

Expression and Characterization of the Flocculin Flo11/Muc1, a *Saccharomyces cerevisiae* Mannoprotein with Homotypic Properties of Adhesion[∇]

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The Flo11/Muc1 flocculin has diverse phenotypic effects. *Saccharomyces cerevisiae* cells of strain background Σ 1278b require Flo11p to form pseudohyphae, invade agar, adhere to plastic, and develop biofilms, but they do not flocculate. We show that *S. cerevisiae* var. *diastaticus* strains, on the other hand, exhibit Flo11-dependent flocculation and biofilm formation but do not invade agar or form pseudohyphae. In order to study the nature of the Flo11p proteins produced by these two types of strains, we examined secreted Flo11p, encoded by a plasmid-borne gene, in which the glycosylphosphatidylinositol anchor sequences had been replaced by a histidine tag. A protein of approximately 196 kDa was secreted from both strains, which upon purification and concentration, aggregated into a form with a very high molecular mass. When secreted Flo11p was covalently attached to microscopic beads, it conferred the ability to specifically bind to *S. cerevisiae* var. *diastaticus* cells, which flocculate, but not to Σ 1278b cells, which do not flocculate. This was true for the 196-kDa form as well as the high-molecular-weight form of Flo11p, regardless of the strain source. The coated beads bound to *S. cerevisiae* var. *diastaticus* cells expressing *FLO11* and failed to bind to cells with a deletion of *FLO11*, demonstrating a homotypic adhesive mechanism. Flo11p was shown to be a mannoprotein. Bead-to-cell adhesion was inhibited by mannose, which also inhibits Flo11-dependent flocculation in vivo, further suggesting that this in vitro system is a useful model for the study of fungal adhesion.

The fungal adhesins are a family of cell surface proteins that mediate adherence to environmental substrates or to other cells (7, 45). Adhesins are critically important in the initial steps of fungal pathogenicity, when fungal cells must adhere to host tissue. For the common human pathogens *Candida albicans* and *Candida glabrata*, the involvement of multiple adhesins in the adherence of fungal cells to host tissue has been demonstrated (4, 5, 18, 26, 43).

Among the adhesins is the flocculin family of *Saccharomyces cerevisiae* cell wall proteins that mediate flocculation, which is asexual calcium-dependent cell-cell aggregation. The most recently described member of the yeast flocculin gene family, *FLO11/MUC1* (24, 30), is the only flocculin expressed in the Σ 1278b strain of *S. cerevisiae* (17), and it exhibits a wide variety of phenotypes. Some of these phenotypes are strain specific. Yeast cells of strain background Σ 1278b have been shown to require *FLO11* for invasive growth (23, 30), the development of pseudohyphae (24, 29), and the formation of biofilms on plastic (36), but they do not flocculate. On the other hand, the variant strain *S. cerevisiae* var. *diastaticus*, which is highly flocculent, has been shown to require *FLO11* for flocculation (30).

FLO11 is also required in Σ 1278b strains for the formation of mats with hub and spoke structures on semisolid agar (36). The common laboratory strain background S288C does not express *FLO11* due to a nonsense mutation in the transcriptional activator *FLO8* (28). In some industrial strains, *FLO11* mediates formation of the specialized biofilms called flors that are necessary for the production of sherry wine (19, 48). The common feature of all these phenotypes is adhesion. Commensurate with the many different pathways that regulate its expression, *FLO11* has been shown to have a promoter that is among the largest described for yeast, at over 3 kb (38). Much more is known about gene regulation of *FLO11* (for reviews, see references 11, 25, and 32) than about the structure and function of the protein.

We further investigated the *FLO11*-dependent phenotypes of *S. cerevisiae* var. *diastaticus* and found that it also differs from Σ 1278b in that the haploids do not invade agar and the diploids do not form pseudohyphae. In order to investigate these strain differences in the phenotypes of *FLO11* we expressed and purified the Flo11 proteins from *S. cerevisiae* var. *diastaticus* and from Σ 1278b and examined their properties. An in vitro system was created for studying the adhesive characteristics of the expressed Flo11 by attaching the protein to microscopic beads and testing the adhesive properties of the beads.

Microscopic beads that can be coated covalently with proteins or ligands have been used to simplify several complex biological processes. For example, Gaur and Klotz used microscopic magnetic beads coated with extracellular matrix proteins to isolate a *C. albicans* adhesin gene, *ALS5*, by expression cloning in *S. cerevisiae* (13). Further work using such beads

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TABLE 1. Yeast strains and plasmids used in this study

Yeast strain or plasmid	Strain background	Genotype or description	Reference or source
Strains			
L5487	Σ1278b	<i>MATα leu2::hisG ura3-52</i>	37
L5487 <i>flo11Δ</i>	Σ1278b	<i>MATα leu2::hisG ura3-52 flo11::URA3</i>	29
L5489	Σ1278b	<i>MATα leu2::hisG/leu2::hisG ura3-52/ura3-52</i>	C. Styles and G. Fink
YIY345	<i>S. cerevisiae</i> var. <i>diastaticus</i>	<i>MATα ura3 leu2-3,112 his4 sta⁰</i>	47
YIY345 <i>flo-1</i>	<i>S. cerevisiae</i> var. <i>diastaticus</i>	<i>MATα ura3 leu2-3,112 his4 sta⁰ flo11::URA3</i>	30
YeAD5	<i>S. cerevisiae</i> var. <i>diastaticus</i>	<i>MATα leu2-3,112 ura3-Δ227 arg4 STA1</i>	30
M1800D × YIY319	<i>S. cerevisiae</i> var. <i>diastaticus</i>	<i>MATα leu2-3,112/LEU2 arg4/ARG4 ura3/URA3</i>	6
FY86	S288C	<i>MATα his3Δ200 ura3-52 leu2Δ1</i>	46
Plasmids			
pFLO11		YEplac181-PGK1p-MUC1	8
pFLO11-GPIA		YEplac181-PGK1p-MUC1-GPIA	This work; derived from YEplac181-PGK1p-MUC1 (8)

resulted in characterization of the adhesion properties of Ala1p and Als1p (12, 14, 15, 21, 22, 35). In this study, we have used this approach to study the in vitro properties of purified Flo11 proteins from two different strains.

MATERIALS AND METHODS

Flo11 phenotypic assays. Flocculation was assayed by culturing cells to mid-log phase in yeast extract-peptone-dextrose medium (1), diluting them to equal cell densities, swirling them vigorously with a Vortex mixer, and photographing them immediately and at specified time intervals. Invasion of agar (37) and formation of biofilms on plastic (36) by haploid strains were assayed as described previously. Diploid strains were tested for pseudohyphae development on nitrogen starvation medium as described previously (16).

Construction of pFLO11-GPIA. pFLO11-GPIA was derived from Yeplac181-PGK1p-MUC1 (abbreviated here as pFLO11), which was generously provided by the laboratory of Isak S. Pretorius (8). Yeplac181-PGK1p-MUC1 is a 2 μ m *LEU2* mutant yeast-*Escherichia coli* shuttle vector containing *FLO11* regulated by the constitutive *PGK1* promoter (8). The ultimate source of the *FLO11* gene in this plasmid is cosmid clone ATCC 70895 (American Type Culture Collection). The *FLO11* sequence is identical to that found by the yeast genome sequencing project in strain background S288C (24).

A PstI-Sall fragment of 408 base pairs of the carboxyl-terminal coding region of pFLO11, including the glycosylphosphatidylinositol (GPI) anchor and *PGK1* terminator, was replaced with a synthesized 44-bp insert engineered to contain a sequence of six consecutive histidine residues. The sequences of the oligonucleotides used for the six-His tag were as follows: 5'-GTCCATCACCATCACCA TCACTAAGGCGCGCCTTTTTTTTATG-3' and 5'-TCGACATAAAAAA AAGGCGCGCCTTAGTGATGGTGATGGTGATGGACTGCA-3'.

Growth of yeast strains. The *S. cerevisiae* strains used in this study are listed in Table 1. Strains transformed with either pFLO11 or pFLO11-GPIA were cultured in SC-Leu selective medium (Q-Biogene, Carlsbad, CA) with twice the concentration of amino acid supplements suggested by the manufacturer for growth enhancement (2 \times SC-Leu). Incubation at 30°C was carried out with shaking at 250 rpm until stationary phase was achieved (A_{600} of approximately 4.0, corresponding to approximately 1.5×10^8 cells/ml), when supernatants were collected for analysis of the secreted protein.

Column purification of proteins. Cultures were centrifuged at 5,000 rpm for 10 min, supernatants were collected, and pellets were washed with deflocculation buffer (20 mM sodium citrate-5 mM EDTA) to remove any Flo11p possibly bound to cell surfaces (30). The pH of pooled supernatants and washes was adjusted from a value of ~3.3 to 7.0 with 1 M Tris base. Prepared supernatant was then applied to a Ni-nitrilotriacetic acid-agarose column (Qiagen, Inc., Valencia, CA) of approximately 2.5 cm by 15 cm at 4°C. Wash buffer (phosphate-buffered saline [PBS], pH 7.0, containing 20 mM imidazole) was subsequently applied to the column. Absorbance at 280 nm was monitored until washing was judged to be complete, i.e., A_{280} readings approached zero. Flo11p was then eluted with 250 mM imidazole in PBS, pH 7.0. Two-milliliter fractions were collected, and the purified Flo11p was stored at -80°C.

Purified Flo11p fractions were concentrated using centrifugal filter devices (Microcon YM-30 and Centricon YM-30; Millipore Corp., Billerica, MA). Total protein in the concentrated samples was quantified following a modification of

the Bradford method (Coomassie Plus protein assay reagent; Pierce Biotechnology, Perbio, Rockford, IL).

Western blot analyses. Protein samples were boiled in sodium dodecyl sulfate (SDS) sample buffer that contained 10% SDS and 100 mM dithiothreitol before being separated in discontinuous SDS-polyacrylamide minigels (3.9% separating gel-5% resolving gel; 7.3 \times 8.3 cm). High-molecular-weight protein markers were used (Rainbow; Amersham Life Science, Amersham International, Buckinghamshire, England). Before being blotted, the Coomassie-stained gels were soaked in deionized water for 15 min, equilibrated in 25 mM Tris-192 mM glycine-1% SDS for 1 hour at room temperature, and transferred to 25 mM Tris-192 mM glycine-0.1% SDS for another hour of incubation. Proteins were blotted onto polyvinylidene difluoride (PVDF) membranes (Immobilon Millipore P; Millipore Corp., Billerica, MA), and the membranes were washed as described previously (10). Membranes were incubated with a mouse monoclonal immunoglobulin G primary antibody specific for a consecutive sequence of five histidine residues (penta-His; Qiagen, Inc., Valencia, CA) diluted 1:2,000 in 3% bovine serum albumin in TBS (10 mM Tris-HCl, pH 7.5, 150 mM NaCl), followed by incubation with horseradish peroxidase-conjugated anti-mouse immunoglobulin G secondary antibody diluted 1:2,000 in 10% nonfat dried milk powder in TBS at room temperature.

Antibody binding to His-tagged protein was assessed using a chemiluminescence assay according to the manufacturer's protocol (ECL Western detection reagents; Amersham Pharmacia Biotech, Inc., Piscataway, NJ). Treated blots were exposed to autoradiographic film (HyperfilmECL; Amersham Pharmacia Biotech, Inc., Piscataway, NJ) for intervals of 2 to 5 min.

Assay for terminal mannose. To assess the binding of Flo11p to *Galanthus nivalis* agglutinin (GNA), a plant lectin that specifically interacts with $\alpha(1-3)$ -, $\alpha(1-6)$ -, or $\alpha(1-2)$ -linked terminal mannose residues, protein samples were blotted directly onto a PVDF membrane in a slot blot manifold (Hoefer Scientific Instruments, San Francisco, CA). The membrane was incubated for 30 min in 20 ml of blocking reagent (Roche Diagnostics, Mannheim, Germany), washed two times for 10 min each in 50 ml buffer 1 (TBS-1 mM MgCl₂-1 mM MnCl₂, pH 7.5), and washed one time for 10 min in TBS (0.05 M Tris-HCl-0.15 M NaCl, pH 7.5). Ten microliters of digoxigenin-labeled GNA (Roche Diagnostics, Mannheim, Germany) was added to 10 ml buffer 1, and the blot was incubated in the solution for 1 hour at room temperature. Three 10-min washes with 50 ml of TBS were performed, followed by a 1-h incubation in a solution of 10 μ l anti-digoxigenin antibody linked to alkaline phosphatase (Roche Diagnostics, Mannheim, Germany) added to 10 ml TBS. Washing was again performed three times with 50 ml TBS for 10 min. The blot was immersed in a substrate solution containing 200 μ l 4-nitroblue tetrazolium chloride/5-bromo-4-chloro-3-indolylphosphate (Roche Diagnostics, Mannheim, Germany) and 10 ml buffer 2 (0.1 M Tris-HCl-0.05 M MgCl₂-0.1 M NaCl, pH 9.5) until color development was complete.

Preparation of Flo11-coated beads. Dynal M-450 tosyl-activated beads (Dynal Biotech, Inc., Lake Success, NY) are polystyrene beads that are approximately the size of yeast cells (4.5 μ m in diameter). The surface of each bead is activated through their amino or sulfhydryl groups. Beads at a concentration of 4×10^8 beads/ml were pipetted in 25- μ l aliquots into 0.5-ml tubes. Beads were collected using a strong magnet (Magnetight separation stand; Novagen, Inc., Madison, WI) for 1 to 2 min, and supernatants were discarded. The beads were gently resuspended in 100 μ l of buffer A (0.1 M sodium phosphate, pH 7.4) for 2 min

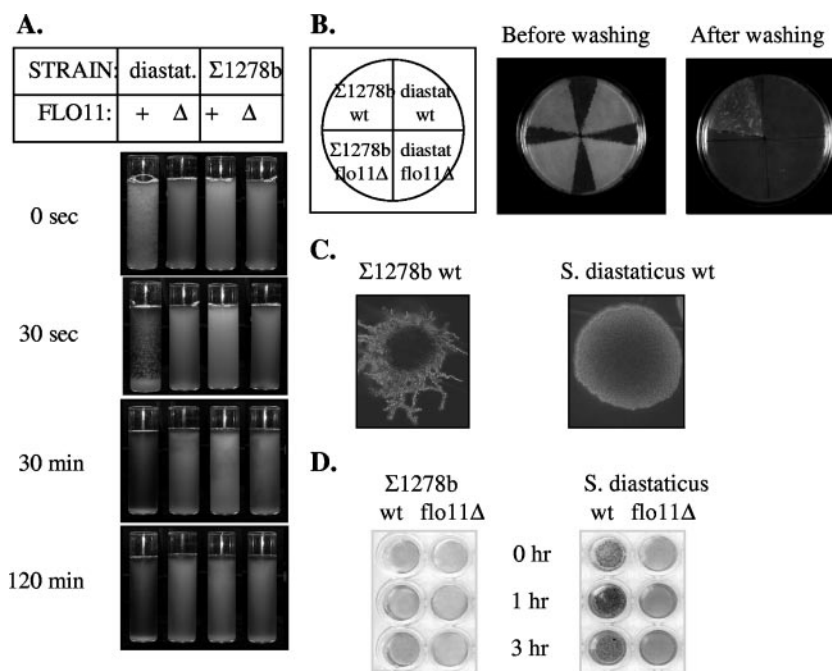


FIG. 1. *FLO11*-dependent phenotypes of yeast cells of *S. cerevisiae* var. *diastaticus* and those of the Σ 1278b strain background. (A) Flocculation assay of *S. cerevisiae* var. *diastaticus* wild-type (+) (strain YIY345) and *FLO11* deletion (Δ) (strain YIY345 *flo11* Δ) strains and of Σ 1278b wild-type (strain L5487) and *FLO11* deletion (strain L5487 *flo11* Δ) strains. Log-phase cultures were adjusted to contain equal cell numbers, vortexed vigorously, and photographed at the indicated time intervals. (B) Agar invasion assay. The strains described above were patched onto yeast extract-peptone-dextrose agar in the arrangement shown on the diagram. They were cultured for 3 days at 30°C, followed by 2 days at room temperature, and photographed before and after being washed with a gentle stream of water. (C) Pseudohyphae development by diploid strains. Cells of each background (Σ 1278b strain L5489 and *S. cerevisiae* var. *diastaticus* strain M1800D \times YIY319) were cultured on nitrogen starvation medium as described previously (16). A representative colony of each strain was photographed. (D) Adherence to polystyrene. Log-phase cultures were suspended in 0.1% glucose, transferred to a 96-well polystyrene plate, incubated at 30°C for the indicated time periods, and stained with crystal violet as previously described (36). The wells were washed with water repeatedly and photographed.

and collected on the magnetic stand for 1 minute. Buffer was pipetted off and discarded. Five micrograms of Flo11p was added to each tube, i.e., to 1×10^7 washed beads, along with buffer A to maintain the initial total volume of 25 μ l. Each tube was vortexed for 2 min and incubated at 30°C for 16 to 24 h with slow, tilt rotation. Beads were also incubated without added Flo11p as a control. Upon completion of incubation, beads were collected for 2 to 3 min in the magnetic stand, and supernatants were removed. Beads were washed twice for 5 min each at 4°C in buffer D, which is PBS, pH 7.4, containing 0.1% fetuin, a glycoprotein which we have shown not to inhibit Flo11-dependent adhesion (data not shown) and which is therefore suitable as a blocking agent. Washing was then performed in buffer E (0.2 M Tris, pH 8.5, with 0.1% fetuin), with incubation at 20°C with slow, tilt rotation for 24 h. Washing was performed in buffer D once for 5 min at 4°C. Flo11p-coated beads and control beads without added Flo11p were stored at a concentration of 4×10^8 beads/ml at 4°C.

Bead adhesion assay. Adhesion of protein-coated beads to cells was assayed by a modification of the method of Gaur et al. (14). Overnight cultures of yeast strains to be combined with the coated beads were diluted to $\sim 2 \times 10^8$ cells/ml, and 500- μ l aliquots were pelleted and washed twice with an equivalent volume of deflocculation buffer. Washed cells were then resuspended in deflocculation buffer containing 20 mM calcium, to promote adhesion (2), combined with 2.5 μ l of coated beads, or 1×10^6 beads, in a total volume of 1 ml. Reaction mixes were vortexed vigorously, and wet mounts were prepared on glass slides for immediate microscopic viewing. Adhesion of yeast cells to beads was assessed by counting beads, utilizing a light microscope with a 40 \times objective (Leica Microsystems, Inc., Allendale, NJ), and scoring them as belonging to one of two categories, i.e., beads bound to yeast cells and beads not bound to yeast cells. Values for each category were calculated as percentages of the total number of beads counted with respect to that category. To assay adhesion of Σ 1278b cells, the cells were first incubated in SC medium plus 0.1% glucose because this strain requires glucose starvation for maximum Flo11-dependent adhesion (36).

To assess the ability of D-mannose to affect the binding of Flo11p-coated beads to yeast cells, the adhesion assay was modified so that 1×10^8 *S. cerevisiae* var.

diastaticus YIY345 cells in 1 ml were placed in a microcentrifuge tube with 2.5 μ l beads coated with column-purified Flo11p derived from *Saccharomyces cerevisiae* var. *diastaticus*. Beads and cells were pelleted and resuspended in 1 ml deflocculation buffer, vortexed, and centrifuged for 5 min at $3,000 \times g$. Buffer was pipetted off, and the procedure was repeated once. D-Mannose at a concentration of 1 M in deflocculation buffer was then added to produce a total volume of 1 ml, and the washed beads and cells were incubated in the sugar for 5 min at 30°C with shaking. Calcium was then added to 20 mM, the tube was vortexed vigorously, and a wet mount was prepared.

RESULTS

Yeast cells from strain backgrounds Σ 1278b and *S. cerevisiae* var. *diastaticus* differ in expression of *FLO11*-dependent phenotypes. It was previously shown that Σ 1278b strains express *FLO11* (29), as do strains of *S. cerevisiae* var. *diastaticus* (30). However, the phenotypic expression of *FLO11* differs in these two strains. *S. cerevisiae* var. *diastaticus* strains flocculate very strongly in a *FLO11*-dependent way, as they begin to settle out of solution immediately after being mixed (Fig. 1A). Σ 1278b strains are virtually nonflocculent in comparison, remaining entirely suspended even 2 h after being mixed (Fig. 1A). Σ 1278b diploid strains, on the other hand, invade agar and form pseudohyphae (16, 17), while *S. cerevisiae* var. *diastaticus* strains do not (Fig. 1B and C). Both *S. cerevisiae* var. *diastaticus* strains and Σ 1278b (36) form biofilms on polystyrene microtiter wells (Fig. 1D). To explore the basis for these

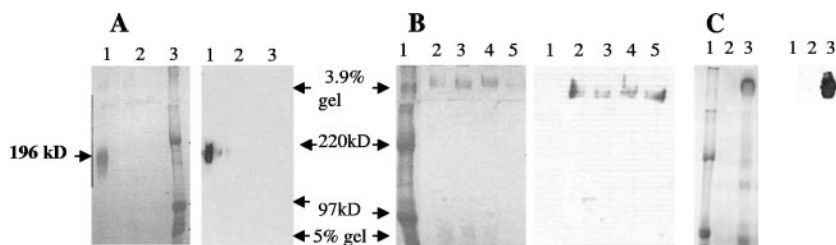


FIG. 2. SDS-polyacrylamide gel electrophoresis of Flo11 proteins secreted from yeast cells of two strain backgrounds. In each case, the Coomassie-stained polyacrylamide discontinuous gel (3.9% stacking gel and 5% resolving gel) is shown on the left, while the corresponding Western blot using anti-six-His antibody is shown on the right. (A) Lane 1, culture supernatant of strain L5487 *flo11Δ* (Σ 1278b background) transformed with pFLO11-GPI Δ ; lane 2, culture supernatant of L5487 *flo11Δ* transformed with pFLO11 (GPI anchor sequences included); lane 3, protein molecular size marker. (B) Purified secreted Flo11p from the Σ 1278b strain aggregated in a form that remained in the stacking gel. Secreted Flo11p from strain L5487 *flo11Δ* transformed with pFLO11-GPI Δ was purified on a nickel column. Lane 1, protein size marker; lane 2, fraction 7; lane 3, fraction 8; lane 4, fractions 9 and 10; lane 5, fractions 13 and 14. (C) Secreted Flo11p from *S. cerevisiae* var. *diastaticus* strain Y1Y345 *flo-1* transformed with pFLO11-GPI Δ produced a large form that remained in the stacking gel. Lane 1, protein size marker; lane 2, culture supernatant of Y1Y345 transformed with pFLO11; lane 3, culture supernatant of strain Y1Y345 transformed with pFLO11-GPI Δ .

strain-specific differences in phenotypic expression, we purified the Flo11 protein from each strain and tested its properties.

A GPI anchor is required for cell surface localization of Flo11. Many fungal adhesins localize to the cell surface through either a cell membrane-anchored or a cell wall-anchored GPI moiety (31). Flo11p has been localized to the yeast cell wall (17, 30). We replaced the GPI anchor with a histidine tag to facilitate secretion of the protein into the medium, from which it could then be purified using nickel columns. When the GPI anchor of *FLO11* was replaced by a six-histidine tag, yeast cells of strain background Σ 1278b transformed with GPI anchorless *FLO11* (pFLO11-GPI Δ) secreted Flo11p (Fig. 2A, lane 1), whereas isogenic yeast strains transformed with wild-type *FLO11* (pFLO11) did not secrete Flo11p (Fig. 2A, lane 2). Using the same plasmids in *S. cerevisiae* var. *diastaticus* also resulted in secretion of the protein (Fig. 2C). This protein was by far the most abundant protein in the concentrated medium, as shown by Coomassie blue staining of the gels in Fig. 2. Therefore, the GPI anchor is critically required for cell surface localization of Flo11p, and its removal results in the accumulation of significant amounts of the protein in the medium.

Secreted Flo11p forms aggregates. The culture supernatant of Σ 1278b strain L5487 *flo11Δ* transformed with pFLO11-GPI Δ contained a protein of approximately 196 kDa (Fig. 2A, lane 1). The identity of the protein as Flo11p was verified by immunodetection with anti-His antibody. However, a much higher-molecular-mass form of Flo11p (>220 kDa) was detected after the same supernatant was purified on a nickel column (Fig. 2B). Fractions collected from the nickel column are shown in Fig. 2B. The higher-molecular-weight form remained in the stacking gel. Total protein quantification showed that purification over Ni-nitrilotriacetic acid resin concentrated Flo11p approximately three times (up to 35 μ g/ml in peak fractions) (data not shown). The culture supernatant of *S. cerevisiae* var. *diastaticus* strain Y1Y345 transformed with pFLO11-GPI Δ most often exhibited the high-molecular-weight form of Flo11p, even before column purification (Fig. 2C, lane 3), although it was also observed in the 196-kDa form (data not shown). Once formed, this high-molecular-weight form of the protein could not be disaggregated by any of the methods tried, which included varying the temperature of incubation of the protein samples, the addition of EDTA at up to

500 mM, and the addition of 50% formamide. Formation of the high-molecular-weight form of Flo11p was also not dependent on the stage of culture growth or the cell density (data not shown). Cell wall-adhesive glycoproteins are known to form aggregates such as this (3, 39, 42).

Flo11p is a mannoprotein. One common characteristic of fungal adhesins is that they contain a large number of residues that are susceptible to N- or O-linked glycosylation (31). Flo11p contains a central domain in which about 50% of the residues are serine and threonine, which are potential O-linked glycosylation sites. Since the predicted molecular mass of Flo11p derived from its amino acid sequence is 137,000 Da, the larger mass observed on the stained gels and immunoblots suggests that the *FLO11* protein product is modified. To test for mannose modification, purified Flo11p samples from YeAD5 (*S. cerevisiae* var. *diastaticus* background) and L5487 *flo11Δ* (Σ 1278b background) transformed with pFLO11-GPI Δ were blotted onto PVDF membranes and treated with digoxigenin-labeled GNA, a plant lectin that specifically binds to α (1-3)-, α (1-6)-, and α (1-2)-linked terminal mannose residues. Figure 3 shows that Flo11p proteins from both yeast strains are mannoproteins possessing terminal mannose residues (Fig. 3, lanes 1 and 2). The positive control for this assay was carboxypeptidase Y (Fig. 3, lane 5), a known mannopro-

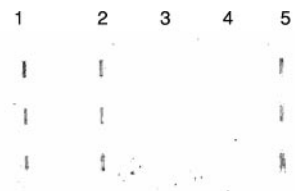


FIG. 3. Flo11 is a mannoprotein. A slot blot is presented, showing purified Flo11 protein applied to a PVDF membrane and treated with a digoxigenin-labeled plant lectin that specifically interacts with terminal mannose residues (GNA). Each lane contains three replicates of a protein sample. Lane 1, nickel column-purified Flo11p from *S. cerevisiae* var. *diastaticus* strain YeAD5 transformed with pFLO11-GPI Δ ; lane 2, column-purified Flo11p from Σ 1278b strain L5487 *flo11Δ* transformed with pFLO11-GPI Δ ; lane 3, fetuin (negative control); lane 4, transferrin (negative control); lane 5, carboxypeptidase Y (positive control).

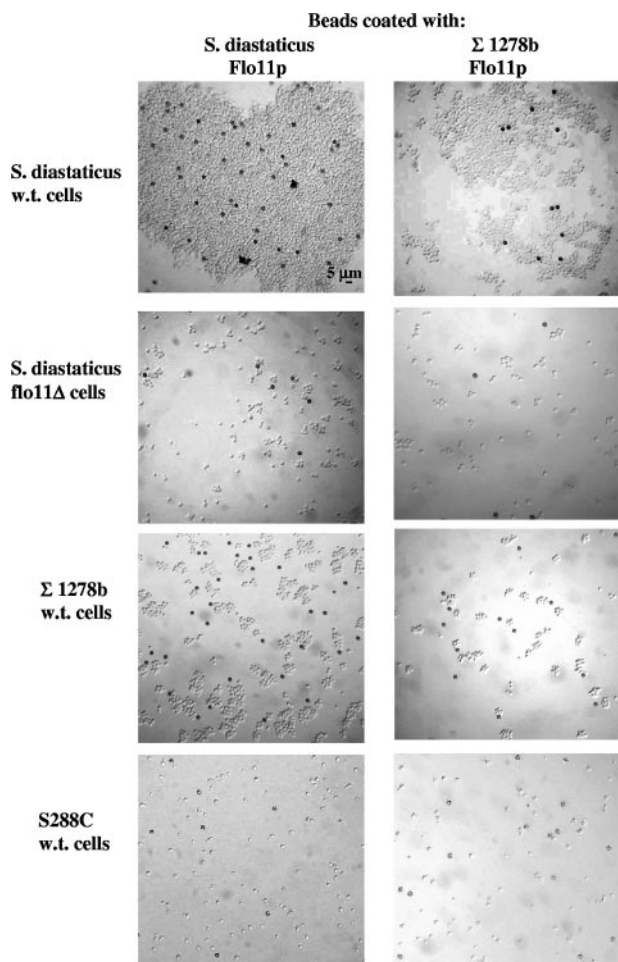


FIG. 4. Bead adhesion assay. Flo11p-coated beads bound specifically to *S. cerevisiae* var. *diastaticus* cells in a *FLO11*-dependent manner. Dynal beads coated with purified secreted Flo11p from either the *S. cerevisiae* var. *diastaticus* or $\Sigma 1278b$ strain background were added to different strains of yeast cells, mixed, and photographed. All photos were taken with a 40 \times objective in the differential interference contrast setting. Beads (approximately 4.5 μm) are visible as dark particles. Cells used were from the following strains: *S. diastaticus* w.t., strain Y1Y345; *S. diastaticus flo11 Δ , strain Y1Y345 *flo-1*; $\Sigma 1278b$ w.t., strain L5487; S288C w.t., strain FY86. A quantitative analysis of this assay is shown in Fig. 5A.*

tein. The glycoproteins fetuin and transferrin, which do not contain mannose, served as negative controls.

Beads coated with Flo11p derived from either $\Sigma 1278b$ or *S. cerevisiae* var. *diastaticus* bind *S. cerevisiae* var. *diastaticus* cells. Yeast strain $\Sigma 1278b$ and *S. cerevisiae* var. *diastaticus* both express the *FLO11* gene (unlike the standard laboratory strain S288C), but only *S. cerevisiae* var. *diastaticus* exhibits Flo11-dependent flocculation. One hypothesis is that the Flo11 proteins produced in these two strains are modified or localized differently, enabling only one form to mediate flocculation. Another possibility is that additional factors in the yeast cell wall are responsible for the different activities of Flo11p in these strains.

In order to examine these hypotheses, an *in vitro* model system was created for the study of Flo11-dependent adhesion by coating plastic beads with Flo11p secreted from the two

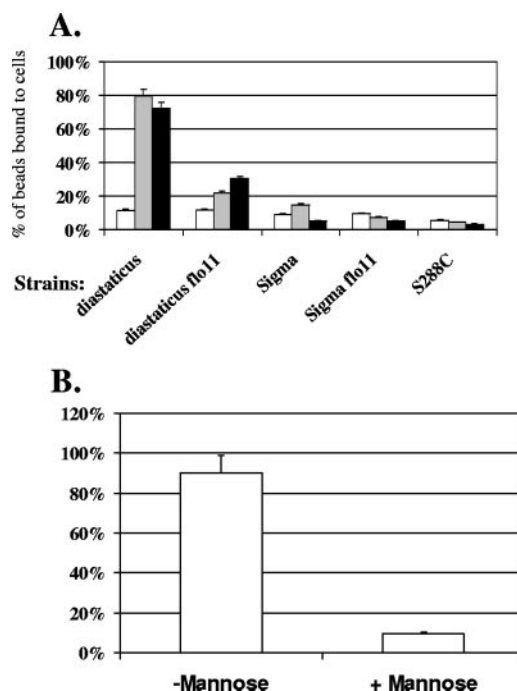


FIG. 5. (A) Quantitation of bead adhesion assay. Each column was based on the mean \pm standard error for three independent experiments, with at least 200 beads counted in each. Uncoated beads (white bars) showed no specific binding to cells; beads coated with purified Flo11p secreted from either *S. cerevisiae* var. *diastaticus* strain Y1Y345 (gray bars) or $\Sigma 1278b$ strain L5487 *flo11 Δ (black bars) showed specific binding to *S. cerevisiae* var. *diastaticus* wild-type cells. (B) Mannose inhibits adhesion of coated beads to yeast cells, just as it inhibits flocculation of these cells. Beads coated with Flo11p derived from *S. cerevisiae* var. *diastaticus* strain Y1Y345 were added to cells of strain Y1Y345, with and without mannose treatment, and the numbers of beads bound and not bound to cells were counted for each case.*

strains of yeast. The Flo11 protein secreted into the medium by *pFLO11*-*GPI* Δ -bearing cells was covalently bound to polystyrene beads of approximately the size of a yeast cell. Dynal M450 tosyl-activated magnetic beads (4.5 μm in diameter) (Dynal Biotech, Inc., Lake Success, NY) are derivatized with *p*-toluenesulfonyl chloride to facilitate the covalent binding of proteins to the surfaces of the beads. The Flo11 protein produced by both $\Sigma 1278b$ and *S. cerevisiae* var. *diastaticus* plasmid-bearing cells was used to coat these beads. Since the secreted Flo11p was encoded by the plasmid, it would have the same primary sequence in both strains, which is identical to the sequence from the S288C strain background, as determined by the yeast genome sequencing project (24). The coated beads were then tested for the ability to adhere to cells of various types.

Uncoated beads did not bind to cells of either strain (see Fig. 5A). However, when they were coated with Flo11p from either yeast strain, the beads adhered to *S. cerevisiae* var. *diastaticus* cells (Fig. 4, top row, and 5A). Therefore, purified Flo11p is sufficient to bind to the cell wall of *S. cerevisiae* var. *diastaticus*. Flo11p produced by the nonflocculent strain $\Sigma 1278b$ was fully capable of mediating bead-to-cell adhesion. Furthermore, the same result was achieved whether the 196-kDa form of Flo11 (from $\Sigma 1278b$) or the large form (>220 kDa; from purified

Σ1278b Flo11 or from *S. cerevisiae* var. *diastaticus*) (data not shown) was used. These data establish Flo11p as an adhesion molecule.

A target of Flo11p adhesion is other Flo11p molecules. We previously established that mannose inhibits Flo11-dependent flocculation (2), leading us to hypothesize that the target of Flo11p adhesion is the mannoprotein layer of the yeast outer cell wall. Since Flo11p itself is a mannoprotein (Fig. 3), we tested the ability of the Flo11-coated beads to adhere to *S. cerevisiae* var. *diastaticus* cells deleted for the *FLO11* gene. Figures 4 (second row) and 5A show that the coated beads failed to adhere to cells lacking Flo11p. Likewise, the coated beads did not bind to cells of strain background S288C (Fig. 4, row 4, and Fig. 5A), which does not express *FLO11*. Therefore, Flo11p itself is an important part of the adhesive target on the cell wall, if not the only target.

Flo11-coated beads did not adhere to yeast cells of strain background Σ1278b, regardless of the source of Flo11p. This result (Fig. 4, row 3) supports the hypothesis that it is differences in the cell wall that are responsible for the different abilities of Σ1278b and *S. cerevisiae* var. *diastaticus* to flocculate in a Flo11-dependent manner. These cell wall differences could be in auxiliary factors required for binding, or they could be differences in the Flo11p receptor molecules anchored in the cell wall.

The adhesion assays were quantitated by counting adherent beads and nonadherent beads in a mixture of cells and beads, using a microscope. The average of three independent assays for each mixture is shown in Fig. 5A. Flo11p from the two strains, i.e., Σ1278b and *S. cerevisiae* var. *diastaticus*, was shown to bind similarly to cells. Among the *in vivo* properties of Flo11-expressing cells is the ability to adhere to agar (24, 29) and to polystyrene (36). These Flo11-coated beads, however, were not observed to bind to agar or plastic (data not shown). It may be that the proper experimental conditions for such binding have not been found, that the specific gravity of the magnetic beads is too high to permit adhesion, or that other factors are involved in binding to these substrates.

Mannose inhibits Flo11-dependent adhesion. Since mannose inhibits Flo11-dependent flocculation *in vivo* (2), an *in vitro* adhesion assay was performed to directly assess the ability of D-mannose to affect the binding of Flo11-coated beads to yeast cells. When beads coated with Flo11p derived from *S. cerevisiae* var. *diastaticus* strain YeAD5 were combined with *S. cerevisiae* var. *diastaticus* Y1Y345 cells and then incubated in 1 M D-mannose, no binding of beads to cells was detected (Fig. 5B). Glucose at the same concentration did not inhibit binding (data not shown). Flo11p is a member of the class of Flo1-type flocculins, whose defining characteristic is inhibition by mannose but not by glucose, maltose, or sucrose (2). Therefore, this *in vitro* system utilizing coated beads to test Flo11p adhesion to cells reflects the *in vivo* properties of mannose inhibition. This finding provides further evidence that mannose is a component to which Flo11p binds on the cell and that Flo11p functions in a lectin-like manner in *S. cerevisiae* var. *diastaticus*.

DISCUSSION

This study directly demonstrates that Flo11p is an adhesion molecule. Yeast cells of three different strain backgrounds

were used in these studies. S288C has been used as a standard laboratory strain of yeast for many years; it does not express *FLO11* and consequently does not form pseudohyphae or exhibit other *FLO11*-dependent properties (28, 38). The first yeast strain shown to form pseudohyphae and to invade a substrate was Σ1278b (16), which exhibits constitutive suppression of the stress response due to high cyclic AMP levels (40). Cells with the Σ1278b background exhibit many *FLO11*-dependent characteristics besides pseudohyphae, including agar invasion (29, 37), biofilm formation, and adhesion to plastic (36), but they do not flocculate. *Saccharomyces cerevisiae* var. *diastaticus*, on the other hand, flocculates very strongly, but when *FLO11* is deleted flocculation is abolished (30). However, this work shows that in spite of its expression of Flo11p, *S. cerevisiae* var. *diastaticus* does not invade agar or form pseudohyphae. These two strains thus represent a naturally occurring experiment to determine the factors that govern Flo11-dependent adhesion. In order to investigate these factors, Flo11 proteins were purified from these two strains and their properties were tested.

When the GPI anchor sequences were removed from Flo11p, the protein accumulated in the extracellular medium. Two forms of Flo11p were observed, with one form of 196 kDa and another, very large form, which did not enter the separating gel. Purification of secreted Flo11p on nickel columns always resulted in conversion of the 196-kDa form to the larger form. We suggest that the large form represents an aggregate of Flo11 protein. Such aggregates have frequently been seen in SDS gels containing membrane and cell wall proteins (3, 9, 33, 39, 42). At least three groups have purified mannose-specific lectins from yeast surfaces, and similar SDS-resistant aggregation properties were displayed by all of them (3, 39, 42). Bony et al. (3) extracted Flo1p from cell walls by using hot SDS-β-mercaptoethanol and observed that the protein migrated in SDS gels as a very high-molecular-mass protein in a highly heterogeneous fashion. The fastest-migrating forms of Flo1p exhibited molecular masses of about 200 kDa, while the largest forms remained at the top and did not enter the resolving gel (3), just as we found for Flo11p. When the GPI anchor sequences of Flo1p were removed, the protein was secreted into the medium in a form that produced a fuzzy, heterogeneous, high-molecular-mass band in gels (3). Treatment with endo-β-N-acetylglucosaminidase H resulted in more protein entering the gel, suggesting that Flo1p is N glycosylated. However, the enzyme-treated Flo1p remained larger than predicted from the amino acid sequence, suggesting O glycosylation as well (3).

Consistent with the model of flocculins as glycoproteins, a homolog of Flo1p has been demonstrated to have a sugar content of 63% (42). This protein also exhibited properties of aggregation; gel filtration studies revealed an active aggregate with an apparent molecular mass of >700 kDa. The present study establishes Flo11p as a mannoprotein.

Flo11p functioned as an adhesin *in vitro* when it was attached to beads. The coated beads bound only to cells of the strain background that exhibits Flo11-dependent flocculation, namely, *S. cerevisiae* var. *diastaticus*, not to Σ1278b strains, which do not flocculate. This binding was not due to nonspecific trapping of the beads in the large flocs of *S. cerevisiae* var. *diastaticus*, since uncoated beads did not bind. Both strains produced Flo11p that is mannosylated, and both produced

Flo11p that adheres to *S. cerevisiae* var. *diastaticus* cells equally well. In vitro model systems such as this one should prove to be very useful for further understanding fungal adhesion.

The work presented here is the first to identify an adhesive target of Flo11p. Since the outer layer of the yeast cell wall consists largely of mannoproteins (20, 27), it seemed likely that Flo11p would bind to the side chains of the cell wall mannoproteins. It has been thought that the specificity of fungal lectins may be quite broad (41). Somewhat surprisingly, we found that an essential component (and perhaps the only component) of the adhesive target of Flo11p is, in fact, other Flo11p molecules. Flo11p receptors on the cell wall are required for adhesion of Flo11-coated beads; it remains to be seen whether they are sufficient. Strain-specific differences in these Flo11p receptors in the cell wall could possibly explain the differential adhesion of *S. cerevisiae* var. *diastaticus* versus Σ 1278b cells to the coated beads. Such receptor differences could be quantitative or qualitative.

Homotypic adhesion has been demonstrated for the ALS adhesins of *Candida albicans* (21, 22). Yeast cell aggregation mediated by cloned ALS proteins has properties characteristic of amyloid protein aggregation (35), and alterations in these properties could, in theory, produce phenotypic variation. Another possible mechanism of fungal adhesion is suggested by the observation that in Als5p, the threonine-rich repeat domain can mediate homotypic adhesion to repeats on the surfaces of apposing cells (34). Such tandem repeats are a hallmark of yeast cell wall proteins and are also found in Flo11p. In the yeast flocculin Flo1p, variation in the length and number of tandem repeats has been shown to occur between strains and to correlate with functional variation in adhesive phenotypes (44). It is possible that mechanisms such as these underlie the strain-specific variation we see in *FLO11*-dependent phenotypes.

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