Transcriptome Profiling of *Botrytis cinerea* Conidial Germination Reveals Upregulation of Infection-Related Genes during the Prepenetration Stage

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Transcriptome studies with *N. crassa*, *Fusarium graminearum*, and *Ustilago maydis* have been performed using spores germinated in minimal (*N. crassa* and *U. maydis*) or full (*F. graminearum*) media in (partly shaken) suspensions, with no permanent surface contact of the spores (17, 18, 19). *Blumeria* microarray-based studies with germinated conidia forming infection structures, including haustoria, on the surface and within host epidermal cells have focused mainly on transcriptomic changes related to metabolic pathways, such as lipid degradation, glycolysis, pentose phosphate pathway, and amino acid biosynthesis (20, 21). Comprehensive studies with *Magnaporthe oryzae* have addressed gene expression changes, after 4 to 12 h, that are mainly correlated with appressorium formation (22, 23). The study presented here focused on gene expression changes during the early stages of conidial germination and appressorium differentiation of *Botrytis cinerea*, in response to a combination of physical (hard surface) and chemical (sugars, plant lipids) signals.

*Botrytis cinerea* (teleomorph *Botryotinia fuckeliana*) is a major necrotrophic pathogen of a wide host range of dicotyledonous

Germination of spores is a fundamental step in fungal development, leading to the conversion of a dormant cell into a growing hypha. It involves the breaking of dormancy by external signals, a pregermination phase that is visibly accompanied by isotropic swelling, and then the formation of a germ tube that marks the establishment of polar growth (1, 2). Basic requirements for the initiation of germination of fungal spores, for example, of the saprotrophic fungi *Neurospora crassa* and *Aspergillus nidulans*, are usually high humidity and the availability of nutrient sources such as sugars (2). Spore germination in response to sugars has been shown in *A. nidulans* to be dependent on Gα subunits, cyclic AMP (cAMP), and protein kinases (3, 4). Similarly, the cAMP-protein kinase A pathway is required for spore germination in *Schizosaccharomyces pombe* (5) and for yeast-to-hypha transition on solid media in the pathogenic yeast *Candida albicans*, a process related to spore germination (6). In plant-pathogenic fungi, including *Blumeria graminis* (7), *Phyllosticta ampelica* (8), *Colletotrichum graminicola* (9), and *Colletotrichum gloeosporioides* sp. *aeschnomyces* (1), breaking of spore dormancy is often induced by surface contact stimuli, such as surface hardness and hydrophobicity, either alone or in combination with nutrient signals. Although some signal transduction components, such as cAMP, Ras2, and Rac1, have been found to be involved in the germination process of these fungi, the regulatory mechanisms are still poorly understood (1, 9, 10). After germination on appropriate surfaces such as host cuticles, germ tube elongation occurs until perception of physical and/or chemical stimuli by the germ tube tip, resulting in the cessation of germ tube elongation and appressorium formation (7, 11, 12, 13, 14, 15). Appressoria, which are morphologically highly diverse in different plant pathogens, initiate the penetration into the plant tissue (16).

Botryotinia fuckeliana causes gray mold on a great number of host plants. Infection is initiated by airborne conidia that invade the host tissue, often by penetration of intact epidermal cells. To mimic the surface properties of natural plant surfaces, conidia were incubated on apple wax-coated surfaces, resulting in rapid germination and appressorium formation. Global changes in gene expression were analyzed by microarray hybridization between conidia incubated for 0 h (dormant), 1 h (pregermination), 2.5 h (postgermination), 4 h (appressoria), and 15 h (early mycelium). Considerable changes were observed, in particular between 0 h and 1 h. Genes induced during germination were enriched in those genes encoding secreted proteins, including lytic enzymes. Comparison of wild-type and a nonpathogenic MAP kinase mutant (bmp1) revealed marked differences in germination-related gene expression, in particular related to secretory proteins. Using promoter-GFP reporter strains, we detected a strictly germination-specific expression pattern of a putative chitin deacetylase gene (*cda1*). In contrast, a cutinase gene (*cutB*) was found to be expressed only in the presence of plant lipids, in a developmentally less stringent pattern. We also identified a coregulated gene cluster possibly involved in secondary metabolite synthesis which was found to be controlled by a transcription factor also encoded in this cluster. Our data demonstrate that early conidial development in *B. cinerea* is accompanied by rapid shifts in gene expression that prepare the fungus for germ tube outgrowth and host cell invasion.
plant species, causing serious pre- and postharvest losses in the production of economically important fruits, vegetables, and ornamental flowers (24). New infections usually occur by wind-dispersed conidia, which germinate on the plant surface and invade the tissue either through wounds or by direct penetration of cuticles and cell walls of epidermal cells (25). The host tissue is rapidly killed and degraded by various mechanisms, such as secretion of cell wall-degrading enzymes and release of toxins and other necrosis-inducing factors (26). Because of the crucial role of *B. cinerea* conidia for epidemiology and infection, a detailed knowledge of the molecular events during the early stages of their development is of great interest not only scientifically but also for the development of new control strategies.

Germination of *B. cinerea* conidia can be induced either in rich media in the absence of any surface, in the presence of carbon sources on a hard surface, or without any nutrients on a hard and hydrophobic surface (27). In the presence of carbon sources, germ tube outgrowth is preceded by rapid turnover of the disaccharide trehalose, which serves as a carbohydrate reserve and as an osmo-protectant (28). Based on the germination phenotypes of mutants, two signaling pathways have been described that are involved in the control of germination induction. For normal carbon-source-induced germination on hard surfaces, an intact cAMP signaling pathway, which includes Ras2, the Gα3 subunit of the heterotrimeric G protein, and adenylate cyclase, is required (27, 29). In contrast, hydrophobic-surface-induced germination is dependent on a functional mitogen-activated protein kinase (MAPK) cascade (Ste11-Ste7-Bmp1) (27, 30). *B. cinerea* mutants lacking either of the three MAPK components, or the putative adaptor protein Ste50, did not germinate on hydrophobic surfaces in pure water (30). After germination, the germ tube elongates on the host surface to different lengths before tip growth stops and penetration is initiated. The appressorium morphology is not very distinct, being characterized by a more or less distinct terminal swelling and, usually, no septum separating it from the germ tube. Penetration into the host tissue is accompanied by rapid death of the invaded plant cells. In addition to their role in germination, the cAMP and the Bmp1 MAPK signaling pathways are required for appressorium differentiation, penetration, and host colonization, similar to the situation in other plant-pathogenic fungi (31, 32). The MAPK cascade ste11, ste7, bmp1, and ste50 mutants did not form appressoria on hard surfaces, they failed to penetrate into host tissue, and they were nonpathogenic (27, 30). Other signaling elements found to be involved in germination are the small GTPase Cdc42 and phospholipase C (Plc1) (33). Taken together, these findings show that germination induction is a complex process involving several signaling pathways which is currently not fully understood.

Transcriptome profiling analyses during germination have been published for *N. crassa* (17), the saprotrophic chytridiomycete *Blastocladiella emersonii* (18), and the wheat pathogen *F. graminearum* (19). These studies revealed major changes of gene expression patterns starting already during the pregermination swelling stage. Genes involved in DNA replication, protein synthesis, and degradation of storage reserves were found to be activated during the early stages of the germination process.

In the present study, we have performed a transcriptome analysis of the early stages of conidal germination of *B. cinerea*. In order to get insight into the molecular events during the germination process as well as during early stages of the infection process, we have chosen optimized conditions for germination and subsequent appressorium formation on a hard, wax-coated surface. To obtain further evidence for genes that might be involved in the early infection process, we included the *bmp1 MAP kinase* mutant in the analysis.

**MATERIALS AND METHODS**

**Fungal strains and growth conditions.** The *Botrytis cinerea* laboratory strain B05.10 (wild type) and the MAP kinase deletion *bmp1* mutant were used for most experiments (27). For construction of green fluorescent protein (GFP) reporter strains, *B. cinerea* strain B05.Hyg-3 was used (34). For maintenance and spore production, *B. cinerea* was cultivated on tomato malt extract agar (TMA) as described previously (16). Conidia were harvested between 9 and 12 days after inoculation from sporulating plates using a glass spatula and 10 ml of distilled water and were washed with distilled water in 15-ml centrifugation tubes by three rounds of centrifugation at 3,500 rpm (27). Freshly harvested conidia (0 h) were used for RNA extraction or resuspended in Gamborg minimal medium (3 g/liter of Gamborg B5; 10 mM KH₂PO₄, pH 5.5) containing 10 mM fructose. Conditions for germination on glass, polyethylene, and polystyrene surfaces were as described previously (27). Conidia were rated as germinated when the spore wall was broken by the emerging germ tube.

**Analysis of pregermination conidial swelling.** For analysis of conidial swelling prior to germination, conidia were incubated in the presence or absence of 10 mM fructose, on apple wax-coated petri dishes, or on round glass coverslips (15-mm diameter; Roth, Karlsruhe, Germany). Time-lapse microscopic observations of individual conidia were made with an Axiosvert Observer A1 inverted microscope (Zeiss, Jena, Germany) equipped with a Neofluar 40×/1.3 oil immersion objective. Using a Canon Power-Shot G9 camera controlled by the program RemoteCapture, pictures were taken at 5-min intervals until germin tube outgrowth was visible. Based on the measured length and width of the spore, the changes in spore volume (ΔV) were calculated according to the formula $V = 4/3 \cdot \pi \cdot a^2 \cdot b^2/4$, where $a$ is length and $b$ is diameter of the spore. For each condition, at least three independent experiments ($n = 30$) were performed.

**Nuclear staining.** For staining of nuclei, the germination medium was removed, and the conidia covered with 50 μl of a fixing solution containing 3% formaldehyde and 10 mg/liter of Hoechst 33342, followed by incubation for 40 min at room temperature in the dark. When the germ tubes had developed for more than 1.5 h, they were fixed in the germination medium by heat treatment (60°C for 15 min) in a humid chamber, followed by 30 min of incubation in a solution containing 5 mg/liter of Hoechst 33342. Stained germ tubes were evaluated with a fluorescence microscope, and pictures were taken at different focal planes for subsequent counting of the nuclei. For each time point, a minimum of 50 spores was analyzed in three independent experiments.

**Preparation of coated surfaces for germination.** Apple wax was prepared by soaking ripe apples (cv. Golden Delicious) three times for 10 s in chloroform. The resulting solution was washed out sequentially with equal volumes of 0.1 N HCl, 0.1 N NaOH, and several changes of distilled water. The chloroform was removed under vacuum in a rotary evaporator at 30°C and dried under a stream of nitrogen gas. The wax was dissolved in hexane at a concentration of 2 g/liter and stored as stock solution at −20°C. Petri dishes (9-cm diameter) were coated by spreading the dissolved wax over the surfaces at a concentration of 10 μg/cm² and subsequently air dried. For preparation of cutin hydrolysate, peeled skins of 20 to 25 ripe apples (cv. Golden Delicious) were prepared, treated with pectinolytic (Frucoxym P; Erbslöh, Germany) and cellulolytic (Rohamet CT; Röhm Enzymtechnologie, Germany) enzymes, and hydrolyzed as described previously (35). The cutin hydrolysate was dissolved in hexane and used at 10 μg/cm² for surface coatings and induction tests. For coatings with trihydroxypalmitic acid (THPA; Sigma), 2-g/liter stock solutions in methanol were prepared.

For germination assays, conidia were inoculated in 30-μl droplets.
containing Gamborg B5 basal salt mixture with 10 mM fructose onto apple wax-coated polystyrene surfaces, if not indicated otherwise below, and incubated at 20°C in the dark.

**Analysis of GFP fluorescence.** For fluorescence microscopy, an Axio Observer A1 (Zeiss, Jena, Germany; Semrock GFP-A basic filter set; excitation wavelength, 469 nm; emission wavelength, 525 nm) was used. For quantification of fluorescence levels of cutB-GFP germlings, ImageJ 1.43u software (National Institutes of Health, Bethesda, MD) was used. Pictures were taken with the same exposure time for all treatments, taking care to avoid overexposure. Fluorescence of the germlings (marked as regions of interest) was quantified, and background levels were subtracted. The obtained values were classified as five levels of intensity, as follows: 0, <10; 1, 10 to 69; 2, 70 to 129; 3, 130 to 190; and 4, >190 (the units are pixel values from 0 [black] to 255 [white]).

**DNA manipulations.** The cdal-GFP and cutB-GFP promoter reporter strains were constructed by using a plasmid encoding a *B. cinerea* codon-optimized GFP (Bc-GFP-1) and a hygromycin resistance cassette for selection (36). For expression of Bc-gfp1 under the control of either cdal or cutB promoter, fragments covering 1,313 bp (cdal) and 1,281 bp (cutB) upstream of the start codons were amplified (primers are listed in Table S1 in the supplemental material) and used to replace the constitutive olcC promoter fragment. The plasmids were linearized with KpnI and used to transform *B. cinerea* B05.0H3-3 (34).

Construction of the rum1 mutant was performed by amplification of the entire coding region of *rum1* with an additional 365 bp of the promoter and 350 bp of the terminator region using the primers *rum1*-KO (see Table S1 in the supplemental material). The fragment was further cloned into the vector pLOBI (16) via Sacl and Apal restriction sites. Natural restriction sites for EcoRV (168 bp downstream of the start codon) and for EcoRI (157 bp upstream of the stop codon) in the coding region of *rum1* were used for replacing the coding region by a hygromycin resistance cassette. Genotypic verification of *rum1* deletion strains was performed by quantitative reverse transcription-PCR (qRT-PCR) analysis of *rum1* expression using the primers P104380fw/rev and PCR analysis of cutB-GFP promoter replacement. The plasmids were linearized with KpnI and used to transform *B. cinerea* B05.0H3-3 (34).

**RNA isolation and qRT-PCR.** For germination on 9-cm-diameter wax-coated petri dishes under standard conditions, 2 × 10⁶ conidia were suspended in 22.5 ml of Gamborg fructose medium and transferred into the dishes (9 cm). After incubation at 20°C in the dark for various times, the medium was poured off, and the germlings were removed from the dishes using a cell scraper. Germlings were centrifuged for 5 min at 4°C and 4,000 rpm, washed once with ice-cold water, and ground in liquid nitrogen to a fine powder. RNA was isolated with the RNeasy plant mini kit (Qiagen, Hilden, Germany).

For quantitative RT-PCR analysis, 1 μg of RNA from each sample was reverse transcribed into cDNA with oligo(dT) primers using a Verso cDNA kit (Thermo Fisher Scientific, Surrey, United Kingdom). PCR was performed using a MyIQ real-time PCR cycler (Bio-Rad, Munich, Germany). Expression of the genes was calculated by the 2-ΔΔCT method (37). For normalization of expression levels, several genes were tested for constant expression during germination (see Table S4 in the supplemental material). Based on the results, ubq1 and actA were subsequently used for normalization. Data are shown as normalized fold expression relative to the expression in ungerminated conidia. Means of three biological replicates, with two technical replicates each, are shown. The primers used in this study are listed in Table S1 in the supplemental material.

**Microarray expression analysis.** Ten micrograms of total RNA was converted into cDNA using a SuperScript II cDNA conversion kit (Invitrogen, Carlsbad, CA). Labeling of double-stranded cDNA with Cy3 or Cy5 namers and microarray hybridization were performed by Nimblegen (Roche NimbleGen, Inc.). For hybridization, Nimblegene 4-plex arrays, containing 4 × 72,000 arrays per slide, were used. Thirty 60-mer oligonucleotides per gene were immobilized for a total number of 20,885 predicted genes of the two sequenced *B. cinerea* strains B05.10 and T4, with 3,189 T4-specific genes and 1,776 B05.10-specific genes (designated BC1G_XXXX) (38). Genes annotated according to strain T4 (http://urgi.versailles.inra.fr/gb2/grayOURSE/BOTRTY7_T4_pub/) are referred to as in the abbreviated form PXXXXXX, instead of BofuT4_PXXXXXXX.1, throughout this article. Of these total genes, predicted genes encoding small coding sequences (<100 amino acids [aa]) and lacking expressed sequence tag (EST) support and any functional or topological domain were excluded, which reduced the number of genes evaluated in our study to 15,738. After hybridization of the arrays with the Cy3-labeled cDNA probes, they were scanned with a GenePix 4000B dual-laser scanner (Axon Instruments, Foster City, CA). Hybridization signals from all data sets, comprising three independent experiments, were normalized together, using the quantile function, and averaged for each gene (39). Thresholds of gene expression were determined by referring the hybridization signals to those of 9,557 random probes. Genes were considered expressed when their signal was ≥2 or higher, within a dynamic range of 2^2 to 2^10. This threshold was selected because for all time points, it was close to the 99th percentile of random probes hybridization signals (see Fig. S1 in the supplemental material). According to this classification, 12,091 (76.8%) of the supported genes were found to be expressed at one or more time points.

For analysis of time-dependent expression changes, genes showing an expression signal of at least 2^2 at one of the time points investigated were included in the analysis. Genes showing at least 2-fold, significant (P < 0.05, Student’s t test) changes in expression at a given time point were considered differentially expressed. Groups of coregulated genes with selected expression profiles across the five time points were generated using ArrayStar software, version 3.0.1 (DNASTAR Inc., Madison, WI). Genes lacking statistical support (P < 0.05) for the chosen expression profile were discarded.

**Gene annotation and categorization.** Functional annotation of *B. cinerea* proteins was performed as described previously (40), with the following modifications. For functional categorization, all expressed genes (12,092) were classified into the supported genes were found be expressed at one or more time points. For analysis of time-dependent expression changes, genes showing an expression signal of at least 2^2 at one of the time points investigated were included in the analysis. Genes showing at least 2-fold, significant (P < 0.05, Student’s t test) changes in expression at a given time point were considered differentially expressed. Groups of coregulated genes with selected expression profiles across the five time points were generated using ArrayStar software, version 3.0.1 (DNASTAR Inc., Madison, WI). Genes lacking statistical support (P < 0.05) for the chosen expression profile were discarded.

**Microarray data accession number.** The entire microarray data set described in this article is available at the Gene Expression Omnibus (GEO) database (http://www.ncbi.nlm.nih.gov/gds/?term = GSE43569) under accession number GSE43569.

**RESULTS**

**Pregeneration swelling, kinetics of germination, and nuclear division during germination.** Germination is a multistep process that starts with initiation of metabolism and the breaking of dormancy. Swelling, i.e., increase in spore volume before germination, is the first cytological evidence of activation of dormant spores in many fungi (42). We checked whether this also holds true for *B. cinerea*. When *B. cinerea* conidia were incubated in the presence of 10 mM fructose on glass slides, pregermination swelling was observed to start after about 1 h and continued until germ tube emergence. Swelling was dependent on active conidia, it did...
not occur in heat-killed conidia, and it was not observed in conidia incubated at high density, which inhibits germination. Surprisingly, no swelling was observed if conidia were incubated on a hydrophobic surface in water, in the absence of nutrients, although these conidia germinated (Fig. 1). Therefore, in B. cinerea conidia, pregermination swelling depends on the availability of nutrients during germination.

In order to achieve optimal germination conditions, similar to those on natural plant surfaces, we compared germination of B05.10 wild-type conidia on different artificial surfaces. For all experiments, conidia suspended in minimal medium containing 10 mM fructose were used. Compared to polystyrene and glass surfaces, surfaces generated by apple wax-coated petri dishes induced the most rapid germination of conidia. On wax surfaces, germination was completed already after 3 h, while on pure plastic and glass surfaces, complete germination was not observed until 4 to 5 h of incubation (Fig. 2A). For the transcriptome analyses shown below, we therefore used wax-coated surfaces. Compared to that of the wild type, germination of the MAP kinase bmp1 mutant on wax-coated surfaces was delayed by approximately 1 h (Fig. 2).

In conidia of N. crassa, A. nidulans, and F. graminearum, DNA replication and nuclear division have been found to start early during germination (43, 44, 45, 46). In B. cinerea, analysis of nuclear division is complicated by the fact that macroconidia are multinucleate. In resting conidia of B. cinerea strain B05.10, we observed between 2 and 7 nuclei (average, 3.7 ± 0.2). When germination was induced in 10 mM fructose on apple wax surfaces, a significant increase in the mean nuclear number was observed after 90 min. At this time, germination had started in approximately 20% of the conidia (Fig. 3). However, it took more than 4 h, when all conidia had germinated, until the average number of nuclei had approximately doubled. During germination, some of the nuclei migrated into the developing germ tube, but no regular pattern in this process was observed. These data indicated that the cell cycle and nuclear division start early during the germination process.

Transcriptome profiling of the germination process. In accordance with our previous biochemical studies and the cytological analyses, the following time points for isolation of RNA were used for microarray hybridization (Fig. 2B). As a reference, freshly harvested, nongerminated conidia (0 h) were used. After 1 h, degradation of trehalose had occurred (28), and conidia started to show swelling (Fig. 1). After 2.5 h, the majority of the conidia had germinated. After 4 h, most of the germ tubes had stopped elongation and formed terminal, appressorium-like thickenings. After 15 h, the germlings had resumed elongation and formed saprotrophic hyphae, with intermittent penetration attempts.

For each time point, microarray hybridizations were performed with three biological replicates of the wild-type strain, using Nimblegene whole-genome arrays representing gene mod-
els of the two sequenced *B. cinerea* strains B05.10 and T4 (40) (GEO database accession number GSE43569). The microarray expression data were first evaluated for stage-specific changes in global transcript abundance by performing pairwise comparisons between different time points (Fig. 4; see also Table S2 in the supplemental material). By far the largest changes were observed between resting conidia and conidia incubated for 1 h (0 h to 1 h), with a total of 2,864 genes being at least 2-fold up- or downregulated. Considerably fewer changes were observed between 1 h and later time points, when most morphological changes, including germ tube outgrowth and appressorium formation, occurred. Interestingly, gene expression changes between 1 h and 4 h were much higher than between 1 h and 2.5 h or between 2.5 h and 4 h. These data indicate that gene expression changes between the three time points largely occur in a continuous, gradual manner. In addition, they confirm the validity and comparability of the data for different time points.

**Correlation of gene expression profiles with gene function.** To identify genes with similar regulation during the germination process, groups of coregulated genes were generated according to the microarray data (Fig. 5; see also Table S3 in the supplemental material). The large majority of these genes (4,949 genes, or 40.1% of the expressed genes) was constitutively expressed (CON groups), showing less than 2-fold variation of expression during the five developmental stages examined. Subdivision of the CON genes into high (CON-H; expression signals of $>2^{14}$), medium (CON-M; $2^{11}$ to $2^{14}$), and low (CON-L; $2^{9}$ to $2^{11}$) expression revealed group CON-L to be the largest, followed by CON-M and CON-H. The next two largest groups, Max0 and Max1-15, comprised genes that showed either maximal or minimal expression in nongerminated conidia, followed by constant expression levels. The sixth largest group, Max15, comprised mycelium-specific genes. The seventh group, Max1-4, is described in more detail below. Four smaller groups are shown in Fig. 5, including one with genes that are upregulated only after 1 h (Max1), at the pregermination state.

The genes that were up- or downregulated between 0 h and 1 h (Fig. 4) and the genes belonging to groups shown in Fig. 5 (except for Max0+15), were analyzed for functional categories of their encoded proteins (Fig. 6; see also Table S3 in the supplemental material). Of the total of 12,092 expressed genes, 4,558 (37.7%) could be functionally categorized. Interestingly, the expression levels of constitutively expressed (CON) genes showed a clear correlation with the proportion of genes that could be assigned to functional categories: while the CON-L group contained only 31.8% genes that could be functionally categorized, the proportions were 47.8% for CON-M and 66.9% for CON-H. Remarkably, genes upregulated after 1 h could be functionally categorized to more than 60% (0-1_up), or even 70% (Max1, Max1-4, and Max1-15), which is clearly above the average of genes (37.7%).

Interesting differences of certain functional groups and their occurrence in specific expression groups were observed. In the CON-L group, an increased frequency of genes encoding secreted proteins was observed, whereas the frequency of genes belonging to cell cycle, transcription, protein synthesis, cellular transport, and other categories was significantly lower than expected. This pattern was largely inverted in groups CON-M and CON-H, which contained relatively few genes encoding secreted proteins but a strong enrichment of genes involved in cellular transport, energy and respiration (CON-H only), C-compound and carbohydrate metabolism, and protein synthesis and processing. Interestingly, all of the groups including genes induced during germination (i.e., genes activated after 1 h or 2.5 h) were enriched in genes encoding secreted proteins and carbohydrate-active enzymes (CAZymes). The only exception was the Max2.5-15 group, which did not contain any CAZyme genes.

**The bmp1 mutant shows downregulation of germination-related secreted proteins.** Germination of the bmp1 mutant conidia was somewhat retarded when compared to the wild type (Fig. 2). In addition, germ tubes continue to elongate on hydrophobic surfaces and never differentiate appressoria, indicating that they are defective in signaling leading to appressorium formation (Fig. 2B).
...upregulated in the wild-type germination-specific genes (32.6% on average) were not profile of the mutant germlings, we aimed to identify genes that are involved in germling development (Fig. 2). A remarkably high proportion of wild type after 2.5 h, representing stages with similar degrees of...bmp1 mutant after 3.5 h was compared to that of the mutant (Table 1; see also Table S3 in the supplemental material). Previously, bmp1 was found to be most stably expressed. We therefore used ubq1 and actA for normalization in the qRT-PCR analyses. In Fig. 7A to C and in Table S4 in the supplemental material, the time course of transcript levels is shown for 14 genes, which were selected because of their distinct, germination-induced expression profiles based on microarray data. For these genes, the microarray data were largely confirmed by qRT-PCR. Ten of these genes encode proteins with predicted signal peptides for secretion. Eight genes that showed weak expression in the bmp1 mutant after 3.5 h in the microarray experiment were confirmed to be downregulated in the bmp1 mutant at all time points analyzed (Fig. 7A).

**Expression analysis of cda1, a germination-specific gene.** cda1 was identified as the most rapidly and strongly induced gene during germination (Fig. 8; see also Table S4 in the supplemental material). The predicted Cda1 protein contains 641 amino acids, including a signal peptide, and shows 49% identity (61% similarity) to the chitin deacetylase of Colletotrichum lindemuthianum, CICDA, a member of the carbohydrate esterase 4 superfamily (47). Expression levels of cda1 in the wild type were maximal after 1 h and thereafter decreased to still high levels at 2.5 and 4 h and to much lower levels at 15 h. In contrast, cda1 expression was very low in the bmp1 mutant (Fig. 8; see also Table S4). To monitor the induction of cda1 in vivo, a cda1-GFP promoter fusion was constructed and transformed into B. cinerea strain B05.Hyg-3 (34). Expression of GFP in conidia was strictly correlated with germination (Fig. 8A). Starting from weak background levels, a strong increase in fluorescence occurred before germ tube outgrowth in all spores (Fig. 8B). Induction of cda1 was independent of the germination conditions and occurred in a similar manner on hard surfaces and in suspensions, in either minimal or full media (data not shown).

**A cutinase-encoding gene (cutB) requires plant lipids for germination-specific expression.** Based on microarray expression data, several genes encoding putative cutinases were found to be induced during germination (Fig. 9; see also Table S4 in the supplemental material; other data not shown). Previously, cutA has been described to be induced early after germination on the host plant surface (48). Based on the microarray data, cutA showed maximal expression between 1 h and 2.5 h, while expression of another cutinase gene, called cutB, showed a peak of expression between 2.5 h and 4 h (Fig. 7; see also Table S4). cutB encodes a 206-amino-acid protein with 34% identity to Fusarium...
solani Cut1 and containing a predicted 18-amo-ino-acid leader peptide. Expression of cutB was analyzed using a B. cinerea cutB-GFP promoter reporter strain. GFP fluorescence was very weak in conidia germinated on artificial surfaces such as polypropylene (Fig. 9B). In contrast, induction of cutB was observed on surfaces covered with plant lipids. While moderate fluorescence appeared in gernmlings on glass surfaces coated with apple wax, strong fluorescence could be observed if the surfaces were coated with cutin hydrolysate or the cutin monomer THPA (Fig. 9B). Interestingly, individual conidia responded differently to the inducing surfaces. While some spores showed bright fluorescence after 3.5 h, other spores in a similar developmental stage remained weakly fluorescing (Fig. 9A and B; other data not shown). This behavior was independent of the concentrations of inducing plant lipids applied to the glass slides (data not shown), indicating that it was due to an intrinsic cell-to-cell variability of the cutB promoter. In the presence of sugars, cutB expression was reduced, but not abolished, by carbon catabolite repression. Fructose was observed to have a weaker suppressive effect than that of glucose, and at higher concentrations of the inducer THPA, repression by the sugars was less effective (Fig. 9B). To verify the physiological relevance of cutB induction, cutB-GFP conidia were germinated on heat-killed onion epidermal cell layers. Many gernmlings showed bright GFP fluorescence covering the spore, the germ tube, and the infection hyphae, indicating that cutB is similarly induced on the host cuticle (Fig. 9C). Again, cutB expression was not uniformly distributed, because some of the gernmlings failed to induce the GFP reporter, although penetration occurred.

A putative secondary metabolite gene cluster showing coordinate regulation during germination. Among the genes showing peak expression at 4 h postinooculation (hpi), we identified eight adjacent genes that could be involved in the synthesis of secondary metabolites (see Fig. S2 in the supplemental material). The putative cluster contained genes encoding a fatty acid elongase, two cytochrome 450 monooxygenases, a transferase, a thioesterase, a small secreted protein of unknown function, a major facilitator superfamily (MFS) transporter, and a transcription factor of the Zn2Cys6 zinc cluster family. When the expression of these genes was reexamined by qRT-PCR, we found three genes on one side of the cluster to be rather stably expressed during germination. In contrast, four of the five other genes, except for the transcription factor-encoding gene (called rum1, for regulator of unknown metabolite 1), were strongly downregulated after 1 h and 2.5 h, followed by a steep increase after 4 h. In contrast to the microarray data, qRT-PCR did not reveal a clear decrease in expression of these genes between 4 and 15 h (data not shown). In order to clarify the role of rum1 in the regulation of the gene cluster, the gene was deleted in B. cinerea strain B05.10. In the rum1 mutant, expression of most the cluster genes was moderately reduced in conidia (0 h) compared to that in the wild type, while a strong downregulation was observed for the four adjacent genes only (see Fig. S2). These data narrow down the gene cluster to five genes, including rum1. Rum1 seems to be required for maintaining high levels of expression of the cluster genes. A preliminary phenotypic analysis of the rum1 mutant did not reveal any differences concerning germination, appressorium formation, or pathogenicity compared to the B05.10 wild-type strain (data not shown). The role of the hypothetical secondary metabolite cluster therefore remains unclear.

DISCUSSION

A transcriptome analysis of the germination process of Botrytis cinerea conidia has been performed under conditions that mimic the situation on the host plant cuticle. On apple wax-coated surfaces and a minimal medium containing fructose, rapid and rather synchronous germination of conidia within 3 h was achieved. The majority of germinated conidia stopped germ tube elongation and formed appressorium-like terminal thickenings within 4 to 5 h, indicating the initiation of penetration. Germination on apple wax was more rapid than on artificial hydrophobic surfaces. The germination-accelerating activity of apple wax might be due to an additional chemical stimulus. Chemical components of host cuticles have been found to affect germination and differentiation of several plant-pathogenic fungi (7, 13, 14, 15). In the powdery mildew Blumeria graminis f. sp. hordei, conidial germination and differentiation were found to be triggered by components of the leaf surface wax, in particular long-chain aldehydes (11).

### TABLE 1 Expression of genes encoding secreted proteins in the wild type versus the bmp1 mutant

<table>
<thead>
<tr>
<th>Category</th>
<th>Total</th>
<th>No. of genes encoding secreted proteins (% of total wild type)</th>
<th>Total no. of bmp1 low genes&lt;sup&gt;a&lt;/sup&gt; (% of total wild type)</th>
<th>No. of bmp1 low genes&lt;sup&gt;b&lt;/sup&gt; encoding secreted proteins (% of wild type)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Max1-2.5</td>
<td>13</td>
<td>5 (38.5)</td>
<td>7 (53.8)</td>
<td>4 (80.0)</td>
</tr>
<tr>
<td>Max2.5 + Max2.5-4</td>
<td>5</td>
<td>2 (40.0)</td>
<td>3 (60.0)</td>
<td>2 (100.0)</td>
</tr>
<tr>
<td>Max1-4</td>
<td>117</td>
<td>21 (17.9)</td>
<td>34 (29.1)</td>
<td>14 (66.7)&lt;sup&gt;***&lt;/sup&gt;</td>
</tr>
<tr>
<td>Total germination specific&lt;sup&gt;c&lt;/sup&gt;</td>
<td>135</td>
<td>28 (20.7)</td>
<td>44 (32.6)</td>
<td>20 (71.4)&lt;sup&gt;***&lt;/sup&gt;</td>
</tr>
<tr>
<td>Max1-15</td>
<td>471</td>
<td>108 (22.9)</td>
<td>102 (21.7)</td>
<td>43 (39.8)&lt;sup&gt;***&lt;/sup&gt;</td>
</tr>
<tr>
<td>Max2.5-15</td>
<td>64</td>
<td>17 (26.6)</td>
<td>39 (60.9)</td>
<td>14 (82.4)</td>
</tr>
<tr>
<td>Total germination induced&lt;sup&gt;c&lt;/sup&gt;</td>
<td>670</td>
<td>153 (22.8)</td>
<td>185 (27.6)</td>
<td>77 (50.3)&lt;sup&gt;***&lt;/sup&gt;</td>
</tr>
<tr>
<td>Max0</td>
<td>552</td>
<td>57 (10.3)</td>
<td>27 (4.9)</td>
<td>0 (0.0)</td>
</tr>
<tr>
<td>Max15</td>
<td>118</td>
<td>20 (16.9)</td>
<td>12 (10.2)</td>
<td>2 (10.0)</td>
</tr>
<tr>
<td>CON</td>
<td>4,949</td>
<td>361 (7.3)</td>
<td>199 (4.0)</td>
<td>19 (3.3)</td>
</tr>
</tbody>
</table>

<sup>a</sup> Genes showing at least 2-fold-lower expression (<i>P < 0.05</i>) in the bmp1 mutant (3.5 h) compared to the wild type (2.5 h). Significant differences (<i>P < 0.001</i>; Student’s <i>t</i> test) between observed and expected frequencies are indicated (**<sup>*></sup>**).

<sup>b</sup> Comprising the Max1-2.5, Max2.5, Max2.5-4, and Max1-4 groups.

<sup>c</sup> Comprising all germination-specific groups and groups Max1-15 and Max2.5-15.
In many fungi, spore germination has been observed to be preceded by swelling (2). Using video microscopy, we observed that pregermination swelling of *B. cinerea* conidia occurred only in the presence of carbohydrate nutrients, while in the absence of nutrients, hydrophobic-surface-induced germination was not accompanied by swelling of the conidia (Fig. 1). This indicates a profound difference in the developmental programs leading to germination either in the presence or in the absence of exogenous nutrients. The major difference in signaling is the predominant role of cAMP for carbon source-induced germination and the essential role of the BMP1 MAP kinase cascade for hydrophobicity-induced germination (27, 28, 30). Carbon source-induced germination, but not hydrophobicity-induced germination in the absence of nutrients, is accompanied by trehalose degradation (28). We therefore speculate that the stronger mobilization of nutrient reserves might be responsible for conidial swelling. In our study,
Germination was accompanied by an increase in the number of nuclei (Fig. 3). However, we did not observe a synchronization of germination or appressorium formation and nuclear division, as has been reported for *M. oryzae* or *C. gloeosporioides* (49, 50, 51). Previously, synchronous division of individual nuclei in germinating *B. cinerea* conidia has been observed, using a *B. cinerea* strain with GFP-expressing nuclei (33). An explanation for this discrepancy from our result is difficult, but it might be related to the different surfaces used, namely, untreated glass slides in the work of Kokkelink et al. (33) and apple wax–coated surfaces in our study, resulting in germination kinetics that differed by more than 1 h (Fig. 2).

For the transcriptome studies, resting conidia (0 h) and conidia incubated for 1, 2.5, 4, and 15 h on apple wax–coated surfaces in the presence of 10 mM fructose were used. Before RNA isolation, conidia were washed, and the germings were scraped off the plates and washed once in ice-cold water. While we cannot exclude that these procedures led to changes in the mRNA composition, the differential expression data obtained were found to be quite robust and did not seem to be due to artifacts. The Nimblegen array represented 15,738 gene models of the two sequenced *B. cinerea* strains, T4 and B0510, that were supported by the size of the open reading frames, the presence of functional or topological domains, or expression data. After normalization and threshold subtraction, 12,092 (76.8%) genes were classified as being expressed during at least one of the five time points analyzed. However, this number is only a rough estimation, because it was difficult to define the threshold that distinguished expression signals from background noise. The threshold of >2^1 was selected because it was beyond the 99th percentile values of the random probes for all time points except one. Extensive changes of expression, equaling 23.7% of the expressed genes, were observed during the first hour of incubation, prior to germ tube outgrowth. In contrast, during the time period when most of the morphological changes occurred, between 1 h and 4 h, much lower changes of gene expression were observed (Fig. 4). In *F. graminearum*, pregermination changes in gene expression (from 0 to 2 h) have been reported to be even larger, with 3,421 (66.5%) of 5,146 expressed genes, also exceeding gene expression changes during later development (19). In *M. oryzae* conidia, 2,154 (approximately 21%) of all genes were found to be up- or downregulated after complete germination under non-appressorium-inducing conditions after 7 h (22). Although these data are not completely comparable to each other, they confirm data in the literature on a rapid mRNA turnover in *F. graminearum* (42). In *B. cinerea* and other fungi, the first biochemical evidence for the breaking of dormancy and initiation of the germination program is the breakdown of trehalose, detectable after 30 min and almost completed after 1 h in the presence of fructose on a glass surface (28). It remains to be shown when transcriptional changes start to occur after perception of the germination stimuli. From the expressed genes, 11 major groups of
coregulated genes were extracted. As expected, the majority of the genes (4,949, or 40.9% of the expressed genes) belonged to the three constitutive groups (CON-L, CON-M, and CON-H) (Fig. 5). Furthermore, many genes showed preferential expression in the dormant stage (Max0; 552 genes) and in the mycelium stage (Max15; 471 genes). As already indicated by the large changes observed between 0 and 1 h, a considerable number of germination-induced genes with specific expression patterns (including Max1, Max1-4, Max2.5-15, and Max1-15; all together, 692) were identified. Overall, the gene expression patterns determined by the microarrays were confirmed by qRT-PCR analysis, although some quantitative differences were observed between the two methods (Fig. 7; see also Table S4 in the supplemental material).

The groups revealed interesting differences in the distribution of functional categories. By far the lowest proportion of genes which could be assigned to functional categories (31.8%) were found in the CON-L group. This illustrates a particular lack of knowledge about the function of fungal genes that are weakly expressed during all growth stages. In contrast, the high proportion of genes with functional evidence (66.9%) in the CON-H group was correlated with a prevalence of well-known genes involved in major functions of primary metabolism, including cell cycle and DNA processing, transcription, protein synthesis and processing, and respiration. Consistent with the analysis in N. crassa and F. graminearum, functional categories of cell cycle, DNA processing, and protein synthesis were also found to be overrepresented in B. cinerea conidia among the genes induced between 0 and 1 h (17, 19). Genes induced 1 h after germination could be functionally categorized to more than 60% (0-1_up), or even more than 70% (Max1, Max1-4, and Max1-15). Furthermore, all of these categories were enriched in genes encoding secreted proteins (20.7% of the total number of expressed genes), whereas a lower frequency of genes encoding secreted proteins were observed to be expressed in dormant spores (10.3%). This peak of secretory activity during early germination has not been reported for N. crassa or Fusarium oxysporum (17, 19). In M. oryzae, genes encoding secreted proteins have been found to be induced mainly during appressorium formation. Some of these proteins, such as the cutinase Cut2, the hydrophobin MPG1, and the SnodprotI homolog MSPI, have been shown to be involved in virulence (22).

A B. cinerea early secretome, derived from conidia germinated for 16 h in a minimal medium enriched with low-molecular-weight host plant compounds, has been published recently (52). Of the 105 secreted proteins identified, 14 are encoded by the germination-specific and 21 by the germination-induced genes identified in our study. In addition, 35 of the proteins corresponded to constitutively expressed genes. Overall, these data document a significant overlap of the proteomic and our transcriptomic data. Five proteins in the secretome have been described as virulence factors, namely, the endopolygalacturonases PG1 and PG2 (53, 54), the pectin methyltransferase PME1 (55, 56), the endo-β-1,4-xylanase Xyn11A (57), and the cerato-platanin family protein SPL1 (58). They were found in the Max1-15 (PG1, PG2, and PME1), Max1 (Xyn11A), and Max15 (SPL1) groups. Eleven of the secreted proteins, including PG1 and PG2, were encoded by genes that are downregulated in the bmp1 mutant, and 9 of them belonged to the germination-induced gene groups. The bmp1 mutant also showed a clear downregulation of the PG3-encoding gene (P094200) which is expressed Max1-15 in the wild type. While PG3 has not been described as a virulence factor, previous expression data revealed a massive induction of PG3 in infected apples (59).

In M. oryzae, 357 genes (22), or even 1,026 genes (23), have been found to be differentially expressed during appressorium initiation or maturation. This large number can be explained by the fact that M. oryzae appressoria are morphologically highly distinct and specialized structures. In contrast, B. cinerea appressoria are morphologically much less differentiated and usually not separated from the germ tube by a septum. Comparison of the genes induced during germination and appressorium formation between B. cinerea and M. oryzae did not reveal clear similarities (23). For example, of the 31 genes involved in acetyl coenzyme A (acetyl-CoA) utilization or production, 13 were found in M. oryzae to be upregulated and six downregulated during appressorium formation. In contrast, only three of these genes were germination-specifically expressed in B. cinerea, and only one of them was an ortholog of an appressorium-specific gene in M. oryzae. Similarly, 17 out of 22 genes involved in fatty acid oxidation and glyoxylate cycle were specifically induced in appressoria of M. oryzae, but none of their orthologs were induced during B. cinerea germination. This suggests that appressorium formation in M. oryzae differs from that in B. cinerea significantly also at the molecular level. It also indicates that fatty acid mobilization and the glyoxylate cycle, which are essential for providing the high concentrations of glycerol in the M. oryzae appressoria that are required for high turgor pressure-mediated penetration, do not play a similarly important role in B. cinerea. Nevertheless, a common feature of both fungi was the high frequency of genes encoding secreted proteins, including CAZymes, which were found to be upregulated both in M. oryzae appressoria and in B. cinerea germings.

Comparison of gene expression in germlings of the wild type and bmp1 mutant revealed a high proportion of genes that were induced during germination in the wild type and that appeared to be dependent for their induction on the presence of the MAP kinase BMP1 (Table 1). Even more strikingly, 50.3% of the germination-induced genes encoding secreted proteins were downregulated in the bmp1 mutant. In contrast, only 5.3% of CON genes encoding secreted proteins were BMP1 regulated. However, these data are based on microarray analysis of only a single time point (3.5 h) for the bmp1 mutant. Nevertheless, for eight genes, dependence on BMP1 was confirmed by a qRT-PCR time course analysis with the wild-type and the bmp1 mutant (Fig. 7A; see also Table S2 in the supplemental material). These results indicate a significantly lower induction of secretory activity during germination in the bmp1 mutant. In M. oryzae, the BMP1-orthologous MAP kinase PMK1 is also essential for appressorium differentiation and infection. Transcriptome studies have revealed a total of 481 genes that are PMK1 regulated. Of these genes, 238 and 174 are induced during early and late appressorium development, respectively (23). However, only 13 orthologs of these genes were found to be downregulated in the B. cinerea bmp1 mutant, and only 5 of these were upregulated during germination and appressorium differentiation. In the M. oryzae genome, a total of 206 genes encode CAZymes. Seventy-two (35%) of these genes were significantly upregulated in appressoria, but only 9 (12.5%) of these genes were downregulated in the bmp1 mutant. Of the 316 genes encoding CAZymes in B. cinerea, 48 (15%) were upregulated during germination, and 30 (62.5%) of these were downregulated in the bmp1
mutant. Overall, this comparison revealed a rather small overlap between the two fungi of the genes that are upregulated during early stages of infection, and differences in their regulation by the BMP1/PMK1 MAP kinase pathway. The most rapidly and strongly induced gene during germination, cda1, showed more than 700-fold increased expression within the first hour of spore incubation. Analysis of a cda1-gfp promoter reporter strain confirmed that the appearance of GFP fluorescence was strictly correlated with germination, reaching strong levels before germ tube appearance. Germination-specific expression of cda1 was found to be independent of the germination conditions and strongly reduced in the bmp1 mutant. cda1 is a member of a gene family encoding secreted proteins similar to chitin deacetylases and chitin binding proteins in other fungi (47). In the rust fungus Uromyces fabae, differentiation-specific induction of chitin deacetylase activity has been speculated to be involved in protecting the fungal hyphae from degradation by plant chitinases (60). Functional analysis of cda1 is difficult, because it is a member of a family of three cda genes, including cda1, cda2 (P079220/BC1G_03291), and cda3 (P005177/BC1G_06509), which possibly have redundant functions. While cda2 shows a rather stable expression during development, both cda1 and cda3 showed maximum expression at 1 to 4 h (see Table S2 in the supplemental material; other data not shown).

Because germination was induced on apple wax-coated surfaces, cutinase genes that require the presence of host plant lipids were observed in the microarrays among the germination-specific genes. cutA has previously been shown to be induced in B. cinerea mycelium in the presence of cutin monomers and repressed by glucose (48, 61, 62). We found that cutA and cutB were induced during germination with different kinetics (Fig. 7). With a cutB promoter reporter strain, GFP fluorescence could be observed in germinating spores only on surfaces covered with plant lipids such as apple wax or cutin monomers. Similar to cutA, cutB was subject to catabolite repression. The degree of repression by glucose was stronger than that by fructose and was reduced with increasing concentrations of the inducing lipid (THPA) (Fig. 9B). In contrast to the highly synchronized expression observed for cda1, expression of cutB was found to be variable in different germings. The delayed induction of cutB in some germings, sometimes even after penetration, indicates that perception of the inducing plant lipids occurs with variable efficiency and kinetics, which is in striking contrast to the strictly controlled induction of cda1 during early germination. In addition to cutA and cutB, the B. cinerea genome encodes several more cutinase genes which show different patterns of expression (reference 48 and unpublished data). The redundancy of cutinases makes it difficult to assess the role of their activity for host cell wall penetration by the analysis of mutants (61).

A limitation of our approach to search for genes that are induced during germination is the fact that it does not identify genes with different expression that might be crucial for the germination process, including regulatory genes. Nevertheless, we identified a BMP1-regulated transcription factor gene (P088480; BC1G_06861) with germination-induced expression (Max2.5-15) (see Table S2 in the supplemental material). Targeted knockout mutagenesis, however, did not reveal any obvious phenotype concerning germination, penetration, or infection (data not shown). Another candidate for functional analysis was the zinc cluster transcription factor rum1, which is part of a co-regulated cluster of five genes that appears to be involved in secondary metabolite synthesis (see Fig. S2). Next to this cluster, three more genes are located that might perform a similar function but were only weakly regulated by Rum1. B. cinerea produces two types of secondary metabolites which have been reported to be virulence factors, namely, sesquiterpenoids, including botrydial, and polyketides, including botricin acid (63, 64). Although the rum1 deletion mutant showed a strongly reduced expression of the cluster genes, it did not reveal any defects concerning germination, penetration, or infectivity. A metabolite profiling of the rum1 mutant might provide an idea about the role of this gene cluster.

In summary, our data demonstrate that early conidial development in B. cinerea is accompanied by rapid shifts in gene expression that likely orchestrate germ tube outgrowth, appressorium differentiation, and host cell invasion. For some of the germination-induced genes, a role in infection has already been demonstrated, while for the other genes, this remains to be analyzed. Besides their functional characterization, it will be a challenge to achieve a more comprehensive description of the regulatory processes that are involved in germination and early infection of B. cinerea.

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