing high concentrations of calcium or on nonfermentable carbon sources (17). It also exhibits increased cold sensitivity (51) and reduced resistance to a variety of stress conditions (16). We thus plated C. albicans tetR-VMA3 cells after 24 h of growth in unbuffered YPD with and without doxycycline on the following: (i) CSM plus 200 mM CaCl2, both unbuffered and buffered to pH 7.5, (ii) unbuffered YEP plus 2% glycerol as the sole carbon source, or (iii) CSM, pH 4, containing the antifungals caspofungin, fluconazole, and amphotericin B. Repression of C. albicans VMA3 expression significantly reduced tetR-VMA3 growth under most conditions (Fig. 3A). Notably, tetR-VMA3 cells had increased sensitivity to caspofungin (0.025 μg/ml) but did not have increased sensitivity to fluconazole (5 μg/ml) and had modestly increased sensitivity to amphotericin B (0.0125 μg/ml) (Fig. 3A). The dependence of V-ATPase function on ergosterol has been elucidated previously (52). Finally, we tested the ability of the tetR-VMA3 strain to grow at various temperatures after 24 h of growth in unbuffered YPD with and without doxycycline. Like S. cerevisiae vmaΔ mutants, the tetR-VMA3 strain showed enhanced sensitivity to lower temperatures under repressing conditions (Fig. 3B). tetR-VMA3 growth was also tested on medium containing 1.5 M NaCl, 50 μg/ml calcofluor white, or 200 μg/ml Congo red; no difference in growth compared to that of the wild type was observed (data not shown).

VMA3 is necessary for vacuolar acidification. V-ATPase proton transport acidifies the vacuolar lumen, and we anticipated that the tetR-VMA3 vacuolar pH would be altered when VMA3 was repressed. We stained the tetR-VMA3 strain with quinacrine, a basic dye that accumulates inside acidic compartments, such as the vacuole (44), to determine whether vacuolar acidification was defective. After 24 h of growth in unbuffered YPD, the vacuoles of tetR-VMA3 cells accumulated quinacrine comparably to results for the THE1-C1p10 wild-type control (Fig. 4A). After 24 h of
growth in unbuffered YPD with doxycycline, tetR-VMA3 vacuoles did not stain with quinacrine. Fluorometric vacuolar pH measurements using BCECF, a pH-sensitive fluorophore that accumulates in the fungal vacuole (42, 53, 54), validated the results obtained with quinacrine (Fig. 4B). Depletion of Vma3p in the tetR-VMA3 strain led to vacuolar alkalization, as indicated by an increase in the vacuolar pH from 6.17 to 6.77 upon repression of VMA3 expression. Together, these results indicate that Vma3p is necessary for vacuolar acidification in C. albicans.

**VMA3 is required for V-ATPase assembly and activity.** The vacuolar alkalization measured in tetR-VMA3 cells upon depletion of Vma3p suggests that V-ATPase activity was drastically compromised. To directly establish the effect of VMA3 suppression on V1, V0 complex assembly and catalytic activity, we purified vacuolar membrane vesicles from cells grown in unbuffered YPD with and without doxycycline for 24 h by density gradient centrifugation. Western blots using an antibody against the catalytic subunit A of V1 (V1A) did not detect the V1A subunit in vacuolar membrane fractions under repressing conditions (Fig. 5A), indicating that V-ATPase complexes are not assembled. This assembly defect suggests that as in *S. cerevisiae* (23), deletion of the V0c subunit in *C. albicans* prevents assembly of V0c, and consequently, V1 cannot assemble at the membrane.

The V1 domain is the site of ATP hydrolysis in the V-ATPase complex, whereas the V0 domain is the site of proton transport. The lack of intact V-ATPase complex upon depletion of Vma3p is thus predicted to result in a significant reduction in both ATP hydrolysis and proton transport. We measured ATP hydrolysis spectrophotometrically using a coupled enzymatic assay (41), and proton transport was measured fluorometrically using ACMA (42, 43). Both assays were performed in the presence and absence of 100 nM concanamycin A, a specific V-ATPase inhibitor. Under tetR-VMA3 repression, concanamycin A-sensitive ATP hydrolysis decreased by 88% (Fig. 5B) and proton transport decreased by >99% (Fig. 5C). These results are in agreement with the alkalization of the vacuoles in the tetR-VMA3 strain upon addition of doxycycline, as evidenced by both quinacrine and BCECF experiments, and confirm that VMA3 is required for both ATPase hydrolysis and proton transport by the V-ATPase in *C. albicans*.

** Interruption of VMA3 leads to abnormal vacuolar morphology.** Our results indicate that VMA3 is a fundamental V-ATPase component required for V-ATPase activity and maintenance of vacuolar function in *C. albicans*. We next cultured vacuoles with FM4-64 and CMAC to determine if lack of V-ATPase function also altered vacuolar morphology. FM4-64 is a lipophilic dye that is endocytosed and transported to the vacuole, where it stains vacuolar membranes. CMAC is a dye that is thought to passively permeate the cell membrane and accumulate in the vacuolar lumen via the action of glutathione pumps; CMAC accumulation is independent of pH (55). For tetR-VMA3 cells grown in unbuffered medium under repressing conditions, excessive FM4-64 vacuolar membrane staining coincided with CMAC staining of the lumen, indicating intravacuolar accumulation of endocytosed membranes (Fig. 6A). This phenotype was confirmed by performing Z-stack fluorescence microscopy of FM4-64-stained cells (see Movie S1 in the supplemental material). When treated with doxycycline, tetR-VMA3 cells contained a single spherical or obloid vacuole containing multiple membrane compartments. In contrast, *S. cerevisiae* vma3Δ cells have a single enlarged vacuole (56). The vacuolar morphology of the tetR-VMA3 strain after 24 h of growth in unbuffered YPD with and without doxycycline was further assessed using thin-section electron microscopy. The vacuolar morphology of the THE1-Cip10 control strain, both with and without doxycycline, has been studied previously (14). Under repressing conditions, the tetR-VMA3 strain accumulates folds of vacuolar membrane on the interior of the vacuole (Fig. 6B). We observed single, enlarged vacuoles with interior membrane structures in the majority of cells examined (data not shown), indicative of a vacuolar fission defect (56).

**VMA3 contributes to protease and lipase secretion.** The secretion of degradative enzymes is involved in *C. albicans* pathogenesis (5, 8). We assayed *in vitro* secretion of aspartyl proteases and lipases on unbuffered BSA and unbuffered YNB-Tween 80 media, respectively (Fig. 7). *C. albicans* cells, spotted on medium
containing BSA as the sole nitrogen source, secrete secreted aspar- 
ty proteases (SAPs) that digest the BSA, creating a halo of prote-
olysis around the colony (45). Like the THE1-Clp10 control 
strain, the tetR-VMA3 strain exhibited normal proteolytic activity 
under derepressing conditions. The addition of doxycycline com-
pletely inhibited extracellular proteolytic activity of the tetR-
VMA3 strain (Fig. 7). Similarly, wild-type C. albicans cells secrete 
lipases on YNB-Tween 80 medium, creating a halo of precipita-
tion around the colony. Under repressing conditions, the tetR-
VMA3 strain exhibited decreased lipolytic activity on YNB-Tween 80 agar 
(Fig. 7).

VMA3 is required for filamentation. Since repression of 
VMA3 impairs secretion of degradative enzymes involved in 
pathogenesis, we asked whether other virulence-associated traits 
are associated with V-ATPase function in C. albicans. We assessed 
in vitro filamentation by the tetR-VMA3 strain on solid and in 
liquid media buffered to pH 4.0, allowing us to discriminate be-
tween filamentation defects and pH-specific growth defects. Un-
der repressing conditions, the tetR-VMA3 strain did not filament 
on solid media that are either weak inducers or strong inducers of 
filamentation (Fig. 8A). In contrast, the tetR-VMA3 strain grown 
under nonrepressing conditions produced robust hyphal struc-
tures comparable to those of the THE1-Clp10 control strain. We 
also assessed in vitro filamentation of the tetR-VMA3 strain on standard filamentation media: unbuffered YPD plus FCS, unbuf-
tered M199 (pH 7.5), and unbuffered Spider pH 7.2 agar, as well as 
RPMI agar buffered to pH 7.0 with 165 mM MOPS. The tetR-
VMA3 strain was also unable to filament under these conditions 
(data not shown). Under repressing conditions, filamentation of 
the tetR-VMA3 strain was dramatically reduced in liquid RPMI 
(pH 4.0) at 37°C even at 24 h of incubation (Fig. 8B). To further 
validate the importance of V-ATPase in filamentation, we treated 
the wild-type C. albicans strain SC5314 with specific chemical in-
hibitors of Vma3p. Concanamycin A and bafilomycin A1 are po-
tent V-ATPase inhibitors that bind specifically to Vccoli (Vma3p), 
blocking rotation of the hydrophobic c ring and preventing 
proton transport and ATP hydrolysis (57–59). Treatment with 
concanamycin A (Fig. 8C) and bafilomycin A1 (data not shown) 
inhibited filamentation. This lack of filamentation upon pharma-
cological inhibition of V-ATPase further supports the genetic data 
indicating that V-ATPase-mediated proton transport is essential 
for C. albicans filamentation under a variety of in vitro conditions.

The tetR-VMA3 strain exhibits the vma phenotype in bio-
films. The adoption of a biofilm lifestyle by Candida species on 
both biotic and abiotic surfaces has been identified as a major 
factor in their pathogenicity and virulence (60, 61). Therefore, we 
tested biofilm formation in RPMI (pH 4.0 to 8.5). The THE1-
Clp10 control strain formed robust biofilms in RPMI buffered to 
pH 7.5 or 8.5 and biofilms of lesser metabolic activity in RPMI 
buffered to pH 4.0 or 5.0 in either the presence or absence 
of doxycycline (data not shown). At alkaline pH, the tetR-VMA3 
strain exhibited decreased metabolic activity relative to that of the 
control strain in the presence of doxycycline. However, at acidic 
pH, the tetR-VMA3 strain consistently generated metabolic activ-
ity similar to that of controls when coincubated with doxycycline, 
indicating that tetR-VMA3 exhibits the vma growth phenotype in 
both the biofilm and planktonic states.

The tetR-VMA3 strain is attenuated in macrophage killing. 
Phagocytes, such as macrophages and neutrophils, constitute the 
host’s first line of defense against Candida infection. After phago-
cytosis, survival of C. albicans within the host is dependent upon 
induction of phagocyte death (62). We therefore used a murine 
macrophage killing assay to analyze the contribution of VMA3 to 
in vitro virulence (Fig. 9). We first tested planktonic growth in 
unbuffered DMEM over 30 h; there was no significant difference 
in growth between the tetR-VMA3 strain with and without 
doxycycline (data not shown). Next, after 24 h of coincubation in 
unbuffered DMEM plus 10%FCS, the THE1-Clp10 strain effi-
ciently killed the macrophage cell line, as did the tetR-VMA3 
strain without doxycycline. Upon repression of the VMA3 gene, 
the tetR-VMA3 strain displayed significantly attenuated 
macrophage killing.

Overexpression of key positive regulators involved in fila-
mentation regulation does not rescue the tetR-VMA3 filamen-
tation defect. In order to better understand the role of V-ATPase 
activity in filamentation, we sought to ascertain whether V-
ATPase activity is an absolute requirement for filamentation or if the severe filamentation defect in the tetR-VMA3 strain could be overcome by overexpression of positive transcriptional regulators of filamentation. Thus, we generated strains that overexpressed FIG 5 V-ATPase assembly and activity. (A) Western blot using anti-V1A antibody in vacuolar vesicles purified after 24 h of growth in unbuffered YPD ± doxycycline. Eighty micrograms of vacuolar protein was loaded per lane. The V1A subunit (Vma1p) is not detected in vacuolar membrane vesicles from the tetR-VMA3 strain under repressing conditions, indicating that the V-ATPase complex is not properly assembled at the vacuolar membrane. (B) Concanamycin-A-sensitive ATP hydrolysis in vacuolar vesicles purified after 24 h of growth in unbuffered YPD with or without doxycycline. ATPase-specific activity was measured in purified vacuolar vesicles using a spectrophotometric enzyme assay in which ATP hydrolysis is coupled to NADH oxidation. Loss of Vma3p leads to an 88% decrease in concanamycin-A-sensitive ATP hydrolysis. (C) Proton transport in vacuolar vesicles purified after 24 h of growth in unbuffered YPD with or without doxycycline. ATP-dependent proton transport across purified vacuolar membranes was measured via fluorescence quenching of ACMA upon the addition of ATP and MgSO4. Repression of VMA3 leads to >99% reduction in proton transport activity. Asterisks (*) denote statistical significance, P < 0.05, compared to results for all other treatments.

FIG 6 Vacuolar morphology of the tetR-VMA3 strain. (A) FM4-64 and CMAC double staining of cells grown in unbuffered YPD for 24 h with and without doxycycline. FM4-64 (red) stains vacuolar membranes, and CMAC (blue) stains the vacuolar lumen. Under repressing conditions, FM4-64 and CMAC staining reveal the accumulation of membranous structures on the interior of the vacuole in the tetR-VMA3 strain. (B) Thin-section electron microscopy of the tetR-VMA3 strain after 24 h of growth in unbuffered YPD with and without doxycycline. “N” denotes the nucleus, and “V” denotes the vacuole. Under repressing conditions, the tetR-VMA3 strain displays aberrant vacuolar ultrastructure, indicated by enlarged vacuoles with inclusions of vacuolar membrane on the interior of the vacuole.

ATPase activity is an absolute requirement for filamentation or if the severe filamentation defect in the tetR-VMA3 strain could be overcome by overexpression of positive transcriptional regulators of filamentation. Thus, we generated strains that overexpressed FIG 7 Secretion on BSA and Tween 80 agar. Cells were grown for 24 h in unbuffered YPD with and without doxycycline. Then, secreted aspartyl protease (Sap) secretion was assayed on unbuffered YNB-BSA agar plates (72 h at 30°C), and lipase secretion was determined on unbuffered Tween 80 agar plates (120 h at 37°C). When grown under derepressing conditions, the tetR-VMA3 strain secretes aspartyl proteases and lipases at levels comparable to those for THE1-CIp10, evidenced by halos of clearance surrounding the colony. When grown under repressing conditions, the tetR-VMA3 strain exhibits dramatically reduced Sap and lipase secretion.
the positive regulators of filamentation UME6, RIM8, CST20, MDS3, and EFG1 in the tetR-VMA3 background. Next, we assayed filamentation of these strains on solid filamentation agar with and without doxycycline. Under repressing conditions, overexpression of RIM8, MDS3, CST20, and EFG1 did not rescue the filamentation defect in the tetR-VMA3 strain on unbuffered YPD plus 10% FCS agar (Fig. 10A) or on unbuffered M199 (pH 7.5) agar (data not shown).

UME6 is a key regulator of filamentation; overexpression of UME6 results in constitutive filamentation and can rescue filamentation defects caused by mutations in other genes (63–65), including RIM8 and MDS3, which regulate pH-dependent signaling pathways. In nonrepressing conditions, the tetR-VMA3 strain exhibited substantially decreased hyphal growth. (C) Filamentation by C. albicans SC5314 in the presence or absence of 5 μM concanamycin A1, a V-ATPase inhibitor specific to Vma3p. Strains were grown for 24 h in RPMI–l-glutamine buffered to pH 7 at 37°C, 200 rpm. The addition of concanamycin A1 inhibited wild-type filamentation.

FIG 8 Filamentation on hypha-inducing media. (A) Filamentation on YPD plus 10% FCS, M199, Spider, and RPMI agar plates buffered to pH 4. Cells were grown for 24 h in unbuffered YPD with and without doxycycline and then spotted to agar plates and incubated for 5 days at 37°C. When grown under derepressing conditions, the tetR-VMA3 strain produces filamentous structures comparable to those of the THE1-CIp10 control strain. When grown under repressing conditions, the tetR-VMA3 strain exhibits dramatically reduced filamentation on all media tested. (B) Filamentation in liquid culture after 24 h of incubation. Strains were grown in RPMI–l-glutamine with and without doxycycline buffered to pH 4 at 37°C, 200 rpm. When grown under derepressing conditions, the THE1-CIp10 and tetR-VMA3 strains produced hyphae. When grown under repressing conditions, the tetR-VMA3 strain exhibited substantially decreased hyphal growth. (C) Filamentation by C. albicans SC5314 in the presence or absence of 5 μM concanamycin A1, a V-ATPase inhibitor specific to Vma3p. Strains were grown for 24 h in RPMI–l-glutamine buffered to pH 7 at 37°C, 200 rpm. The addition of concanamycin A1 inhibited wild-type filamentation.

FIG 9 In vitro model of macrophage infection. C. albicans cells were grown for 24 h in YPD with and without doxycycline before coincubation with macrophage cells in unbuffered DMEM plus 10% FCS at an MOI of 2. (A) Counts of live macrophage cells from 12 separate fields after 24 h of coincubation with C. albicans strains. The asterisk denotes statistical significance, P < 0.01, compared to all results for other treatments. Each experiment was performed in triplicate; a representative experiment is shown. (B) Live (green) and dead (red) macrophage cells were costained with calcin AM and ethidium bromide homodimer, respectively, and visualized by fluorescence microscopy. Representative images from the 24-h time point are shown.
VMA3+/NAT1-PENO1-UME6 strain formed pseudohyphae rather than true hyphae in rich medium (Fig. 10B). We then studied the morphology of the cells in filamentation-inducing medium (unbuffered YPD plus 10% FCS). Despite constitutive filamentation in the absence of doxycycline, in the presence of doxycycline, overexpression of UME6 did not rescue the tetR-VMA3 filamentation defect; the majority of cells were observed in yeast rather than pseudohyphal or hyphal form (Fig. 10C). Taken together, these data suggest that V-ATPase activity is a fundamental requirement for filamentation.

DISCUSSION

We have shown that C. albicans VMA3 is functionally similar to S. cerevisiae VMA3. Like S. cerevisiae vmaΔ mutants, the C. albicans tetR-VMA3 strain lacks V-ATPase activity and vacuolar acidification upon depletion of Vma3p, and the V$_i$ subcomplex of the V-ATPase fails to assemble at the vacuolar membrane. The pleiotropic effects associated with loss of V-ATPase activity in both species include an inability to grow on neutral to alkaline media, sensitivity to stress conditions such as high calcium and low temperatures, and growth defects on nonfermentable carbon sources. The centralized role of V-ATPase in stress responses is mainly due to the fact that proton pumping energizes the vacuolar membrane and drives secondary transporters involved in sequestering toxins, such as metal ions and metabolic by-products. For example, previous work in S. cerevisiae has suggested that vma mutants are able to utilize the aerobic glycerol metabolism pathway but have a defect in the sequestration of one or more by-products of this metabolism (26). Similarly, we showed that the tetR-VMA3 strain grows poorly on glycerol-containing medium when VMA3 expression is repressed. Of note, sensitivity of C. albicans to high concentrations of calcium and glycerol upon depletion of Vma3p is less severe than that of S. cerevisiae. One possible explanation for this difference is that adaptation to extreme environmental changes in the host has increased the ability of C. albicans to tolerate various stress conditions relative to that of S. cerevisiae.

We are uncertain why we were unable to generate a C. albicans vma3Δ null mutant despite the viability of the S. cerevisiae vma3 null mutant (25–28) and the C. albicans vma7Δ null mutant (15). It is possible that locus-specific factors markedly reduced the efficiency of our genomic integrations or that the null mutants were unable to tolerate the stresses induced by lithium acetate transformation, making successful recovery difficult.

We next studied the contribution of VMA3 to three major C. albicans virulence traits: (i) secretion of degradative enzymes, (ii) filamentation, and (iii) biofilm formation. Both the secretion of degradative enzymes and hyphal development are impaired upon depletion of Vma3p. Loss of Vma3p revealed a pH-dependent phenotype in which Vma3p-depleted cells form reduced biofilms compared to those of controls at alkaline pH but form biofilms of mass similar to that for controls at acidic pH. To our knowledge, this is the first demonstration that the vma phenotype extends to the biofilm as well as the planktonic form of C. albicans. To more directly study the contribution of Vma3p to pathogenesis, we assayed virulence of the tetR-VMA3 strain in an in vitro model of macrophage infection. In vitro macrophage killing of Vma3p-deficient cells is significantly attenuated, since macrophage survival increases 7-fold after repressing VMA3. This defect in macrophage killing is not simply due to a growth defect, since growth in unbuffered liquid DMEM was normal. Furthermore, macrophage phagosomal pH has been studied extensively; in general, macrophage phagosomal pH becomes acidic (i.e., pH range > 5.0) upon engulfment (66, 67). These data suggest that the V-ATPase complex may play an important role in C. albicans pathogenesis and suggest the potential of the V-ATPase as a target for antifungal therapy.

The V-ATPase is important for both vacuolar membrane fission and fusion in S. cerevisiae, and S. cerevisiae vmaΔ mutants exhibit a vacuolar fission defect evidenced by a single enlarged vacuole (56, 68). We have discovered a striking vacuolar fission defect in C. albicans upon depletion of Vma3p, as evidenced by the presence of a single enlarged vacuole with an excessive accumulation of membrane. Deletion of C. albicans VMA7 results in a similar phenotype (15). Vacuolar fission is important for the response to starvation and osmotic stress (56, 69); therefore, defective fission could hinder the ability of the C. albicans tetR-VMA3 strain to respond to environmental stresses. Notably, membrane fission defects are epistatic to membrane fusion defects (56); therefore, the possibility of a hidden membrane fusion defect in V-ATPase mutants cannot be ruled out and is suggested by previous work in both S. cerevisiae (56) and C. albicans (24). Of note, the cold sensitivity observed in vma mutants may be the result of decreased membrane fusion rates; although fusion through the proton pore of the V-ATPase is halted in vma mutants, spontaneous mem-
brane fusion does occur at low rates. Decreased temperature may lead to a lethal decrease in the rate of spontaneous membrane fusion as secretion is more fully inhibited (68). We have shown that loss of Vma3p in *C. albicans* results in cold sensitivity. These data suggest a cryptic vacuolar fusion defect in the tetR-VMA3 strain, and this hypothesis will be tested in future studies. Importantly, defective membrane fusion could partly explain the defect in Sap and lipase secretion observed upon loss of Vma3p, since fusion has been previously linked to secretion (68). Another possible contributor to the accumulation of vacuolar membrane upon VMA3 repression is fusion of autophagic membrane that cannot be properly degraded due to the inactivity of some degradative proteins at an alkaline vacuolar pH.

The yeast-to-hypha transition is a key element of *C. albicans* pathogenesis (70). Environmental inducers of the yeast-to-hypha transition are compatible with host conditions and include alkaline pH, high temperature, the presence of human serum, and nutrient depletion (71). The response to these environmental triggers requires a highly regulated transcriptional network. The cAMP-PKA pathway, which responds to environmental stimuli, including nitrogen starvation, is one of the major regulatory pathways (72). In this pathway, the adenyl cyclase Cya1 utilizes ATP to synthesize cAMP and activate the PKA complex, which then activates filamentation genes, including the transcription factor Efg1, a central regulator of filamentation (73). Maintenance of filamentation after initial hyphal induction is controlled by the transcription factor Ume6 (64, 65, 74). The response to alkaline environmental pH is mediated primarily by the Rim101 signal transduction pathway; this pathway depends on environmental pH sensing, followed by ubiquitinization of Rim8 and processing of the Rim101 transcription factor into an activated form (75–77). A parallel pathway regulated by Mds3 also contributes to filamentation in response to alkaline pH (78, 79). Lastly, the mitogen-activated protein kinase (MAPK) pathway is a lesser contributor to filamentation and includes the protein kinase Cst20, homologous to *S. cerevisiae* Ste20 (47). We overexpressed one positive regulator from each of these five *C. albicans* pathways in the tetR-VMA3 strain and found that overexpression of Rim8, Cst20, MDS3, or EFG1 does not rescue the filamentation defect in tetR-VMA3 cells. Overexpression of UME6 results in partially restored filamentation in YPD (i.e., pseudohyphae) but not under filamentation-inducing conditions, such as YPD plus serum. The role of V-ATPase in filamentation is therefore downstream of these transcriptional regulators or involves independent regulatory mechanisms. These results suggest that V-ATPase activity is a central requirement for filamentation in *C. albicans*.

Loss of VMA3 more severely inhibits filamentation than loss of the V$_{5a}$ isoform *VPH1* (24), and this difference has implications for the mechanism of the V-ATPase contribution to filamentation. Interruption of VMA3 and *VPH1* leads to similar phenotypes, including an $\sim$90% reduction in ATPase-specific activity and an alkalinized vacuole (24). However, loss of VMA3 expression results in a greater reduction in proton transport levels than *vph1Δ* (99.9% versus 88.0%). The similarity in vacuolar pH between the tetR-VMA3 strain and the *vph1Δ* strain suggests that protons transported into the vacuole in the *vph1Δ* strain are quickly utilized by downstream membrane pumps and shuttled out of the vacuole. Importantly, after transport into the vacuole by the V-ATPase, protons are utilized in a range of important cellular processes, including protein and membrane trafficking (11, 19). Further studies are needed to clarify which, if any, of these downstream functions are required for filamentation. Membrane trafficking and delivery of cargo proteins is a promising possibility, since the integration of new membrane at the hyphal tip is a key step in germ tube formation (71). Another possible mechanism to explain the filamentation defect is cytosolic acidification. Germ tube formation requires alkalinization of the cytoplasm, a process regulated via the Pma1p plasma membrane proton efflux pump, which maintains a neutral-to-alkaline cytosol and an acidic external environment (53, 80). Pma1p activity and expression are upregulated during filamentation in *C. albicans* (81). In *S. cerevisiae*, the V-ATPase regulates the trafficking and activity of Pma1p, and V-ATPase mutants display an abnormally acidified cytosol (19, 53, 82, 83). If the V-ATPase-Pma1p axis is maintained in *C. albicans*, cytoplasmic alkalinization should be defective upon loss of VMA3. In accordance with this model, the differential effect on filamentation upon loss of VMA3 versus *VPH1* could be due to the presence of Stv1p-containing V-ATPase complexes in the *vph1Δ* strain, leading to partial retention of Pma1p activity (24); detailed studies to define these mechanisms are under way.

The importance of V-ATPase to such diverse cellular processes in *C. albicans*, including processes important for pathogenesis, underscores the potential of the V-ATPase as a drug target. Notably, the antifungal activity of azoles, which inhibit ergosterol biosynthesis, is thought to be partially due to decreased V-ATPase activity in ergosterol-deficient vacuolar membranes (52). Amphotericin B, whose antifungal activity is largely due to ergosterol binding, may also decrease V-ATPase activity (84). Further, naturally occurring compounds have evolved to inhibit V-ATPase, including two highly specific and potent inhibitors isolated from *Streptomyces* species, balafolinycin A1 and concanamycin A (57, 85). Unfortunately, these compounds are poor therapeutic candidates, since they cannot discriminate between mammalian and fungal V-ATPase (86, 87). However, the existence of the fungus-specific V-ATPase subunit c$_1$, which is encoded by *VMA11* and lacks a mammalian homolog, further supports the potential of the V-ATPase as a drug target (17). Future studies will focus on the contribution of *VMA11* and related V-ATPase components to *C. albicans* cell biology and virulence.

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