Insight into the Antiadhesive Effect of Yeast Wall Protein 1 of *Candida albicans*

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Ywp1 is a prominent glycosylphosphatidylinositol (GPI)-anchored glycoprotein of the cell wall of *Candida albicans*; it is present in the yeast form of this opportunistic fungal pathogen but absent from filamentous forms and chlamydoconidia. Yeast cells that lack Ywp1 are more adhesive and form thicker biofilms, implying an antiadhesive activity for Ywp1, with a possible role in yeast dispersal. The antiadhesive effect of Ywp1 is translatable from yeast to hyphae, as hyphae that are forced to express YWP1 lose adherence in an *in vitro* assay. Deletion of the GPI anchor results in loss of Ywp1 to the surrounding medium and reduction of the antiadhesive effect, implying an importance of time-dependent residency in the cell wall. Anchor-negative versions of Ywp1 possessing or lacking a C-terminal green fluorescent protein (GFP) tag were created in *C. albicans* and harvested from culture supernatants; in addition to serving as quantifiable markers for Ywp1 secretion, they revealed that the cleaved 11-kDa propeptide of Ywp1 remains strongly but noncovalently associated with the Ywp1 core. This association is resistant to highly acidic and basic solutions, 8 M urea, and 1% SDS (below 45°C). Above 50°C, SDS dissociates the isolated complex, but even higher temperatures are required to dissociate the propeptide from native Ywp1 that is anchored in a cell wall. This property has permitted detection, for the first time, of orthologs of Ywp1 in other members of the *Candida* clade. The cleaved propeptide, which carries the sole N-glycan of Ywp1, must participate in the antiadhesive effect of Ywp1.

*Candida albicans* is the most prevalent fungal pathogen of humans, often existing as a commensal in the alimentary tract but being capable of dispersing and causing disease in diverse anatomical locations when immune defenses are compromised. Superficial infections of skin and mucosae are usually not life-threatening, but systemic infections have a high mortality rate (46). Adhesion and biofilm formation are critical components of the persistence and pathogenesis of this fungus in the human population. Biofilms typically start as adherent yeast forms (also termed blastospores or blastoconidia) that develop into filamentous (hyphal and pseudohyphal) forms as they proliferate; the cells become enmeshed in a loose polymeric matrix and develop enhanced drug resistance (3, 8, 13) in a complex and dynamic process that is influenced by a diverse array of regulatory and surface molecules (14). Cells may adhere to tissue surfaces as well as catheters and medical prostheses *in vivo* and to various substrates *in vitro*; the adhesion properties of yeast forms are relevant to both initial adhesion and later dispersal and thus have an important role in the commensalism and pathogenesis of this organism.

Just outside the plasma membrane of each *Candida* cell is an enclosing wall that provides physical protection from hypo-osmosis and other environmental perils and is also the point of contact between the fungus and the host. It is composed of a durable network of β-1,3-glucans cross-linked by β-1,6-glucans and reinforced with chitin (32). Glycoproteins embedded in the plasma membrane or covalently attached to the glucans make up the major protein component of the wall. Rich in mannose-containing glucans, these mannoproteins are prevalent at the outer surface of the wall and are responsible for functions such as recognition, adhesion, and biofilm formation (7). Many of these proteins acquire a permanent or temporary glycosylphosphatidylinositol (GPI) anchor that dictates their wall association (11). This paper focuses on one such protein that has a structure similar to that of the known cell wall adhesins of *C. albicans* but, unexpectedly, has an antiadhesive effect.

Ywp1 (Pga24) is a GPI-anchored glycoprotein of the *C. albicans* cell wall, based on predictions from its sequence (11, 35, 62) and on investigations of the protein itself (10, 23, 52). The 533-amino-acid (aa) polypeptide encoded by YWP1 has the following features: (i) an N-terminal signal peptide of 21 aa that directs the nascent polypeptide into the secretory pathway but is then cleaved and lost; (ii) a tribasic sequence (RRR) followed closely by a dibasic sequence (KR), both of which are cleaved to create an N-terminal propeptide of about 100 aa (at least one of these cleavages is predicted to occur intracellularly, prior to secretion); (iii) a segment of about 360 aa that is especially rich in threonine, with a third of the residues in this segment being either threonine or serine (this segment contains three tandem copies of a 42- to 51-aa motif that is also found in several other wall proteins [23, 39, 54]); and (iv) a hydrophobic C terminus and adjacent signal that result in replacement of the C-terminal 22 aa with a GPI moiety anchored in the plasma membrane. The GPI anchor is subsequently cleaved, and the 378-aa core of Ywp1 becomes attached covalently to the glucan network of the cell wall through a lipid-free remnant of the GPI anchor. A single N-glycan attachment site is present in the Ywp1 polypeptide, near the C terminus of the cleaved propeptide. The Ser/Thr-rich central region of Ywp1 is predicted to be heavily O-glycosylated, consistent with mass spectroscopic analysis of two tryptic peptides from this region (10).
YWPI, the gene encoding Ywp1, is located on chromosome 2 of *C. albicans* (2, 30), about 6 kb downstream of a nonexpressed pseudogene paralog (2, 5, 23, 64). YWPI is highly expressed during exponential-phase growth of yeast forms (23) and is strongly suppressed under conditions that promote filamentation (23, 41, 56). It continues to be regarded as a specific marker for the yeast form of *C. albicans* (18). Deletion of both alleles of YWPI results in strains with normal growth rates, morphologies, and physicochemical sensitivities (23, 43, 47) but increased adhesiveness of the yeast forms (23). Deletion of just one allele of YWPI results in a slight increase in adhesiveness, suggesting a nonlinear dosage effect (23). Ywp1 is thus abundant in the yeast cell wall but not the hyphal cell wall, correlating with its demonstrated effect on yeast cell adhesion but not hyphal cell adhesion. In systemically infected mice, YWPI deletion has little effect on kidney colonization (43) or mouse survival times (23), but numerous other aspects of infection and pathogenicity have yet to be examined.

The molecular mechanism of the antiadhesive effect of Ywp1 is currently unknown. The amino acid sequence of Ywp1 reveals no obvious binding or enzymatic activities that might be important for this effect. Orthologs of YWPI are present in other members of the *Candida* clade (6), both pathogenic and nonpathogenic, but not in the *Saccharomyces* clade or more distant genera; since Ywp1 has so far been identified and studied only in *C. albicans*, there is currently no information from homologs that might provide clues about its activities. YWPI expression patterns have been revealed by dozens of published transcript profiling and mass spectroscopy experiments with *C. albicans* (see File S1 in the supplemental material), but the conditions that have been tested to date represent a small fraction of those that might be encountered by *C. albicans* in *vitro* and *in vivo*; collectively, clear correlations of YWPI expression have not yet emerged for phenotypes other than the proportion of actively growing yeast forms (as opposed to filamentous or filamenting forms) in the populations that have been sampled, and none of the findings are inconsistent with an antiadhesive role for Ywp1. This paper focuses on the Ywp1 protein itself and reveals several functional and structural features that promote a better understanding of how Ywp1 might effect its adhesion-inhibiting activities.

**MATERIALS AND METHODS**

**Strains and media.** *Candida* strains (detailed in Table S1 in the supplemental material) were routinely grown as yeast cells in well-aerated shaking flask cultures at 30°C in a defined minimal medium containing 100 mM glucose and 80 mM ammonium chloride as the carbon and nitrogen sources (medium 13 in reference 23). Supplemental arginine, histidine, and uridine were added as needed, and phosphate was limited in some experiments by reducing its concentration from 5 mM to 0.2 to 0.3 mM. These and other modifications are specified below or in individual figure legends.

Mycelial growth of *C. albicans* for adhesion assays was performed in multiwell polystyrene tissue culture plates containing RPMI medium at 37 to 38°C. Similar results were observed for Thermomax and polyvinyl chloride substrates. A version of RPMI 1640 (Sigma) that contains 11 mM glucose but no bicarbonate or phenol red was supplemented with 25 mM HEPES, 10 to 15 mM NaOH (to raise the pH to 7.5 to 7.7), 0.1 to 0.5 mM uridine, and 0.02% bovine serum albumin (BSA). For growth of mycelia that were utilized for protein analyses, the RPMI medium was supplemented as described above, except with an extra 14 mM glucose (to bring the total to 25 mM) and no BSA.

Mammalian cell lines were grown in Dulbecco’s modified Eagle’s medium containing 6% heat-inactivated fetal bovine serum at 37°C in a humidified atmosphere containing 5% CO₂. Lines included CV-1 (African green monkey kidney), NRK (normal rat kidney), HeLa (human cervical carcinoma), and CHO (Chinese hamster ovary) cells.

**Genetic transformations.** For the ectopic (hyphal) expression of YWPI, the initial coding sequence of one allele of *HWPI* was replaced, by homologous recombination (67), with one of four YWPI-HIS1 cassettes (detailed in Table S2 in the supplemental material). Lithium acetate-polyethylene glycol (PEG) transfections (20) of the PCR amplicons into *C. albicans* strains BWPI7 (YWPI/YWPI) and Cal12 (ywpi1/ywpi1), as well as subsequent selections and genetic analyses, were performed as described previously (23). Selected transformants of strain Cal12, which previously had YWPI coding nucleotides +64 to +815 replaced by ARG4 in one allele and by a recyclable URA3-dpl cassette in the other allele (23), were grown in the presence of 5-fluoroorotic acid (5-FOA) to select progeny that had lost the inserted URA3 cassette; this allowed disruption of the ectopic YWPI gene with a new URA3 cassette designed to replace coding nucleotides +173 to +435 of the ectopic YWPI gene (see Table S2 in the supplemental material).

To create an anchor-negative version of Ywp1, a HIS1 cassette was designed to place a stop codon just upstream of the GPI attachment site codon of YWPI (detailed in Table S2 in the supplemental material). This cassette was transfected into three strains of *C. albicans* that were derivatives of strain BWPI7, with either two wild-type YWPI alleles or codons 22 to 271 of one YWPI allele replaced with either ARG4 or URA3. The STOP/HIS1 cassette was targeted to (and homologously recombined with) the wild-type YWPI alleles as well as the previously disrupted (∆) YWPI alleles, giving transformants (histidine prototrophs) with all combinations of wild-type (WT) and anchor-negative (anchor-minus [AM]) Ywp1: WT/AM, WT/∆, and ∆/AM. Along with the parental strains (WT/WT and WT/∆), this made comparison of several independent transformants possessing various auxotrophies possible, thus strengthening the attribution of phenotypic changes to anchor-negative Ywp1. The engineered genetic changes were confirmed by numerous PCR analyses; for one transformant, the genomic DNA that encompassed the introduced stop codon was amplified and confirmed by sequencing.

To create YWPI-GFP chimera, GFP-HIS1 ampcilons (23) were used to transform *C. albicans* strain BWPI7 and a URA3 derivative of BWPI7. The GFP portion of these constructs was the yeast enhanced green fluorescent protein (EGFP) construct (yEGFP; encodes S65G and S72A enhancements and is optimized for *Candida* codon usage) of Cormack et al. (9) and was targeted to codons 119, 165, 520, and 533 of YWPI (see Table S2 in the supplemental material). Transformants (histidine prototrophs) were screened for correct insertion into YWPI by PCR analyses and by fluorescence detection of secreted GFP; the success rate was 55% (10/18 transformants) for Ywp165-GFP and 69% (9/13 transformants) for Ywp520-GFP; these values are much higher than those typically seen with only 58 to 63 bp of matched flanking sequence (21).

**Protein analysis.** Mannoproteins that were present in culture supernatants or extracted from live cells by use of agents such as SDS and disulfide-reducing agents (dithiothreitol [DTT] and Tris-carboxyethyl phosphine [TCEP]) were concentrated by precipitation with 1 to 2 volumes of ethanol (23); secreted Ywp1–GFP chimeras were harvested from culture supernatants by centrifugal ultrafiltration in devices with nominal cutoffs of 10 to 50 kDa. Unlike yeast cells, mycelia could not be centrifuged into compact pellets, so they were separated from surrounding media and from subsequent washes and extracts by filtration in small, disposable columns with porous frits. β-1,3-Glucanase (Quantzyme ygl) digestions of cell walls previously extracted with DTT, hot SDS, and saturated urea were performed as described previously (23), as were mannoprotein digestions with peptide N-glycanase F (PNGase F; New England BioLabs).

SDS-polyacrylamide gel electrophoresis (SDS-PAGE) utilized the discontinuous Tris-glycine-chloride system of Laemmli (34) or the Tris-Tricine–chloride modification (for Fig. 4 and 7A) of Schagger and von Jagow (51); samples in Fig. 1 and 3 were reduced with DTT prior to electrophoresis. For the Ywp1–GFP fusion proteins, the Laemmli system
was modified as described in File S2 in the supplemental material. Western blotting was performed as described previously (23), using polyvinylidene difluoride (PVDF) filters and chromogenic detection for Fig. 3. For Fig. 1, secondary goat anti-mouse IgG was conjugated to horseradish peroxidase, and detection involved enhanced chemiluminescence capture by X-ray film. In both cases, mouse antisera elicited by DNA vaccination against Ywp1 amino acids 51 to 197 and 105 to 161 were used (23). The brightness and contrast of all images were adjusted uniformly with Adobe Photoshop.

N-terminal sequencing by Edman degradation of a polypeptide electroblotted onto a PVDF filter was performed at the Iowa State University Protein Facility. Mass spectrometry was performed at the Montana State University Mass Spectrometry and Proteomics Facility, as follows. Gel-eluted polypeptides were bound to an Agilent 1100 reverse-phase high-performance liquid chromatography (HPLC) column in 0.1% formic acid, eluted with a gradient of acetonitrile, and fed into a Bruker micrOTOF instrument with electrospray ionization (set at 100 V) and a time-of-flight detector.

RESULTS

YWP1 expression correlates with cellular morphology. C. albicans YWP1 is expressed abundantly in yeast but is downregulated upon filamentation; this has been shown by transcript profiling (31, 40, 41, 56) and by using GFP as a reporter of YWP1 gene expression (23). The latter technique has also revealed that YWP1 is not expressed by suspensor cells or chlamydospores (see Fig. S1 in the supplemental material). Even though the rate of YWP1 downregulation and functional mRNA loss upon filamentation can be high (31, 40, 41, 56), contributions from the diffusion of existing mRNA molecules or from incompletely shut-down YWP1 in early daughter filaments are likely low but have not been ruled out.

Ectopic expression of YWP1. Deletion of both alleles of YWP1 was previously found to increase C. albicans yeast adhesiveness, implying that the presence of Ywp1 in the yeast cell wall results in decreased adhesiveness (23). If this is indeed due to an effect of Ywp1 in the cell wall, then ectopic Ywp1 might similarly decrease the adhesiveness of hyphae, which normally lack this protein. To test this, YWP1 was placed under the control of the endogenous hyphal wall protein 1 gene (HWP1) promoter so that it would be expressed in filamentous forms of C. albicans rather than in yeast forms (61); the adhesiveness of the filamentous forms was then examined.

Strain Ca#12, which previously had both of its YWP1 alleles disrupted (23), was transformed with a series of constructs that targeted cloned versions of YWP1 to the HWP1 locus. The start codon of one allele of HWP1, along with 32 or 42 downstream HWP1 codons, was replaced with the coding sequence of YWP1, resulting in strains that drove expression of YWP1 exclusively with the natural HWP1 promoter. The other HWP1 allele remained unaltered. Six independent strains were created, representing both alleles of YWP1 and different extents of HWP1 replacement; all gave similar results (described below), lessening the possibility that the observed changes were due to unforeseen mutations. For further confirmation and to ensure that the observed effects were not due simply to loss of one of the two HWP1 alleles, the ectopic YWP1 gene was disrupted with URA3 in two of the strains. The data presented here are from one representative set of three strains, denoted HY (hyphal YWP1 strain), HYΔY (hyphal YWP1 strain in which the ectopic YWP1 gene was disrupted with URA3), and HYU (hyphal YWP1 strain in which the URA3 cassette was inserted elsewhere in the genome, leaving the ectopic YWP1 gene intact).

Ectopic hyphal Ywp1 was found to have properties similar to those of yeast Ywp1. (The term “hyphal” is used here to include hyphae and pseudohyphae found in the filamentous growth of mycelia.) Figure 1 shows that ectopic Ywp1 is detectable by Western blotting reveals the presence of Ywp1 in culture supernatants, in disulfide-reduced (TCEP) extracts of intact cells, and in β-1,3-glucanase extracts of cell walls, as indicated. Strains (all derived from C. albicans SC5314) included the wild type (WT; DAY185), a ywp1/ywp1 knockout (ΔΔY; 4L1), and a ywp1/ ywp1 knockout that subsequently had YWP1 placed under the control of the HWP1 promoter (HY); a transfected URA3 cassette then either disrupted the ectopic YWP1 gene in HY (giving HYΔY) or was inserted randomly into the genome (giving HYU). Cells were grown as yeast or hyphae, as indicated. Nonspecifically labeled marker bands provide size references of 46, 58, 80, and 175 kDa. The bar denotes the junction of stacking and resolving gels.

![FIG 1](image)

**FIG 1** Normal (yeast) and ectopic (hyphal) production of Ywp1. Western blotting reveals the presence of Ywp1 in culture supernatants, in disulfide-reduced (TCEP) extracts of intact cells, and in β-1,3-glucanase extracts of cell walls, as indicated. Strains (all derived from C. albicans SC5314) included the wild type (WT; DAY185), a ywp1/ywp1 knockout (ΔΔY; 4L1), and a ywp1/ywp1 knockout that subsequently had YWP1 placed under the control of the HWP1 promoter (HY); a transfected URA3 cassette then either disrupted the ectopic YWP1 gene in HY (giving HYΔY) or was inserted randomly into the genome (giving HYU). Cells were grown as yeast or hyphae, as indicated. Nonspecifically labeled marker bands provide size references of 46, 58, 80, and 175 kDa. The bar denotes the junction of stacking and resolving gels.
pressed ectopic Ywp1 but left the other colonies adherent (Fig. 2C). The dislodged colonies retained their hemispherical shapes. Thus, the antiadhesive effect of Ywp1 appears to be transplantable from yeast to hyphae.

Under identical conditions, two independent strains lacking Als3 (42, 69) had unaltered colony morphology and initial attachment to the substrate, but the stellate colonies showed eventual loss of adhesion, typically several hours sooner than the colonies expressing ectopic hyphal Ywp1 (data not shown). This detachment was not observed in a parental control strain (69), and detachment was inhibited when ALS3 was restored (68). In contrast, strains lacking Hyr1, Cht2, Ece1, Rbt5, Ecm331 (42), or Als1 (16) all maintained adhesion of their stellate colonies (data not shown). Thus, among the few GPI-anchored proteins tested, only the absence of Als3 and the ectopic presence of Ywp1 resulted in loss of hyphal adhesion under these conditions.

Interestingly, none of the strains formed adherent microcolonies unless exogenous protein was present, either in the culture medium or preadsorbed to the plastic. BSA was typically included in the culture medium at 0.02% for these assays, and gelatin and fetal bovine serum could substitute for the BSA. One of four tested brands of gelatin, however, as well as heating of the BSA to 80°C, negated this effect (resulted in loss of adhesion for all strains) (data not shown). Even though the in vitro growth assays that have been devised to document the antiadhesive effect of Ywp1 are not identical to any growth condition in vivo, they are nevertheless robust and convincingly document a role for Ywp1 in reduced cellular adhesion.

Similar antiadhesion effects of ectopic hyphal Ywp1 were observed on a more complex substrate, namely, live monolayers of various mammalian cell lines (Fig. 2D). To cultures of confluent or nearly confluent monolayers of CV-1, NRK, HeLa, and CHO cells were added a small number of wild-type or HY C. albicans yeast cells, which were allowed to grow undisturbed for 1 to 2 days. Poorly adherent microcolonies were dislodged with gentle agitation, and the adherent cells were fixed with ethanol and stained with aqueous crystal violet (shown in Fig. 2D for CV-1 and NRK cell monolayers). As observed for microcolony growth on bare plastic, hyphal expression of YWP1 inhibited adhesion to the mammalian cells. The mammalian cells continued to look healthy throughout the culture period, but there was diminished cell mass directly beneath the fungal microcolonies, visualized most readily by diminished staining where microcolonies had detached from the NRK cell monolayer (Fig. 2D). Similar culture conditions have resulted in minimal disturbance to a variety of primary human cell monolayers (60). The extent of fungal penetration through the mammalian cell monolayers to the plastic substrate was not determined; details of this adhesion are therefore uncertain, but it nonetheless appears to be inhibited by ectopic Ywp1.

Hyphal expression of YWP1 was also engineered into strain BWP17, which has two unaltered YWP1 alleles, resulting in strains that exhibited both normal yeast expression of YWP1 and ectopic hyphal expression of YWP1. Phenotypically, the adhesion properties of the hyphae were the same as those described above, i.e., apparently unaffected by the wild-type YWP1 alleles (data not shown). Since YWP1 and HWP1 have largely complementary and exclusive expression patterns (in yeast and hyphae, respectively), the adhesion properties of the two morphologies appear to be independently assayable. Nevertheless, since mycelial cultures can also generate blastoconidia under various conditions, the strains possessing just hyphal Ywp1 ruled out the possibility that yeast Ywp1 or its effects might have influenced the results of these experiments.

Anchor-negative Ywp1. An anchor-negative version of Ywp1 was created by inserting a stop codon immediately upstream of the “omega” (glycine 511) codon of the GPI attachment site of YWP1, thus eliminating the C-terminal 24 amino acids of Ywp1. Multiple strains were engineered for comparison, differing in their auxo-

FIG 2 Hyphal microcolony growth and adhesion. Sparsely plated yeast cells grown under filament-inducing conditions generated adherent hyphae regardless of YWP1 expression. (A) Strain HY after 8 h of growth. (B) Within 2 days, macroscopic, radially stellate colonies developed, and they remained adherent unless YWP1 was expressed ectopically. (C) Gentle agitation of a 6-well plate, followed by removal of nonadherent colonies, demonstrated the release of strains expressing hyphal YWP1 (strain HYU in this case) and the continued adherence of both wild-type strains and those in which the ectopic YWP1 gene had been disrupted (strain HYΔY in this case). (D) Growth of colonies on monolayers of mammalian cells (monkey CV-1 and rat NRK cells in this case) resulted in similar adhesion properties, with hyphal expression of YWP1 resulting in loss of adherence. The wells were stained with crystal violet after gentle removal of nonadherent colonies, resulting in dark fungal colonies and lighter thinning of the NRK monolayer under dislodged colonies.
No Ywp1 immunoreactivity has been seen in any sample from increased the size and diminished the mobility of this form of Ywp1. quantitatively over time through visualization of the deglycosylated negative Ywp1 in the culture medium were monitored semiquan-

Western blotting with anti-Ywp1 revealed that virtually all of the anchor-negative Ywp1 was secreted into the medium, with faint signal in the lanes for the anchor-negative allele only [lanes 4 and 6]. Marker proteins migrated as indicated (masses in kDa), and the horizontal bar marks the top of the resolving gel.

The appearance and accumulation of wild-type and anchor-negative Ywp1 in the culture medium were monitored semiquantitatively over time through visualization of the deglycosylated propeptide of Ywp1. (Propeptide cleavage is predicted to occur intracellularly, giving a 1:1 stoichiometry for the Ywp1 core and the propeptide.) Figure 4A shows that anchor-negative Ywp1 was prevalent in the culture medium at the end of exponential-phase growth (25 h), whereas wild-type Ywp1 was not. After 42 h, both forms had increased in quantity, but there was a greater fold increase in wild-type Ywp1. (Aside from the case for Ywp1, which can be identified in protein-stained gels only after deglycosylation, no consistent differences in band patterns have been observed between WT/WT and ΔYWP1 strains [23; our unpublished observations].) Numerous analyses such as these have suggested that a substantial fraction (half or more) of the wild-type Ywp1 that is initially anchored to the cell eventually ends up in the medium in 3- to 4-day-old stationary-phase yeast cultures. However, this loss ("shedding") appears to be pH dependent, as shown in Fig. 4B. Using the anchor-negative form of Ywp1 as a quantitative standard, it was apparent that the wild-type form accumulated in the medium when the culture was unbuffered, attaining a stationary-phase pH of about 2, but was nearly absent from the medium when the buffer kept the pH close to 6. Parallel cultures buffered at pH 7 to 8 also showed little or no wild-type Ywp1 in the medium.
poorly mimics wild-type Ywp1 in this antiadhesive effect. Several independent strains with various auxotrophies, making WT and/or AM Ywp1, were compared using this assay. Strains making wild-type Ywp1 showed little yeast biofilm adhesion, while strains making only anchor-negative Ywp1 showed nearly as much adhesion as strains that made no Ywp1 of either sort. Results were independent of requirements for arginine, histidine, and uridine for growth. Enhanced adhesion in the absence of any Ywp1 was found to largely disappear when the growth temperature was raised from 30°C to 37°C, indicating that other (as yet unidentified) antiadhesion mechanisms must become operational in yeast cells at 37°C (23). Figure 5B shows that the same holds true for strains making only anchor-negative Ywp1. Thus, it appears that the antiadhesive effect of Ywp1 in yeast cells is largely dependent upon anchorage of Ywp1 at the cell surface or in the cell wall; mere extracellular presence and accumulation in the medium are not sufficient to markedly reduce yeast biofilm adhesion.

**Ywp1-GFP chimeras.** Four Ywp1-GFP chimeras were created by transfection of *C. albicans* with selectable GFP-HIS1 PCR amplicons targeted to different positions in YWP1. The resulting chimeras included the N-terminal 119, 165, 520, or 533 amino acids of Ywp1 fused to C-terminal GFP. The Ywp119-GFP and Ywp533-GFP chimeras were barely detectable and were not investigated further. The Ywp165-GFP and Ywp520-GFP chimeras were more abundant and were secreted into the yeast culture medium, where they were easily harvested by ultrafiltration. Significantly, these chimeras represented the predominant protein species in these samples. By microscopy, the fluorescence of both chimeras could be visualized in transit through the secretory pathway, but neither showed detectable association with the cell wall after secretion (data not shown).

Ywp165-GFP included 32 amino acids downstream from the dibasic site of Ywp1 (...KRDQIDFIASIENTEGTALEGSTLEVVDYVPGS-GFP; dibasic site is underlined); N-terminal sequence analysis of the gel-purified GFP moiety gave the sequence “DQIDD...,” confirming that cleavage had taken place immediately downstream from the dibasic site. The Ywp520-GFP chimera consisted of a replacement of the C-terminal 13 amino acids of Ywp1 (VLALALIPLAYFI), encompassing the hydrophobic GPI signal anchor, with GFP, giving a splice site sequence of ...FEAGAADASAGAS-GFP, where the underlined G is the “omega” site for GPI attachment. This substitution thus disrupted the normal GPI attachment and anchor cleavage process. The resulting chimera included the repetitive and heavily O-glycosylated core stalk region of Ywp1.

Perhaps the most striking feature of the secreted Ywp1-GFP chimeras was that the cleaved Ywp1 propeptide remained strongly but noncovalently associated with the downstream (core) Ywp1 sequences. In the case of Ywp165-GFP, a segment of only 32 downstream Ywp1 amino acids evidently maintained this association. Remarkably, this association persisted at pH values between 1 and 12, at temperatures up to 65°C, in 8 M urea, and in 1% SDS at temperatures up to 45°C (see Fig. S2 and S3 and File S2 in the supplemental material). The properties of the Ywp1-GFP chimeras are summarized schematically in Fig. 6.

Independent analyses using mass spectrometry, carboxypeptidase B digestion, and electrophoretic mobility shifts all revealed that the Ywp165-GFP preparations always possessed similar amounts of two versions of the propeptide, differing by a C-terminal arginine (the first arginine of the trasic site). Upon pro-
longed storage, limited end nibbling of the propeptide by endogenous proteases also became evident (see Fig. S2 in the supplemental material). The Ywp1 core region of Ywp520-GFP (387 aa) was more prone to cleavage upon storage than that of Ywp165-GFP (32 aa), as it afforded either specific sites or more opportunities for cleavage by endogenous proteases (see Fig. S3 in the supplemental material).

Ywp1 in the walls of wild-type cells. The Ywp1-GFP chimera data suggested that in wild-type C. albicans, the Ywp1 propeptide might be dissociable from the cell wall by use of SDS at 50°C, leaving the downstream Ywp1 core covalently anchored within the wall matrix. To test this, whole wild-type and ywp1/ywp1 knockout yeast cells were washed and sequentially treated with 1% SDS at increasing temperatures. The original culture supernatant, wash, and SDS extracts were precipitated with ethanol, digested with PNGase F, and resolved by SDS-PAGE. Strikingly, there was little evidence of any cytosolic protein being extracted by the SDS; the predominant SDS-extracted protein was the Ywp1 propeptide (as evidenced by its deglycosylated form, which was absent from the knockout strain) (Fig. 7A). Instead of dissociating from the wall-anchored Ywp1 core at 50°C, however, as expected from the Ywp1-GFP chimera results, it dissociated at 70°C. Additional experiments (not shown) revealed that a minor proportion was extractable at 60°C, while most was released at 70°C. Identical results were obtained for five laboratory strains of C. albicans (SC5314, 3153A, 64550, CA-1, and A9), as well as a recent oral isolate (JG1). A strain lacking cell wall phosphomannan (CDH5 [MNN4/H9004]) (27) gave the same result in these experiments as a wild-type strain (SC5314) and a strain with MNN4 restored (CDH13) (data not shown), indicating that simple electrostatic repulsion of the anionic dodecyl sulfate by anionic wall phosphodiester cannot explain the resistance of the Ywp1 propeptide to dissociation. In addition, when hyphae that expressed ectopic YWP1 were tested (namely, strains HY and HYU versus the Ywp1-free strain HYAJ), the Ywp1 propeptide was liberated by SDS at 70°C but not at 50°C (data not shown).

Finally, regardless of their growth pH, when intact SC5314 yeast cells were subsequently treated with HCl at pH 2, no Ywp1 propeptide was extracted (data not shown); this is consistent with the inability of HCl at pH 1 to dissociate the propeptide from isolated Ywp165-GFP (see Fig. S2 in the supplemental material) and confirms that the low pH attained by unbuffered yeast cultures does not liberate the Ywp1 propeptide.

Ywp1 orthologs. Orthologs of YWP1 can be identified in the genome sequences of other species of Candida and in closely related genera (6), and the N-terminal domains of the corresponding translations are highly conserved (see Fig. S4 in the supplemental material). In contrast to the genes, none of the non-C. albicans protein orthologs has yet been shown to exist. Our antisera that specifically bind to the C. albicans Ywp1 polypeptide (23) have not shown cross-reactivity against equivalent samples from non-C. albicans species, but candidate Ywp1 propeptide bands have been observed on gels containing samples similar to those that originally revealed C. albicans Ywp1 (unpublished observations). In light of the results shown in Fig. 7A, we tested non-C. albicans Candida species for the presence of candidate propeptides extractable from whole cells by use of hot SDS. We found that all of the tested species indeed exhibited such a protein; most of it was liberated from the cells by SDS at 60 to 70°C (but not at 50°C), and it appeared as a sharp band with a size similar to that of the C. albicans Ywp1 propeptide only after N-glycan removal by PNGase F. Figure 7B and C show a selection of the data for C. dubliniensis MYA-578, C. lusitaniae 64125, and C. krusei M-103 in comparison to C. albicans SC5314; C. parapsilosis 90018 and C. tropicalis 750.
showed similar patterns, but with lesser amounts of these putative Ywp1 propeptide bands (data not shown). Except for *C. krusei*, whose genome has not been sequenced, the *Candida* Ywp1 propeptides are all predicted to have unglycosylated masses of 10.7 to 11.5 kDa, consistent with the observed bands.

**DISCUSSION**

The *YWP1* gene is expressed primarily, if not exclusively, by the budding yeast form of *C. albicans*. The product of this gene, Ywp1, is a prominent glycoprotein of the yeast cell wall. Deletion of *YWP1* results in greater adhesiveness and biofilm accumulation of the yeast morphotype *in vitro*, implying an antiadhesive effect of Ywp1. Evidence is presented here to show that this effect is attributable to the sustained presence of Ywp1 in the yeast cell wall. The effect is largely but not completely lost if Ywp1 lacks an anchor and passes through the wall without becoming incorporated, suggesting a time-dependent component of its antiadhesive activity. Furthermore, the antiadhesive effect is transplatable, as shown by the ectopic expression of *YWP1* in hyphae. Detachment of such hyphae was observed only after many hours, again suggesting a requirement for a prolonged presence of Ywp1 in the wall. The molecular mechanism of this effect remains unknown but conceivably involves binding or enzymatic modification of other components of the cell wall. A cleaved propeptide that carries the sole N-glycan of Ywp1 must have a role in these effects, as it remains strongly but noncovalently associated with the Ywp1 core polypeptide in the cell wall.

The antiadhesive effect of Ywp1 presumably has an advantageous role in the growth or survival of *C. albicans* in mammalian hosts. Yeast forms and *YWP1* expression are prominent during alimentary commensalism (50, 53, 66), and yeast forms usually coexist with filamentous forms at sites of pathogenic infection (44). Certain routes of dissemination to new growth sites may be favored by the yeast form, due to differences in size, shape, and interactions with host cells (22, 24, 29). The immune system responds differentially to yeast and hyphae (22), but no evidence has yet been presented to show that Ywp1 is a component of that recognition or response. Biofilms that grow on tissue surfaces or implanted medical devices have features that are similar to those of *in vitro* models, which are typically composed of basal yeast layers and overlaid filamentous layers; as such, they may depend on subsequent production of low-adherence yeast to effect dispersal (65). A recent report (49) suggested that phosphate limitation, which increases *YWP1* expression in yeast cells (23), attenuates *C. albicans* SC5314 biofilm formation *in vitro* and *in vivo* and may lead to greater dissemination from the mouse cecum to the kidney, liver, and lung. The yeast cell adhesion that Ywp1 inhibits appears to be a highly complex process, considering that 30 transcription factors have been identified as promoting yeast adhesion to a silicone substrate (14). The postulated role for Ywp1 in yeast dispersal does not rule out other phenotypic traits that have not yet been discerned experimentally, however.

Ywp1 is prominent as a shed, thiol-extractable, SDS-extractable, and unanchored truncated component of the yeast secretome (23; this study) and is prominent in yeast secreted samples and wall samples that have been prepared for mass spectroscopy (26, 37, 57–59). This abundance appears to be an essential feature of the function of Ywp1, as reductions by half or more result in yeast cells that are clearly more adhesive (23). Loss of Ywp1 has no obvious effect on wall resistance properties (23, 47), suggesting that Ywp1 does not have a major structural role in the wall. The extensive shedding of Ywp1 and its accumulation in stationary-
phase yeast cultures (23) appear to be largely a function of the low pH attained by these cultures and may be mediated by acid hydro-lases (17, 33). GPI-anchored Sap9 may have a role in this shedding (52; our unpublished observations). Compared to anchor-negative Ywp1, shed wild-type Ywp1 has increased mobility during SDS-PAGE, suggesting that one or more proteolytic cleavage events simultaneously liberate and shorten it. Shedding of GPI-anchored proteins may also result from cell wall remodeling or from washout during the membrane-to-glucan transfer process (12, 33, 37, 52, 58). Notably, the extractability of Ywp1 with disulfide-reducing agents increases as yeast cultures age (23), suggesting that a subfraction of the shed (as well as unshed) molecules may become disulfide bonded to other polypeptides within the wall and thiol extractable only if the link to β-1,6-glucan (through the GPI remnant) has been severed. Ywp1 has nine cysteines in its 378-aa core, at least one of which is not involved in an intramolecular disulfide bond, and similar cysteines in another GPI-anchored protein have been shown to mediate wall anchorage (4). Formation of such disulfides may require prolonged residence times of the polypeptides in the wall, as the anchor-negative version of Ywp1 and the Ywp520-GFP chimera pass through the wall without obvious trapping by disulfide bond formation. Based on the results presented here, extensive shedding of Ywp1 may be largely an artifact of highly acidic cultures in vitro and is likely less prevalent in more moderate environments in vivo.

Cleaved propeptides often serve as intramolecular chaperones, folding catalysts, signaling molecules, or regulated inactivators. The current data do not reveal the function of the Ywp1 propeptide but do reveal the surprising persistence with which it remains associated with the Ywp1 core. As few as 32 aa of the N terminus of the Ywp1 core are required for this noncovalent association. Whether there are nonadenaturing, nonproteolytic conditions that allow dissociation of the Ywp1 propeptide from the core as part of the normal functioning of Ywp1 remains to be determined. Nevertheless, no physical condition that C. albicans is likely to encounter in the human body, including passage through the stomach, is likely to dissociate the Ywp1 propeptide from the cell wall.

Formation of the 11-kDa propeptide by cleavage of Ywp1 at its tribasic site requires Kex2 and is predicted to occur intracellularly (23). The resulting propeptide exists predominantly as two versions that are present in similar amounts, retaining either one or none of the three arginines of the tribasic site (23; this study). Interestingly, Saccharomyces cerevisiae Kex2 itself has a cleaved propeptide, and C-terminal basic residues on this propeptide are critical for Kex2 activity (36); by extension, the C-terminal arginine on a subpopulation of the Ywp1 propeptides conceivably has an important regulatory function.

In C. albicans, Sap9 and Sap10 are surface GPI-anchored proteins whose proteolytic functions may overlap those of intracellular Kex2 (1, 52). The proteolytic activities of Sap9 and Sap10 span a broad pH range (pH 5 to 8), and although these proteins show a preference for basic and dibasic sites, their cleavage targets include a wide range of sequences (52). When applied to SDS-stripped cell walls, soluble recombinant Sap9 is capable of cleaving the Ywp1 core just 23 aa from its N terminus; Ywp1 that is shed into the culture medium from mutants lacking Sap9 and Sap10 shows no such cleavage upon incubation with recombinant Sap9 or Sap10, presumably because of conformational differences or carbohydrate masking (52). Similarly, no such cleavage appears to occur in intact cells, as no differences have been found for the 11-kDa Ywp1 propeptide or the 378-aa Ywp1 core in comparing wild-type cells with mutants lacking Sap9 and Sap10 (52). The possibility that Sap9 or Sap10 cleaves the Ywp1 core near its C terminus, possibly contributing to some of the observed shedding of Ywp1, has not been ruled out, however; such a process would resemble the liberation of GPI-anchored proteins (such as the adhesin Epa1) from the cell wall of C. glabrata by the GPI-anchored Yps proteases (33). Still other proteases might be responsible for such cleavage, however, as shown for shedding of the extracellular domain of C. albicans Msb2 (63). Cleavage of Ywp1 at its tribasic site (but not its dibasic site) depends on Kex2 (23), but whether Sap9 or Sap10 has a role in cleavage at the dibasic site remains to be determined.

Many C. albicans wall adhesins contain short (4 to 9 aa) segments with high β-aggregation potential; these are thought to form amyloids that are important for cell adhesion (19, 45, 48). These segments are rich in the amino acids Val, Thr, and Ile (average of 77%) and poor in aromatic amino acids, and they have high scores for β-aggregation potential (34 to 100%). Analysis of the Ywp1 sequence reveals five 6- to 8-aa segments with β-aggregation potentials of 94 to 98%, two located in the propeptide and three located in the core region; two of those in the core region are devoid of aromatics and have Val-plus-Thr-plus-Ile contents of 83% and 100%. Reconciling this with the antiadhesive effects of Ywp1 may require that these sequences serve some other function, not be positioned to form amyloids, or, more interestingly, interfere with the adhesive functions of amyloid-forming adhesins.

Cell surface hydrophobicity can promote adhesiveness of C. albicans yeast under certain conditions (25). It is thus conceivable that Ywp1 decreases yeast adhesiveness simply by decreasing cellular hydrophobicity. Various assays have suggested that deletion of YWP1 increases the surface hydrophobicity of yeast cells (unpublished observations), but efforts to reliably quantify this effect have been unsuccessful, in part because wild-type parental strain SC5314 is already relatively hydrophobic (38). Such an antihydrophobic effect might be due to the abundance of Ywp1 if Ywp1 is indeed hydrophilic, but the mechanistic relationship between glycans, hydrophobicity, and adhesion remains complex and enigmatic at this time (38, 55). Alternatively, the effect may be more specific. For example, adhesins of the Als family possess tandem repeats of a 36-aa threonine-rich segment (28) that exhibits hydrophobic patches and mediates binding to similar domains in other proteins as well as to hydrophobic substrates (15); although these repeats have no obvious homology to Ywp1, it is conceivable that Ywp1 might normally block hydrophobic components of the wall and thus result in a more hydrophobic surface when absent.

In conclusion, the existing body of knowledge about Ywp1 suggests that it is made predominantly by rapidly proliferating yeast forms and that it exerts an antiadhesive effect as a resident of the C. albicans cell wall. It will be of interest to determine whether Ywp1 orthologs in other members of the Candida clade share these features. Knowledge of the complex biosynthetic pathway and structural properties of Ywp1 provides a firm foundation for further exploration of the molecular mechanisms of its antiadhesive effect in C. albicans.

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