

The plant hormone indoleacetic acid induces invasive growth in *Saccharomyces cerevisiae*

Reeta Prusty, Paula Grisafi, and Gerald R. Fink*

Whitehead Institute for Biomedical Research, Nine Cambridge Center, Cambridge, MA 02142

Contributed by Gerald R. Fink, January 29, 2004

Fungi must recognize plant-specific signals to initiate subsequent morphogenetic events such as filamentation that lead to infection. Here we show that the plant hormone indoleacetic acid (IAA) induces adhesion and filamentation of *Saccharomyces cerevisiae*. Genome expression profiling of cells treated with IAA identified Yap1, a fungal specific transcription factor, as a key mediator of this response. Strains lacking *YAP1* (*yap1-1*) are hypersensitive to growth on IAA because they accumulate more IAA than can wild type. Members of a family of transporters the amino acid/auxin:proton symport permeases with homology to *AUX1*, a putative IAA transporter from plants, are up-regulated in the *yap1-1* mutant. Deletion of any one of these transporters makes *yap1-1* mutants more resistant to IAA by decreasing its uptake. The permease mutants are defective in IAA perception and filamentation. The ability of a fungus to perceive a plant hormone that causes it to differentiate into an invasive form has important implications for plant-pathogen interactions.

Infection of a plant by a fungus involves mutual chemical recognition between the host and the pathogen. Plants recognize and respond to compounds such as chitin, which are unique signatures of fungal and insect pathogens (1). Reciprocally, fungi recognize chemical cues that signal the presence of the plant host and induce the subsequent morphogenetic changes that enable invasion. Fungi alight on a plant in a cellular form either as a yeast cell or spore and, after sensing the plant, proliferate into an invasive filamentous form. For example, spores of *Colletotrichum gloeosporioides* elaborate an invasion structure (appressorium) in response to ethylene, a plant hormone that promotes fruit ripening (2). By homing in on the plant hormone, the fungus synchronizes its invasion with a nutritionally advantageous stage in plant development.

Indoleacetic acid (IAA) is a molecule that is synthesized by plants and a few microbes (3, 4). In plants, IAA plays a key role in both root and shoot development. The hormone moves from one part of the plant to another by a designated importer (*AUX1*) and efflux pumps (*PINI-7*) (5, 6). In peas, wounding or application of IAA induces the expression of *ACS2*, an IAA-responsive gene, suggesting that IAA may be present at the site of a wound (7). The presence of IAA at the wound site would make it a distinctive attractant for fungi; however, there is limited evidence for such a role. One difficulty in establishing the connection between IAA and pathogenesis is the lack of an *in planta* assay for the concentration of IAA at the foci of infection.

In this study, we take a different approach and ask whether the fungus *Saccharomyces cerevisiae* is able to perceive the phytohormone IAA. Our results show that *Saccharomyces* distinguishes IAA from other closely related compounds. At high concentrations, IAA inhibits growth, whereas at lower concentrations, it induces filamentation and adhesion. These responses are mediated by a family of transporters and the fungal transcription factor Yap1.

Materials and Methods

Strains and Media. Growth and media conditions were those described (8, 9). The *S. cerevisiae* strains are isogenic to BY4742 [the S288c background that was used for the Yeast Genome

Sequencing project and the deletion project (10)]. Strains in the S288c background have a *flo8* mutation, which prevents pseudohyphal growth; therefore, pseudohyphal growth and microarray studies were performed by using cognate deletions made in the Σ 1278b background. Both S288c and Σ 1278b are sensitive to IAA. Deletion strains were either from the Euroscarf deletion library or were constructed by replacement of the relevant ORF with the G418 resistance marker (11). The *Ustilago maydis* strain, (*a1 b1*, wild-type), used in this study was provided by S. Gold (University of Georgia, Athens).

Compounds Tested. More than 50 compounds were tested for differential growth inhibition (for a complete list, see *Data Set 1*, which is published as supporting information on the PNAS web site). These compounds included plant hormones (such as salicylic acid, ethylene, cytokinin, gibberellic acid, and abscisic acid), several IAA-conjugated compounds, and other indoles (such as indole, indole-3-pyruvic acid, and indole-3-acetamide), mimics and inhibitors of IAA transport (such as 2,4-(dichlorophenoxyacetic) acid and 2,3,5-triiodobenzoic acid), weak acids (sorbic acid and benzoic acid), and agents known to arrest the growth of yeast (azetidine, canavanine, and ethionine).

IAA Uptake Assay. The whole-cell assay used yeast strains grown at 30°C to an OD₆₀₀ of ≈ 0.4 in minimal media supplemented with required amino acids. To measure IAA uptake, cells were harvested by centrifugation at 1,000 $\times g$ and were resuspended in Mes buffer at pH 4.6 with 2% glucose (12). Duplicate samples were removed after the addition of cells to medium containing ³H-labeled IAA (Amersham Pharmacia, Piscataway, NJ) 220 μ Ci/mmol (1 Ci = 37 GBq). Cells were collected on glass-fiber filters and were washed extensively with buffer. The filters were placed in glass scintillation vials, were treated with 350 μ l of 1 M NaOH for 10 min, and the solution was neutralized with 400 μ l of 1 M acetic acid. Uptake of labeled nutrients into cells was determined by using 5 ml of liquid scintillation fluid. The radioactive material extracted from cells comigrates with pure IAA in thin-layer chromatography. Uptake of IAA is not due to nonspecific binding of IAA because only live cells are capable of its transport. For 2,4-dinitrophenol (DNP) or carbonyl cyanide *m*-chlorophenylhydrazone (CCCP) treatment, cells were exposed to 1 mM DNP or 500 μ M CCCP for 45 min before the uptake assay. Tryptophan uptake was measured in the same whole cell assay, except 20 μ Ci/mmol ³H-tryptophan was used instead of ³H-IAA. The *yap1-1* and *avt yap1-1* double mutants show the same level of tryptophan uptake as wild-type cells; therefore, the *yap1-1* mutation does not lead to a general elevation in the uptake of indoles (Fig. 5, which is published as supporting information on the PNAS web site).

Protein Localization. GFP was placed at the C terminus of Avt3 and Avt4 by site-directed PCR. The resulting in-frame fusion

Abbreviations: IAA, indoleacetic acid; AAAP, amino acid/auxin:proton symport permeases; DNP, 2,4-dinitrophenol; CCCP, carbonyl cyanide *m*-chlorophenylhydrazone.

*To whom correspondence should be addressed. E-mail: gfink@wi.mit.edu.

© 2004 by The National Academy of Sciences of the USA

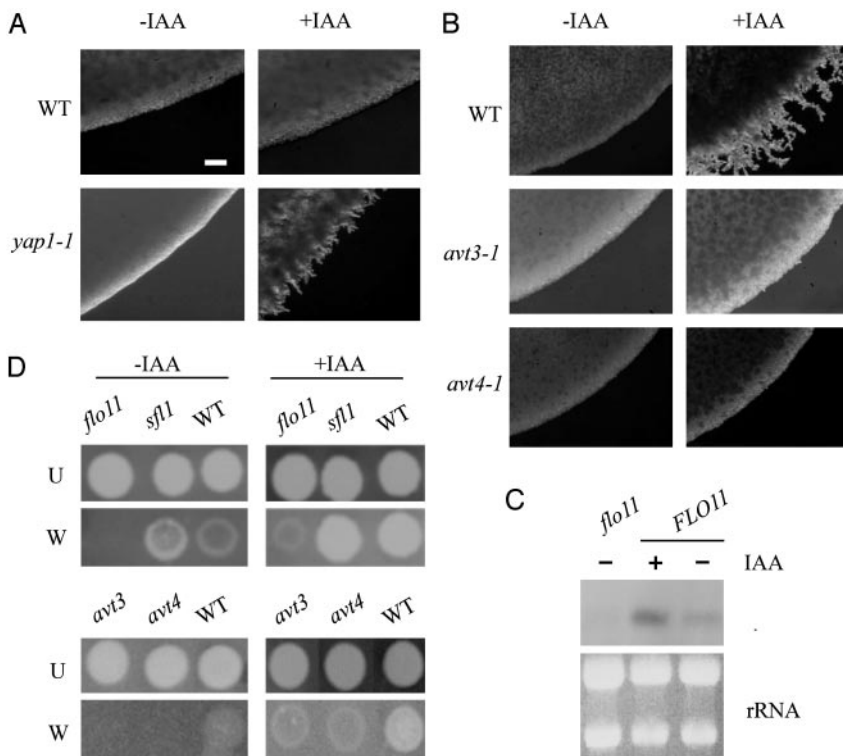


Fig. 1. IAA enhances filamentation and surface adhesion. *A* and *B* show the edge of a yeast patch ($\Sigma 1278b$) growing on filamentation-inducing media (8) with xylose as the carbon source. Plates were incubated in the dark to prevent photodegradation of IAA. For the purpose of comparing the images, *A* and *B* were taken at the same magnification. (Bar in *A*, 10 μm .) (*A*) The *yap1-1* mutant filaments at low levels of IAA (2 μM), and wild type does not. (*B*) Wild-type filaments at higher concentrations of IAA (50 μM). IAA-enhanced filamentation is abolished in all seven *avt* mutants (*avt3-1* and *avt4-1* are shown). (*C*) Northern blot analysis shows that IAA induces expression of *FLO11*. (*D*) IAA enhances haploid invasive growth of *S. cerevisiae*. Cells were patched on media containing IAA (120 μM). U, unwashed; W, washed.

proteins, *Avt3::GFP* and *Avt4::GFP*, were shown to be functional because they suppressed growth inhibition by IAA and restored IAA uptake in *avt* mutants. Strains were grown on minimal media and were visualized on a Zeiss LSM 510 confocal microscope.

Results

IAA Induces Invasive Growth in *S. cerevisiae*. A number of plant-specific compounds were tested for their ability to inhibit either the growth of yeast or to induce filamentation (see *Data Set 1* for a complete list of compounds tested). Diploid yeast cells develop into a multicellular, pseudohyphal form (invasive filaments of elongated cells) under conditions of nitrogen starvation. Haploid cells adhere to the agar surface of rich media plates (haploid invasion). Of the many compounds tested, only IAA had effects in these assays. IAA enhances both haploid invasion and diploid pseudohyphal growth (Fig. 1 *B* and *D*). The enhancement of these phenotypes by IAA is optimal when xylose is the carbon source. The recognition of IAA by yeast is specific; other plant hormones and indoles closely related in structure to IAA fail to show induction of these phenotypes.

Induction of morphogenesis by IAA requires the function of Flo11, a cell-surface protein known to be required for the nutritional induction of both diploid filamentation and haploid invasive growth (13). Growth of cells in the presence of IAA leads to an increase in *FLO11* mRNA (>3-fold, Fig. 1*C*), suggesting that IAA induces morphogenetic changes by elevating *FLO11* transcription. Induction of adhesion and pseudohyphal growth by IAA is dramatically reduced by mutations that diminish *FLO11* expression (*flo11*, *ste20*, *ste11*, and *tec1*; data not shown) and enhanced in mutants that have elevated levels of *FLO11* (*sfl1*, Fig. 1*D*).

At higher concentrations, IAA, but not other indoles tested, inhibits the growth of yeast (Fig. 2*A*) both in liquid ($\text{EC}_{50} = 250 \mu\text{M}$) and on solid media (Fig. 2*B* and *C*). IAA arrests growth in all stages of the cell cycle and, after removal from IAA, cells resume growth. Several distinct laboratory strains, including

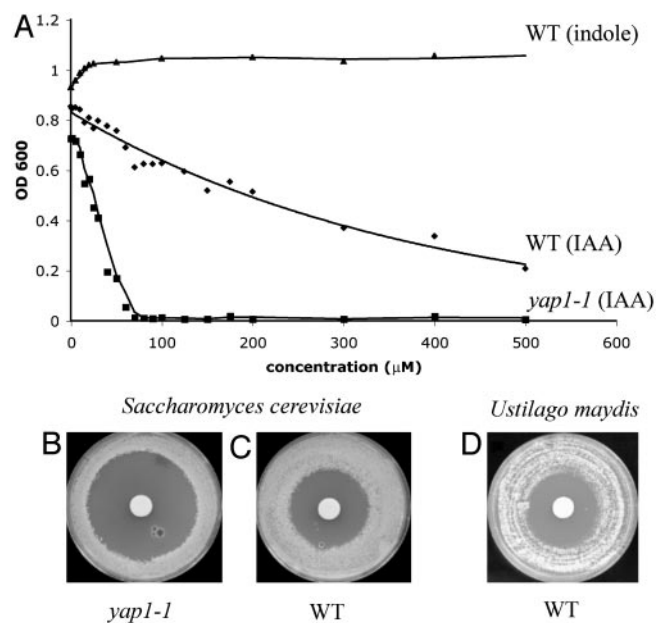


Fig. 2. IAA growth inhibition in liquid. (*A*) Cells were treated with various concentrations of IAA or other compounds. The EC_{50} is the concentration of IAA at which growth is the half-maximum obtainable in the absence of the drug. Cells were inoculated into medium in 96-well plates. The OD at 600 nm was measured after 2 days of incubation. Each data point is the average of two samples that varied by <10%. The *yap1-1* mutant is hypersensitive to IAA ($\text{EC}_{50} = 25 \mu\text{M}$) compared with wild type ($\text{EC}_{50} = 250 \mu\text{M}$). Indole and many other indolic compounds tested do not inhibit growth. (*B–D*) IAA growth inhibition on Petri plates. A filter disk saturated with IAA was placed on a lawn of cells. Plates were incubated for 3 days in the dark. The clear area indicates the zone of growth inhibition. The *yap1-1* strain (*B*: *yap1-1*, *C*: wild type) is hypersensitive to IAA. Approximately 50 related compounds were tested by using this assay (see *Data Set 1*). Of these compounds, the *yap1-1* mutant is only slightly sensitive to fluoro-IAA, indole pyruvate, and benzoic acid. (*D*) IAA-inhibition of *Ustilago maydis* growth.

S288c and $\Sigma 1278b$, were sensitive at approximately the same IAA concentrations. The inhibition by IAA is specific because other plant hormones and indolic compounds, even those structurally very similar to IAA, such as indole butyric acid and 5-hydroxy IAA, do not inhibit growth at comparable concentrations (see *Data Set 1*). IAA also inhibits the growth of the *Ustilago maydis*, a fungal pathogen of corn (Fig. 2D).

Transcription Profiling and Deletion Library Screen Implicate Yap1 in the IAA Response. Whole-genome arrays show that the transcription profile of cells growing in the presence of subtoxic levels of IAA is unique and distinct from those of cells exposed to many other diverse environmental stress conditions (Fig. 5). Of the 126 stress-induced genes identified in two independent studies (14, 15), only four are also induced after IAA exposure (Fig. 3A). Of the 99 genes whose transcripts were induced by IAA, 28 contain a consensus Yap1-binding site within 1 kb of the transcription start site (Fig. 3B). Of these genes, 10 have been shown previously to be regulated by Yap1 (asterisks, Fig. 3B, see the *Saccharomyces* Genome Database, which can be accessed at www.yeastgenome.org), whereas the remaining 18 genes have perfect matches to the Yap1-binding site, TTACTAA (open circles, Fig. 3B).

A screen of the complete yeast deletion library (10) for IAA-hypersensitive mutants identified *yap1-1*; no other mutant in the entire library is as IAA-sensitive as *yap1-1* ($EC_{50} = 20 \mu M$, Fig. 2A). In particular, mutations in paralogs of *YAP1* (especially *YAP2*) are not hypersensitive. Furthermore, the *yap1-1* mutant is induced to filament at concentrations of IAA ($2 \mu M$) that are >20-fold lower than those required to induce filamentation in wild type ($50 \mu M$) (compare Fig. 1A and B). These data suggest that Yap1 is the key mediator of the response to IAA.

The Avt Proteins Are IAA Transporters. The specificity of IAA recognition for morphogenesis and growth inhibition suggests that yeast has transporters that recognize IAA. A search of the yeast genome for possible components of an IAA uptake system identified a family of seven genes [*AVT1-7* (16)] annotated as the amino acid/auxin:proton symport permease (AAP) family, which is part of a larger family that includes *AUX1*, an Arabidopsis gene implicated in IAA transport in plants (17, 18) and γ -aminobutyric acid transporters in *Caenorhabditis elegans* (19). Deletion of any one of the yeast *AVT* family members makes *yap1-1* mutants more resistant to IAA (Fig. 6, which is published as supporting information on the PNAS web site), suggesting that deletion of these putative transporters reduces uptake of this compound. Mutations in other permease genes (the general amino acid permease *GAP1*, the tryptophan permease *TAT2*, or the pleiotropic drug-resistant pump *PDR12*) do not alter the sensitivity of *yap1-1* mutants to IAA (Fig. 4C).

To determine whether the *AVT* genes mediate the transport of IAA, we designed an assay to measure the ability of intact yeast cells to take up IAA from the medium. In this assay, the *yap1-1* mutant shows increased uptake of 3H -IAA, as compared with wild type (Fig. 4A). The *yap1-1* mutation does not lead to a general elevation in the uptake of indoles (Fig. 6B). This enhanced uptake of IAA in the *yap1-1* mutant is reduced to wild-type levels in the presence of DNP or CCCP, compounds that collapse the proton gradient (Fig. 4B). These data suggest that the uptake of IAA has two components, one of which requires an intact membrane potential.

The steady-state mRNA levels for *AVT3* and *AVT4* are elevated (>2 fold) in *yap1-1* strains, as compared with wild type (Fig. 3C), which could account for the increased IAA uptake of *yap1-1*. As predicted by this model, the *avt yap1-1* double mutant strains show a decreased rate of IAA uptake as compared with the *yap1-1* mutants (Fig. 4A). It is the energy-dependent component of uptake that is missing in the *avt yap1-1* strains (Fig.

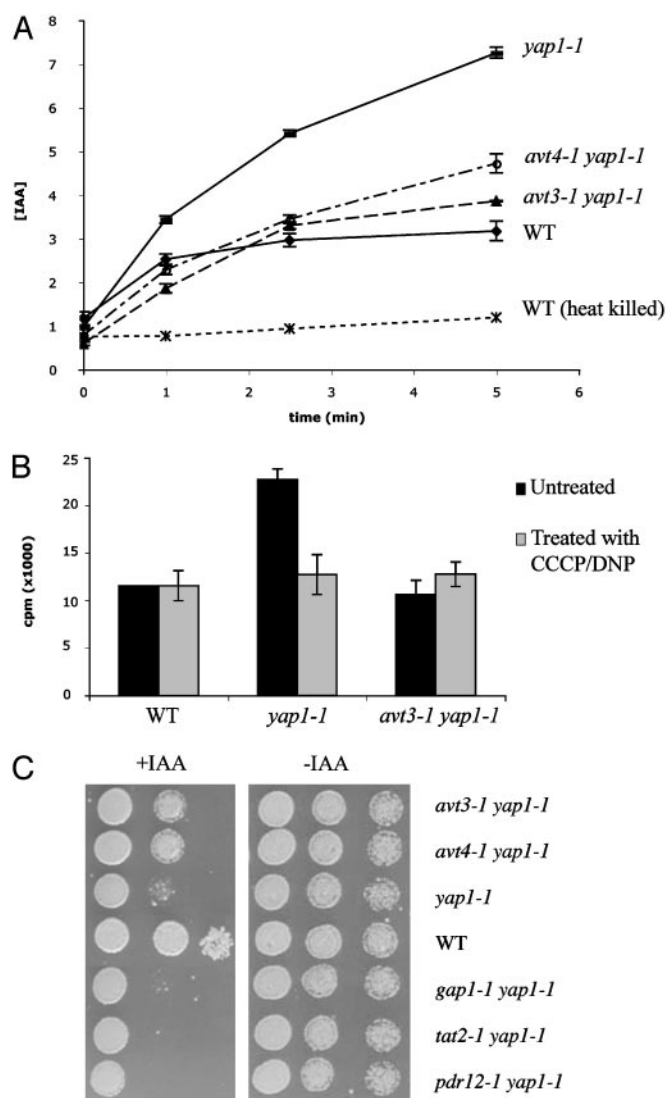


Fig. 4. Uptake of 3H IAA into yeast cells. Cells grown to log phase in minimal media (9) were harvested and incubated in the presence of 3H -IAA for various lengths of time. Cells were filtered, and total counts retained on glass filters were monitored. (A) The *yap1-1* strain accumulates more IAA than wild type; heat-killed (HK) cells show no IAA uptake. The *avt3-1 yap1-1* and *avt4-1 yap1-1* double mutants (dashed lines) show decreased IAA uptake. Other *avt yap1-1* double mutants also reduce IAA uptake (Fig. 6A). IAA concentration is represented in μ mol of IAA transported. (B) IAA accumulation depends on an intact membrane potential. Cells were treated with CCCP or DNP for 45 min at room temperature before IAA exposure. The bar graph compares IAA accumulation at 5 min with and without drug treatment. 3H -IAA transported is represented in cpm. (C) Deletion of any *AVT* gene makes *yap1-1* mutants more resistant to IAA. Deletion of other transporters, such as the general amino acid permease (*GAP1*), the high-affinity tryptophan permease (*TAT2*), or the pleiotropic drug resistant pump (*PDR12*), does not alter the IAA sensitivity of the *yap1-1* strain.

4B). The reduced IAA uptake in the *avt yap1-1* strains correlates with their reduced sensitivity to IAA. Consistent with their reduced uptake, the *avt* mutants fail to show the IAA induction of either haploid adherence (Fig. 1D) or diploid filamentation (Fig. 1B). Mutation in any one of the seven *AVT* genes abolishes the effects of IAA. This lack of redundancy suggests that the Avt proteins either function in the same pathway or are components of a multiprotein complex involved in some aspect of IAA transport into yeast.

