Tight Control of Trehalose Content Is Required for Efficient Heat-induced Cell Elongation in Candida albicans*

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Background: Heat-induced morphogenetic switch is Hsp90- and PKA-dependent.
Results: The G protein-coupled receptor Gpr1, upstream of PKA, regulates trehalose levels in glucose-grown cells.
Conclusion: The link between the PKA pathway and Hsp90-mediated regulation of heat-induced morphogenesis is trehalose.
Significance: Tight control of trehalose levels is required for morphogenesis in Candida albicans.

The ability to form hyphae in the human pathogenic fungus Candida albicans is a prerequisite for virulence. It contributes to tissue infection, biofilm formation, as well as escape from phagocytes. Cell elongation triggered by human body temperature involves the essential heat shock protein Hsp90, which negatively governs a filamentation program dependent upon the Ras-protein kinase A (PKA) pathway. Tight regulation of Hsp90 function is required to ensure fast appropriate response and maintenance of a wide range of regulatory and signaling proteins. Client protein activation by Hsp90 relies on a conformational change of the chaperone, whose ATPase activity is competitively inhibited by geldanamycin. We demonstrate a novel regulatory mechanism of heat- and Hsp90-dependent induced morphogenesis, whereby the nonreducing disaccharide trehalose acts as a negative regulator of Hsp90 release. By means of a mutant strain deleted for Gpr1, the G protein-coupled receptor upstream of PKA, we demonstrate that elevated trehalose content in that strain, resulting from misregulation of enzymatic activities involved in trehalose metabolism, disrupts the filamentation program in response to heat. Addition of geldanamycin does not result in hyphal extensions at 30 °C in the gpr1Δ/gpr1Δ mutant as it does in wild type cells. In addition, validamycin, a specific inhibitor of trehalase, the trehalose-degrading enzyme, inhibits cell elongation in response to heat and geldanamycin. These results place Gpr1 as a regulator of trehalose metabolism in C. albicans and illustrate that trehalose modulates Hsp90-dependent activation of client proteins and signaling pathways leading to filamentation in the human fungal pathogen.

Candida albicans is an opportunistic human fungal pathogen that can cause superficial and severe invasive infections. Virulence of the fungus is in part dependent on the morphogenic plasticity of C. albicans, as strains that are trapped in a particular growth form are avirulent (1–4). The complex trait of filamentous regulation is under the control of multiple signaling pathways, including the Ras-protein kinase A (PKA) pathway (5). This pathway regulates several biological processes in response to nutrient availability, such as stress resistance, growth, cell differentiation, and trehalose metabolism (6–8). In Saccharomyces cerevisiae, PKA activity is low, and trehalose levels increase upon nutrient starvation, whereas activation of the pathway by glucose or sucrose, via the G protein-coupled receptor Gpr1, results in low trehalose levels (9, 10). In C. albicans, the nature of the ligand of the homologous receptor Gpr1 remains controversial as the receptor is not required for sugar-induced cAMP signaling (7). Trehalose is a nonreducing disaccharide that can be found in bacteria, fungi, plants, and insects. It is synthesized by two consecutive enzymatic reactions. The enzyme trehalose-6-phosphate synthase (TPS),3 encoded by TPS1, produces trehalose-6-phosphate, which is then converted into trehalose by the trehalose-6-phosphate phosphatase (TPP) enzyme encoded by TPS2 (11, 12). Trehalase biosynthesis plays a role in the virulence of different human and plant pathogenic fungi, as typically, deletion of the Tps1 and Tps2 orthologs leads to absence of trehalase formation, reduction or absence of virulence, and sensitivity to stress (13–21). However, molecular links among trehalase, the filamentous programs, and virulence have never been identified.

Human body temperatures induce filamentous growth in C. albicans, and this process is evoked by the release of repression by the essential molecular chaperone Hsp90 on its target proteins (22). The effect of Hsp90 on filamentation was originally suggested to be mediated via the Ras-PKA pathway, as strains deleted for positive regulators of the pathway are unable to form filaments upon compromise of Hsp90 function by its inhibitor geldanamycin, whereas strains deleted for negative regulators easily show the morphogenetic response (22). However, a clear point of interaction between the Ras-PKA pathway and Hsp90 could not be identified (23, 24).

In the current study, we provide evidence that supports a model of regulation of heat-induced filamentation by the

3 The abbreviations used are: TPS, trehalose-6-phosphate synthase; TPP, trehalose-6-phosphate phosphatase; Tricine, N-[2-hydroxy-1,1-bis(hydroxymethyl)ethyl]glycine.
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chemical chaperone trehalose. In this model, the Ras-PKA pathway participates in the release of function of Hsp90 by regulating trehalose content, hence providing clues on the role of the signaling pathway in the Hsp90-dependent filamentation program. Heat-induced filamentation is severely reduced by the absence of Gpr1. We show that this defect is caused by increased intracellular trehalose content. In addition, validamycin, a trehalose analog, promotes trehalose accumulation and blocks filamentation in response to heat and geldanamycin. Our results clearly establish trehalose as a negative regulator of the heat-induced cell elongation program.

EXPERIMENTAL PROCEDURES

Strains and Growth Conditions—All C. albicans and S. cerevisiae strains used in this study are listed in Table 1. Under continuous shaking, C. albicans strains were grown at 37 °C, S. cerevisiae strains were grown at 30 °C. Strains were grown on rich medium (YP: 1% yeast extract, 2% bactopeptone) supplemented with 2% glucose (YPD) or 3% glycerol (YPG), or on synthetic complete medium (SC: 0.17% yeast nitrogen base without amino acids and without ammonium, supplemented with synthetic drop-out amino acid and nucleotide mixture as required, 0.5% ammonium sulfate) and supplemented with 2% glucose (SCD). Solid media contained 2% (w/v) Difco-agar in addition. When applicable, geldanamycin or validamycin was added to a concentration of 10 μM, unless stated otherwise.

Construction of Plasmids and Strains—The C. albicans GPR1 gene was amplified with primers pCaGPR1tF and pCaGPR1tR (Table 1). The resulting pCaGPR1t PCR fragment was co-transformed with a BglII/MluI linearized YEplac112-pScGPR1t fragment to S. cerevisiae. pCaGPR1t and pScGPR1t were cloned in pYX012KanMX for genomic integration by digestion with EcoRI/SphI and SacI/SphI, respectively, and ligation. pYX012KanMX for genomic integration was obtained with a BglII/MluI linearized YEplac112-p

Intracellular levels of glucose 6-phosphate and ATP were determined as described previously (26). 

| TABLE 1 | C. albicans and S. cerevisiae strains and oligonucleotides used in this study |
| Genotype | Source or reference |
|------------------|-----------------------|
| S. cerevisiae strains |          |
| JS4 | H9262::URA3::hisG-URA3::hisG-ura3 This study |
| JS16 | H9262::URA3::hisG-URA3::hisG-gpr1 This study |
| JS4 | H9004::hisG/gpr1 |
| JS16 | H11001::hisG-URA3::hisG \( \Delta \) gpr1 |
| LK41 | W303-1A |
| LR2 | H9004::hisG-URA3::hisG-URA3/ura3 |
| CR216 | H9004::hisG-URA3::hisG-URA3/ura3 |
| C. albicans strains |          |
| HT10 | H9261::ura3/ura3 |
| BSC7 | H9004::FRT/ura3 |
| BSI3 | H9004::FRT |
| CAH1+URA4 | H9261::ura3/ura3 |
| GPR1-URA3 | H9004::FRT/ura3 |
| MTR1-URA3 | H9004::FRT/ura3 |
| MTR2-URA3 | H9004::FRT/ura3 |
| MTR3-URA3 | H9004::FRT/ura3 |
| MTR4-URA3 | H9004::FRT/ura3 |
| MTR5-URA3 | H9004::FRT/ura3 |
| MTR6-URA3 | H9004::FRT/ura3 |
| MTR7-URA3 | H9004::FRT/ura3 |
| MTR8-URA3 | H9004::FRT/ura3 |

Biochemical Determinations—Intracellular levels of cAMP and trehalose were determined as described previously (25). Intracellular levels of glucose 6-phosphate and ATP were determined as described previously (26).
For determination of extracellular trehalose content, the Waters Breeze HPLC (Waters Corporation) was used. Samples were taken over time and cells were centrifuged and discarded. Extracellular medium was used in the analysis. Samples were analyzed at a flow rate of 1 ml/min, using 5 mM H2SO4 as eluant. Results were processed with the Breeze software. Trehalose content was determined as mMtrehalose.

**Enzymatic Determinations**—Trehalase activity was determined as described previously (27). Hexokinase activity was determined as described previously (28).

For TPS and TPP activity determination, cells were harvested at indicated time points and washed with and extracted in 50 mM imidazole (pH 6.3) supplemented with 1 mM EDTA, 2 mM MgCl2, and protease inhibitor and desalted on Sephadex columns. For TPS activity determinations, samples were added to an assay mixture solution, containing (at final concentration) 40 mM Tricine (pH 7), 10 mM MgCl2, 4 mM uridine diphospho-glucose, and 8 mM glucose 6-phosphate. In the control samples, glucose 6-phosphate was omitted. Formed trehalose 6-phosphate was determined as the addition of phosphoenolpyruvate, NADH, and lactate dehydrogenase and measured as a difference in A340 nm before and after addition of pyruvate kinase. The amount of protein in the samples was determined as described by Lowry et al. (29). TPS activity was expressed as nanokatals of TPS (g of protein)−1. For TPP activity determinations, samples were added to an assay mixture solution, containing (at final concentration) 40 mM Tricine (pH 7), 10 mM MgCl2, and 2.5 mM trehalose 6-phosphate. In the control samples, trehalose 6-phosphate was omitted. Formed trehalose was determined by detection of glucose formed after hydrolysis by trehalase. The amount of protein in the samples was determined as described by Lowry et al. (29). TPP activity was expressed as nanokatals of TPP (g of protein)−1.

**Microscopy**—Cells were grown in synthetic complete medium containing glucose at 37 °C. Experiments with geldanamycin were conducted as described previously (22). Cells were observed with a Zeiss Axioscope 2 fluorescence microscope, and images were captured with an Axioscam HRm camera by using Axiovision 3.0 software (Carl Zeiss, Thornwood, NY).

**Gene Expression Analysis**—Total RNA was extracted using the TRIzol reagent method. Complementary DNA was prepared from DNase-treated RNA samples with a Reverse Transcription kit (Promega). Quantitative PCR was performed using a StepOnePlus real-time PCR system (Applied Biosystems). Reactions were prepared with the Kapa SYBR Fast kit (KapaBiosystems) according to the manufacturer’s instructions. The fold regulation of the target gene was calculated using the comparative Ct method, using TEF1 Ct to normalize the data.

**RESULTS**

**Gpr1 Is Required for Lowering Trehalose Content during Active Growth in C. albicans**—Previous findings have shown that absence of GPR1 in C. albicans leads to increased trehalose content as well as activity of the Tps1 enzyme, especially in heat shock conditions (30). We investigated the role of Gpr1 on trehalose content in C. albicans in conditions of 37 °C, the physiological temperature that is not experienced as a stress in the pathogenic fungus. Deletion of GPR1 resulted in increased levels of trehalose in exponentially growing cells (Fig. 1A), a growth phase during which trehalose does not normally accumulate. Significantly higher amounts of trehalose were measured in the deleted strain compared with wild type and GPR1-reintegrand strains. This peculiar accumulation occurred in glucose-containing growth conditions, however not in glycerol-containing media (data not shown). Defective export of tre-
Trehalose in the surrounding medium was ruled out by HPLC analysis, which showed equal amounts of trehalose accumulating in the medium for all strains (data not shown). Higher trehalose content in the \textit{gpr1}^{−}/\textit{H9004}^{−}/\textit{gpr1}^{−}/\textit{H9004}^{−} strain did not impair the carbohydrate mobilization upon addition of glucose to sugar-deprived cells (Fig. 1B). We speculated that activation of trehalase, the enzyme that breaks down molecules of trehalose into glucose, would therefore occur prior to mobilization. Reports on neutral trehalase activity in \textit{C. albicans} exist (31), but activation by resupplementation of missing nutrients has never been documented. In wild type \textit{S. cerevisiae}, activation of neutral trehalase could be observed 2–5 min after glucose addition (Fig. 1C). However, only basal trehalase activity and no activation could be detected in \textit{C. albicans} strains.

Gpr1 in \textit{C. albicans} is required to control basal trehalose levels in particular when cells are metabolically active and growing in favorable conditions. Absence of the receptor seems to mimic a nutrient-poor environment for the cells, leading to trehalose accumulation. However, Gpr1 does not play a role in the induction of trehalose breakdown, typically observed upon sugar addition, which differs from the situation in \textit{S. cerevisiae}.

The substrates of trehalose are uridine diphosphoglucose and glucose 6-phosphate. They are converted into trehalose 6-phosphate in the first step of trehalose accumulation by the Tps1 enzyme (11). We aimed at determining whether increased trehalose content in \textit{gpr1}^{−}/\textit{gpr1}^{Δ} cells correlated with increased levels of glucose or glucose 6-phosphate. In none of the assays we performed we could detect a lower or higher level of these substrates in cells lacking \textit{GPR1}, either before or after induction with glucose (Fig. 2). Specifically, Gpr1 does not play a role in the transport of glucose (Fig. 2A), and it does not control the rise of glucose phosphorylation following glucose addition (Fig. 2B). The transient increase of glucose 6-phosphate coincides with a drop of ATP levels, which was observed in both wild type and \textit{GPR1}-deleted strains (Fig. 2C). The upregulated conversion to trehalose could mask any increased glucose 6-phosphate production, possibly resulting from elevated hexokinase activity. However, the total hexo- and glucokinase activities were not significantly different between the two strains (Fig. 2D). Although these data were required to determine possible routes of regulation of trehalose metabolism via Gpr1, they evidently suggest other level(s) of regula-

**FIGURE 2.** Increased trehalose levels in \textit{gpr1}^{Δ}/\textit{gpr1}^{Δ} are not associated with altered glucose transport, glucose 6-phosphate, and ATP levels, and hexokinase activity. \textbf{A}, glucose transport is measured by adding increasing concentrations of radioactively labeled glucose. Progressive shading from black to white represents increasing glucose concentrations (1, 5, 10, 20, 50, and 100 mM). Standard deviations (error bars) are calculated from three independent biological repeats. \textbf{B} and \textbf{C}, glucose 6-phosphate (\textbf{B}) and ATP (\textbf{C}) levels were determined after addition of 100 mM glucose. ●, WT; ○, \textit{gpr1}^{Δ}/\textit{gpr1}^{Δ}; ▲, \textit{gpr1}^{Δ}/\textit{gpr1}^{Δ} + \textit{gpr1}; †, \textit{gpr1}^{Δ}/\textit{gpr1}^{Δ} + \textit{GPR1}. \textbf{D}, basal hexokinase activity is determined. Standard deviations are calculated from three independent biological repeats.
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FIGURE 3. Increased trehalose levels in gpr1Δ/Δ correlate with aberrant activities of the trehalose metabolism and catabolism enzymes. TPS (A), TPP (B), and trehalase (C) activities are monitored over time. Standard deviations (error bars) are calculated from three independent biological repeats. An asterisk (*) indicates significant difference compared with wild type (p < 0.05). The shaded bars represent, from black to white: wild type, gpr1Δ/gpr1Δ, gpr1Δ/gpr1Δ + riGPR1.

Gpr1 Strongly Regulates TPS Enzymatic Activity—We measured TPS and TPP activities, the two main enzymatic activities responsible for trehalose synthesis, in *C. albicans* cells growing in glucose-containing medium for 3–72 h. A sharp rise in TPS activity was observed in gpr1Δ/gpr1Δ cells at 9 and 24 h of growth, coinciding with increased trehalose content at the same time of growth (Fig. 3A). The dephosphorylation step of trehalose 6-phosphate into trehalose by TPP was somewhat increased as well, but statistical significance could never be obtained (Fig. 3B). The steadily increased TPS activity is likely necessary to ensure trehalose 6-phosphate to be continuously converted into trehalose, as accumulation of this intermediate molecule is toxic to cells (19, 32). Transcriptional regulation of the TPS1 gene was not modulated differentially in wild type and gpr1Δ/gpr1Δ cells (data not shown), suggesting a post-translational regulatory mechanism. An increase of trehalose content in *C. albicans* is mostly accompanied by a decrease in neutral trehalase activity, at least as a response to heat or oxidative stress (33, 34). In nonstress conditions, neutral trehalase activity increased over time during growth of wild type cells (Fig. 3C). In absence of GPR1, trehalase activity could never be demonstrated as significantly different from wild type levels. Nevertheless, partial impairment of this activity was always observed in the mutant strain. To corroborate these findings, a strong and opposite direct effect on trehalose metabolism was apparent upon expression of *C. albicans* GPR1 in a strain of *S. cerevisiae* deleted for the homologous receptor (Fig. 4). Trehalose content was significantly reduced as well as TPS activity in this strain. Similar to the assays performed in *C. albicans* cells, trehalase and TPP activities seemed to be differentially regulated in the presence of *C. albicans* Gpr1, but the difference could not be determined as significant. Hence, the aberrant trehalose accumulation observed in *C. albicans* gpr1Δ/gpr1Δ cells seems to result from a higher rate of synthesis, mainly governed by post-transcriptional regulation of Tps1 and a lower catabolism.

Heat-induced Morphogenesis Is Impaired by Increased Trehalose Levels—*C. albicans* wild type cells typically undergo a morphogenetic switch from yeast to filamentous forms at elevated temperature (>35 °C) in growth conditions with neutral pH (35). We had previously shown that deletion of GPR1 impaired invasive growth on solid media but not in liquid cultures in presence of serum (7). However, we show here that the gpr1Δ/gpr1Δ mutant was found to be impaired in the morphogenetic switch induced by elevated temperature (Fig. 5A, panels 1 and 2). In the conditions tested, the mutant did not display any reduced growth rate.

We hypothesized that presence of elevated trehalose content in gpr1Δ/gpr1Δ cells at 37 °C may alter the heat-induced filamentation phenotype in the mutant strain. To further assess the role of trehalose in *C. albicans* cell elongation, we made use of validamycin, a specific inhibitor of trehalase, the enzyme breaking down trehalose. Validamycin has *in vitro* activity against several fungal trehalases, including the one of *S. cerevisiae* (36) and *C. albicans* (31). *In vivo* activity has been demonstrated against the trehalase enzyme of *Rhizoctonia solani*, the pathogen responsible for sheath light of rice plants (37). We demonstrate here that addition of increasing amounts of the trehalose analog led to increasing amounts of trehalose being accumulated *in vivo* (Fig. 5B), without altering cell growth rate (data not shown). As little as 5 μM compound ensured a 100% increase in intracellular trehalose levels. The increase in treha-
Trehalose coincided with a reduction up to 75% of elongated cell phenotypes observed at 37 °C, as the concentration of the inhibitor increased. Wild type cells treated with validamycin had a clear reduction in cell elongation (Fig. 5A, panel 3). These data demonstrate for the first time that trehalose is a potent inhibitor of morphogenesis in *C. albicans*.

**Trehalose Restrains Cell Elongation by Release of Hsp90 Inhibitor** —The specific condition of cellular transition caused by elevated temperature is orchestrated by the molecular chaperone Hsp90 in *C. albicans* (22). Compromising Hsp90 function, by elevated temperature or chemical inhibitors, such as geldanamycin, leads to release of inhibition imposed on the morphogenetic program. We speculated that elevated trehalose levels could lead to a defect in the filamentation program governed by Hsp90 when active at 37 °C. We therefore examined whether the induction of filamentation resulting from Hsp90 inhibition by means other than temperature was also impaired in *gpr1Δ* cells. As *gpr1Δ* cells accumulate trehalose even at 30 °C while actively growing (data not shown and Ref. 30), we investigated whether geldanamycin, leads to release of inhibition imposed on the morphogenetic program. We speculated that elevated trehalose levels could lead to a defect in the filamentation program governed by Hsp90, as it impaired cell elongation at 37 °C. We therefore examined whether the induction of filamentation resulting from Hsp90 inhibition by means other than temperature was also impaired in *gpr1Δ* cells. As *gpr1Δ* cells accumulate trehalose even at 30 °C while actively growing (data not shown and Ref. 30), we investigated whether geldanamycin was able to induce the filamentation phenotype in the *gpr1Δ* mutant strain. Whereas extrusions of the cells were readily observed in wild type upon addition of geldanamycin at the noninducing temperature of 30 °C, the *gpr1Δ* mutant was partially defective in this response (Fig. 5A, panels 4 and 5). In addition, the observed filamentation in response to the Hsp90 inhibitor in an *ira2Δ* mutant and absence thereof in a *cdc25Δ* or *cdc35Δ* strain and avirulence of a *gpr1Δ* strain suggested a connection between the G protein-coupled receptor and trehalose metabolism, a feature suggested to be downstream of PKA (30). It is now clear from the present findings that these phenotypes were caused by the role of Gpr1 in negatively regulating the trehalose biosynthesis activity when cells are actively growing under optimal conditions and do not require trehalose as a compatible solute. Typically, trehalose is synthesized at the onset of nutri-

**DISCUSSION**

*C. albicans Gpr1 Participates in the Maintenance of Trehalose Homeostasis at the Human Body Temperature* —The G protein-coupled receptor Gpr1 in *C. albicans* was identified based on its high level of amino acid sequence identity with the ortholog of the model yeast *S. cerevisiae*. Despite the high degree of sequence homology between the two, their functions have diverged substantially, the most striking difference being the apparent loss of its role in fast glucose signaling to adenylate cyclase for cAMP production (7). A mild induction of TPS activity at 43 °C in the *gpr1Δ* strain and avirulence of a *gpr1Δ* strain suggested a connection between the G protein-coupled receptor and trehalose metabolism, a feature suggested to be downstream of PKA (30). It is now clear from the present findings that these phenotypes were caused by the role of Gpr1 in negatively regulating the trehalose biosynthesis activity when cells are actively growing under optimal conditions and do not require trehalose as a compatible solute. Typically, trehalose is synthesized at the onset of nutri-

**FIGURE 4**. Expression of *C. albicans GPR1* in a *S. cerevisiae gpr1Δ* strain mainly results in absence of trehalose accumulation and a reduced TPS activity. Trehalose levels (A), TPS (B), TPP (C), and trehalase activity (D) are measured in cells grown for 24 h in rich medium. Standard deviations (error bars) are calculated from three independent biological repeats. An asterisk (*) indicates significant difference compared with wild type (*p* < 0.05).
ent depletion. Hydrolysis of trehalose upon readdition of nutrients is equally important, as it is an essential event in many cellular processes such as fungal spore germination, insect flight, and the resumption of growth in resting cells (38), as well as germ tube formation in *C. albicans* (39). This tight regulation of trehalose content is perturbed in a *gpr1Δ/gpr1Δ* strain, due
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to an increased activity of the trehalose synthesis enzymes Tps1 and Tps2 and a concomitant inactivation of neutral trehalase. One hypothesis is that in absence of Gpr1, the activity of the Ras-PKA pathway of *C. albicans* is lowered during exponential growth, which in turn reduces the repression of the trehalose biosynthesis enzymes. To gain support for this hypothesis is the finding that several mutants of components of the pathway display an aberrant trehalose content too. Mutants in genes that activate the pathway, such as CDC25 and CDC35, display a higher trehalose content than wild type, whereas mutants of inhibitors of the pathway, such as *IRA2*, present lower levels of trehalose. Contrary to previous reports (20), we could identify four and six potential PKA phosphorylation sites of the consensus sequence (R/K)X1–2(S/T) in Tps1 and Tps2, respectively, with two in each protein having the second most abundant consensus sequence KKX(S/T) (40). The pathway may therefore regulate enzymatic activities via phosphorylation by PKA rather than gene expression in *C. albicans*.

It is noteworthy to state that this phenomenon of trehalose accumulation in *gpr1Δ/gpr1Δ* cells is a glucose-dependent process, in contrast to the well characterized accumulation of the carbohydrate when glucose gets depleted. Gpr1 therefore seems to behave as a glucose sensor, yet not involved in fast signaling of glucose, but rather involved in the long term maintenance of glucose signaling leading to trehalose metabolism regulation.

**Tight Control of Trehalose Levels Is Required for Filamentation**—We used the trehalase inhibitor validamycin to validate the finding that trehalose is the causative agent that diminishes the potency of *gpr1Δ/gpr1Δ* cells to elongate. We establish the potency of validamycin to increase trehalose content in vivo in *C. albicans*. Addition of validamycin results in 100% increase in trehalose levels and simultaneous reduction of 75% in cell elongation. This links morphogenetic differences between wild type and *gpr1Δ/gpr1Δ* cells to trehalose content. Indeed, the observed reduction of cell elongation as well as trehalose accumulation of the *gpr1Δ/gpr1Δ* strain at 37 °C was comparable with that of the wild type strain treated with validamycin. The data here demonstrate that although high levels of trehalose in the *gpr1Δ/gpr1Δ* strain are not detrimental to cell growth, they diminish the ability of the mutant cells to undergo the morphogenetic switch. Hence, increased trehalose content correlates with inefficient cell elongation at 37 °C in *C. albicans*. Coincidentally, other mutants of the Ras-PKA pathway also display altered phenotypes at 37 °C, which remarkably correlates with their trehalose content (22). These findings emphasize the biological connection among the Ras-PKA pathway, filamentation, and trehalose content.

**A Biological Link among the Storage Carbohydrate, Filamentation, and Virulence Originates from the Membrane Receptor Gpr1 in *C. albicans***—Heat-induced filamentation in *C. albicans* is centered on the relief of the heat shock chaperone Hsp90 from the morphological signaling machinery (22). It can be mimicked efficiently at 30 °C by addition of geldanamycin, an Hsp90 inhibitor. Here, we propose that trehalose serves as a novel regulatory factor of the Hsp90-mediated cell elongation program. Down-regulation of PKA signaling leads to trehalose accumulation, which in turn accentuates Hsp90 function. As an important part of the heat shock response, trehalose binds to partially folded intermediate conformations of proteins (41). It has been reported that trehalose binds to heat shock proteins and can chaperone their activity (41, 42). However, whether trehalose aids the function of the chaperone, or vice versa, remains unclear. In the context of heat-induced filamentation, it becomes apparent from our findings that trehalose reinforces the chaperone function of Hsp90 on some of its client proteins belonging to the cell elongation pathway. The possibility that trehalose directly inhibits components of the Ras-PKA pathway cannot be excluded. However, serum, which activates PKA as well induces the morphogenetic program, is a potent activator of cell elongation in the *gpr1Δ/gpr1Δ* strain (7). This tends to indicate that the effect of trehalose is specific on Hsp90, which governs filamentation in response to heat.

Studies on trehalose relate to its physical and chemical properties or to the regulation of its biosynthesis and degradation, without directly addressing the interaction point between the carbohydrate and the targeted molecular process. In other words, we often know that trehalose is involved, but rarely do we know how and at which molecular level it plays its function. In the present study, we have identified a clear biological link among the Ras-PKA pathway, the increased trehalose synthesis, and the filamentation process governed in response to the human body temperature. We can also extrapolate from our findings on filamentation that this molecular link is likely to be the chaperone protein Hsp90.

**How May Trehalose Inhibit the Morphogenetic Switch?**—The present findings raise the interesting question of how a sugar molecule can modulate the signaling process leading to cell differentiation. At high temperature, trehalose can serve the role of a chaperone, which protects proteins from heat-induced denaturation and aggregation. Trehalose is a very potent protein stabilizer, in that it keeps proteins in native states in unfavorable conditions (43). Hence, overproduction of trehalose may overwhelm normal protein functioning (44). The obvious stabilized target in this study is the chaperone Hsp90, the central player in heat-induced filamentation in *C. albicans* (22). In that context, the presence of high trehalose content at inappropriate timing in a *gpr1Δ/gpr1Δ* strain or other positive regulators of the PKA pathway may maintain Hsp90 in a stable conformation, ensuring effective inhibition of the filamentation program even at the permissive temperature. Stable Hsp90 binds to its client proteins, disabling their function (45).

Second, the influence of trehalose on Hsp90 may be mediated through Hsf1, rather than Hsp90 itself. The hyperphosphorylation of Hsf1 leading to HSP expression, generally seen after heat shock, has been shown to be dependent upon mild increase of trehalose levels in *S. cerevisiae* (46). In *C. albicans*, 86% of up-regulated genes after heat shock, especially the HSP family genes, depend on Hsf1 for their up-regulation (47). In *C. albicans*, the increase in trehalose levels may lead to the repression of Hsp90 relief through the phosphorylation status of Hsf1. As Hsf1 participates in the basal expression of 75 genes even in the absence of heat shock, the essential transcription factor

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