Differential Gene Expression During Conidiation in the Grape Powdery Mildew Pathogen, *Erysiphe necator*

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ABSTRACT

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Asexual sporulation (conidiation) is coordinately regulated in the grape powdery mildew pathogen *Erysiphe necator* but nothing is known about its genetic regulation. We hypothesized that genes required for conidiation in other fungi would be upregulated at conidiophore initiation or full conidiation (relative to pre-conidiation vegetative growth and development of mature ascocarps), and that the obligate biotrophic lifestyle of *E. necator* would necessitate some novel gene regulation. cDNA amplified fragment length polymorphism analysis with 45 selective primer combinations produced ≈1,600 transcript-derived fragments (TDFs), of which 620 (39%) showed differential expression. TDF sequences were annotated using BLAST analysis of GenBank and of a reference transcriptome for *E. necator* developed by 454-FLX pyrosequencing of a normalized cDNA library. One-fourth of the differentially

expressed, annotated sequences had similarity to fungal genes of unknown function. The remaining genes had annotated function in metabolism, signaling, transcription, transport, and protein fate. As expected, a portion of orthologs known in other fungi to be involved in developmental regulation was upregulated immediately prior to or during conidiation; particularly noteworthy were several genes associated with the light-dependent VeA regulatory system, G-protein signaling (Pth11 and a kelch repeat), and nuclear transport (importin-β and Ran). This work represents the first investigation into differential gene expression during morphogenesis in *E. necator* and identifies candidate genes and hypotheses for characterization in powdery mildews. Our results indicate that, although control of conidiation in powdery mildews may share some basic elements with established systems, there are significant points of divergence as well, perhaps due, in part, to the obligate biotrophic lifestyle of powdery mildews.

Additional keywords: expression analysis, *Uncinula*.

In powdery mildews, rapid production of asexual conidia enables dispersal and expansion of population size, while sexual ascocarps are required for survival of abiotic stresses and of host dormancy in the absence of living green tissues. For *Erysiphe necator* (syn. *Uncinula necator*), the causal agent of grapevine powdery mildew, the developmental cues for conidiation are dependent on a number of factors, including inoculum density, light, temperature, and humidity (12,15,32,41). In spite of the gradient in age across a colony, *E. necator* conidiation occurs synchronously at a defined and predictable time, suggesting the presence of a signal to coordinate conidiation (41). Further evidence of the coordinate regulation of conidiation in *E. necator* occurs after pairing of opposite mating types and ascocarp formation, when conidial production ceases (20).

To date, however, no work has been done to uncover the genetic basis for developmental signals or how these signals might control powdery mildew development in general. In contrast, significant progress has been accomplished with several other fungi (13,19,21,22), including a major effort in the model fungus *Aspergillus nidulans* to elucidate regulatory genes and signaling pathways for developmental control (10,18,21,44). Functional analyses have uncovered roles for several transcription factors, including the C2H2 transcription factors *brlA*, *FlbC*, and *SteA* (2,27,41), a single Myb transcription factor *FlbD* (42), and *abaA* and *wetA*, which, along with *brlA*, form a central regulatory pathway for conidiation (1). These and other pathway regulators activate 1,200 conidiation-specific genes (38). For example, *abaA* coordinates regulation of a number of structural genes, including the chitin synthases *chsA* and *chsC*; the Δ *chsA* Δ *chsC* double mutant mimics the aberrant conidiophore morphology of mutants defective in *abaA* or *brlA* (24). In powdery mildews, many of these transcription factors and structural genes could be involved in the developmental commitment to conidiation.

A significant barrier to expression analyses in powdery mildews has been the paucity of publicly available gene sequences, with the notable exception of *Blumeria graminis*. Outside of the phylogenetically distinct tribe of powdery mildews that contains only *B. graminis* (7), only 23 expressed sequence tags have been deposited at GenBank for the Erysiphales to date (28). This lack of expression data is partially due to the inability to maintain the fungus in the absence of the host. However, recent advances have enabled the isolation of high-quality *E. necator* RNA without contaminating host RNA (9). Still, without sequence data, approaches

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to gene expression analysis are limited to open-platform technologies that discover gene sequences. The cDNA amplified fragment length polymorphism (AFLP) technique allows the simultaneous comparison of expression of uncharacterized genes across multiple treatments or developmental stages, the discovery of new gene sequences, and higher stringency than other differential-display methods, thus reducing some potential for false positives (4).

The goal of the present research was to identify gene sequences differentially expressed during vegetative growth and throughout asexual and sexual reproduction in *E. necator*. Previously, we have used microsurgery to restart the *E. necator* developmental clock by excising the colony center precisely before conidiation (20). By sequentially excising the centers of colonies that were (i) at a presporulation stage or (ii) sporulation competent, we were able to define precisely when a purported sporulation signal might be propagated within the colony. With this detailed knowledge of the timing of signal initiation for conidiation, we were able here to study expression precisely before conidiation signaling (preconidiation), afterward at conidiophore initiation and full conidiation, and during ascocarp development when conidiation is again halted. We hypothesized that homologs of genes required for conidiation in other fungi would be upregulated at conidiophore initiation or full conidiation, and that the obligate biotrophic lifestyle and synchronous conidiation of *E. necator* would necessitate some novel gene regulation.

MATERIALS AND METHODS

Plant material, inoculum, and pathogen growth. Seedlings were grown from seed collected from susceptible open-pollinated *Vitis vinifera* vines according to Ficke et al. (17). The seedlings were grown in constant 23°C with a 12-h photoperiod until the five- to six-leaf stage under mildew-free conditions. Two isolates of *E. necator*, 10-18 and 10-36, of opposite mating type were collected from New York vineyards and maintained on seedlings under the same conditions. For growth of the asexual stages, conidia were harvested from colonies ≈8 days postinoculation (dpi) by shaking infected leaves in distilled water containing 0.005% Tween 20. The suspensions were adjusted to a concentration of 40,000/ml by counting with a hemacytometer, and inoculated as described below. For colonies where sexual reproduction was desired, conidial suspensions of both isolates were prepared as above, mixed in equal proportion, and vortexed to create one suspension of both mating types.

Colonies for RNA extraction were prepared by placing two dozen 5-μl drops of the conidial suspension (200 conidia/drop) across the surface of the youngest fully expanded leaf of a grape seedling. The suspension was allowed to dry and then the seedlings were placed in a mildew-free growth chamber at 23°C with a 12-h photoperiod. When the colonies reached the desired stage, all inoculated leaves were observed under a dissecting microscope for uniformity of development. Mildew colonies showing at least 90% uniformity in terms of relative radial growth of colonies and appearance of the appropriate conidiation stage

(i.e., for colonies in the conidiophore development stage, >90% of colonies on selected leaves showed conidiophore emergence but no fully developed conidia) were selected for RNA extraction.

RNA isolation. RNA was extracted from powdery mildew colonies at each of four stages: preconidiation (3 dpi), conidiophore initiation (≈5 dpi), full conidiation (8 dpi), and development of mature ascocarps (≈4 weeks post inoculation). All RNA extractions were performed according to Cadle-Davidson et al. (9). For the three asexual stages, two independent isolations were performed from each of two isolates. For the sexual stage, four independent isolations were performed of the same pair of isolates. Within each stage, equal amounts of total RNA were pooled from the four independent extractions to form one pool per stage of interest. Polyadenylated mRNA was isolated from total RNA using the PolyATtract system (Promega Corp., Madison, WI). The mRNA was not eluted from the beads in the final step, and the beads were used directly in the iScript cDNA Synthesis Kit (Bio-Rad, Hercules, CA) to convert the mRNA into cDNA.

cDNA-AFLP analysis. The cDNA-AFLP protocol was adapted from Bachem et al. (4), using the same adapters and primers described therein. cDNA from each stage was digested with *TaqI* followed by *AseI* digestion. The digested fragments were then ligated to *TaqI* and *AseI* adapters prior to preamplification with *TaqI*+0 and *AseI*+0 primers. Amplification conditions were as follows: 4 min of denaturation at 94°C and then 27 cycles of 30 s of denaturation at 94°C, 30 s of annealing at 55°C, and 60 s of extension at 72°C. After preamplification, products were diluted to 1 ng/μl (approximately fivefold dilution).

Diluted preamplification product (5 μl) was used as template for selective amplifications. Selective amplification was carried out with 45 primer combinations using two selective nucleotides for each primer (e.g., *TaqI*+2 and *AseI*+2) for the selective nucleotide combinations listed in Table 1. The polymerase chain reaction (PCR) amplification program was as follows: denaturation at 94°C for 4 min; then 13 cycles of denaturation at 94°C for 30 s, annealing at 65°C for 30 s (reduced 0.7°C per cycle), and extension at 72°C for 60 s; followed by 32 cycles of 30 s of denaturation at 94°C, 30 s of annealing at 56°C, and 60 s of extension at 72°C.

Selective amplification products were separated by electrophoresis on a 5% polyacrylamide gel in a Bio-Rad Protean II xi Gel System at 40 W and 45°C for 3.5 h. Following separation, fragments were visualized by silver staining according to the protocol by Echt et al. (16). Fragments of interest were picked from the gel using a sterile pipette tip, transferred to 100 μl of distilled water in a microcentrifuge tube, and heated to 95°C for 10 min. This eluted DNA (1 μl) was used as a template for reamplification using the preamplification primers under the following conditions: 4 min of denaturation at 94°C; followed by 8 cycles of 30 s of denaturation at 94°C, 30 s of annealing at 61°C (reduced by 0.7°C per cycle), and 60 s of extension at 72°C; followed by 24 cycles of 30 s of denaturation at 94°C, 30 s of annealing at 56°C, and 60 s of extension at 72°C. PCR samples (15 μl) were electrophoresed in a 1% agarose gel and visualized using UV excitation of SYBR Green in the loading dye. PCR products with visible, single bands were purified using Sephadex columns and Sanger sequenced directly using each preamplification primer at Cornell University's Core Laboratories Center (CLC).

Normalized cDNA library and 454 pyrosequencing. *E. necator* isolate G14 was collected from a no-spray plot of the *Vitis* interspecific hybrid Rosette in an experimental vineyard in Geneva, NY, in September 2007. RNA was isolated from cultures of G14 according to Cadle-Davidson et al. (9) at 17 dpi and provided to Bio S&T (Montreal, CA) for preparation of an uncloned normalized cDNA library for 454 pyrosequencing and a cloned library for quality assessment. A sodium chloride precipitation was completed at Bio S&T prior to library construction due

TABLE 1. Forty-five selective nucleotide primer combinations used in the cDNA amplified fragment length polymorphism analysis of differential expression across four developmental time points in *Erysiphe necator*

TaqI+2	AseI+2
CA	GA, GG, GT
CC	AG, GA, GT
CG	GA, GG, GT
CT	GA, GG, GT
GA	AG, GA, GC, GG, GT, TG
GC	AA, AG, CC, GA, GC, GG, GT, TG
GG	AA, AG, GA, GC, GG, GT, TG
GT	AG, CC, CT, GA, GC, GG, GT, TA
TG	GA, GC, GG, GT

to a low 260/230 ratio. Purified cDNAs were digested after normalization with *Sfi*I for directional cloning. After purification, the digested cDNA fragments were ligated into pBluescriptII SK(–) with *Sfi*I A&B inserts between *Eco*RI and *Xho*I. A portion of the ligation products was transformed into DH10B-T1R (Invitrogen, Carlsbad, CA). Ten colonies were arbitrarily selected from the cloned library and insert sizes were determined by PCR using M13 primers.

Pyrosequencing by a 454-FLX (454 Life Sciences, Branford, CT) was completed at Cornell CLC. After in silico removal of adaptor sequences from the raw sequencing reads for the 5' end (5'-CAGTGGTATCAACGCAGAGTGGCCATTACGGCCTAGTTACGGG-3' and the 3' end (5'-[PolyA]GGCCGCCTCGGC CACTCTGCGTTGATACCACTG-3'), MIRA software (14) was used to assemble the reads into contigs de novo.

Sequence analysis. Similarity searching by blastx and tblastx was carried out against the National Center for Biotechnology Information (NCBI) nonredundant (nr) BLAST database. To increase the probability of finding a match, longer sequences corresponding to the amplified cDNA-AFLP products were found by performing a nucleotide BLAST (blastn) of each sequence against the normalized cDNA database described above. cDNA contigs matched in this search were then searched as six-frame translations against the translated NCBI nr database using tblastx.

Real-time PCR confirmation. To confirm differential expression of the identified sequences, quantitative reverse-transcription (qRT)-PCR was performed using a MyiQ thermal cycler (Bio-Rad), 1.5 μ M Syto 9 for product quantification (Invitrogen), and 10 nM fluorescein (Sigma-Aldrich, St. Louis) for well factor normalization. For preliminary assessment of the isolate-independent differential expression of genes, colonies were grown from two isolates of *E. necator*, Fcon2 and G9. Fcon2 was collected in September 2007 from an abandoned Concord vineyard in Fredonia, NY. G9 was collected in June 2007 from the same no-spray Rosette vineyard as G14. RNA was isolated from colonies of Fcon2 at precondiation, conidiophore initiation, and full conidiation and from paired Fcon2 and G9 at ascocarp initiation. These stages were identical to those used within the cDNA-AFLP experiments, with the exception of the sexually reproducing stage, wherein RNA was isolated from colonies at ascocarp initiation rather than maturation. For the qRT-PCR experiments, only one RNA isolation was performed per stage of interest as opposed to the four used in the cDNA-AFLP analysis. Extraction of the RNA, purification of mRNA, and synthesis of cDNA were performed as described above. Each sample was then normalized to equalize cDNA concentration. Following purification, the cDNA samples and remaining mRNA were analyzed with *E. necator* primers to confirm the absence of *E. necator* genomic DNA.

After analyzing the sequence data from the cDNA-AFLP experiment, 17 sequences with a match in the BLAST searches were arbitrarily selected. Primers for real-time PCR were designed for all 17 sequences and tested for quantitative response to gDNA and specificity by melt curve analysis. Following this initial analysis, seven sequences were selected for further analysis based on a linear response of threshold cycle (C_t) to log-transformed starting quantity and a single sharp peak at the amplicon's melting temperature in the dissociation, or melt curve, analysis. In addition to normalization based on cDNA concentration, an amplification control, histone H3, previously shown to be evenly expressed throughout *B. graminis* f. sp. *hordei* development (6), was included in all qRT-PCR experiments. Three technical qRT-PCR replicates were performed for all samples. The PCR amplification program was as follows for all qRT-PCR experiments: 10 min of denaturation at 95°C; followed by 45 cycles of 15 s of denaturation at 95°C, 45 s of annealing at 58°C, and 60 s of extension at 72°C; this was followed by a melt curve analysis ramping from 55 to 95°C in 15-s increments of

0.4°C. Normalized relative quantity (NRQ) values were calculated as described elsewhere (34) to compare a single cDNA-AFLP upregulated developmental stage relative with background expression in the other stages, normalized by expression of the control gene histone H3 (6). NRQ values >1 reflect higher expression at the expected developmental stage.

RESULTS

cDNA-AFLP sequences. In total, 45 primer combinations were used for cDNA-AFLP analysis, each resulting in 21 to 64 transcript-derived fragments (TDFs) in a size range of $\approx$ 100 to 800 bp (Fig. 1). Across all 45 primer combinations, $\approx$ 1,600 TDFs

Fig. 1. Representation of differential expression of *Erysiphe necator* transcripts across four developmental stages for three cDNA amplified fragment length polymorphism primer combinations. Transcript profiles are shown for three *Ase*I+2 primers (GA, GC, and GG) with the AG *Taq*I+2 primer. cDNA pools within each gel from left to right are column 1, precondiation; column 2, conidiophore initiation; column 3, full conidiation; and column 4, ascocarp maturation. These three gels show a molecular weight range of $\approx$ 100 to 800 bp.

were visualized. Within these, 620 TDFs (39%) were polymorphic among the four stages, suggesting differential expression. Of these polymorphic TDFs, 230 were directly sequenced (Supplemental Table 1). The remaining 390 fragments (out of the 620 TDFs) either gave poor product upon reamplification or gave a degraded product when sequenced, likely due to co-migration of different sequences. When searched against the GenBank database using tblastx, 45 of the 230 sequences (20%) had a significant match (E value ≤ 1 E-03) (Table 2). Only one sequence matched the *V. vinifera* genome, indicating that the extraction method was successful in targeting *E. necator* RNA.

Pyrosequence matches. To improve the ability to find matches, a reference transcriptome for *E. necator* was developed by 454-FLX pyrosequencing of a normalized cDNA library of isolate G14. The average insert size of this library was 2.3 kb (Fig. 2). However, after one sequencing run, insufficient coverage resulted in an average contig size of only 591 bp (Table 3). The TDFs were searched against this *E. necator* cDNA sequence database using blastn. Of the 230 sequences searched against the library, 153 (67%) showed significant matches to cDNA contigs. The 153 matching contigs were searched against the GenBank database, again using tblastx, and 116 (76%) had a significant

TABLE 2. Annotation of select transcript-derived fragments (TDFs) identified by cDNA amplified fragment length polymorphism as differentially regulated during *Erysiphe necator* development

Primers ^a	Fragment ID ^b	Stage ^c	E value cDNA ^d	Annotation ^e	E value GenBank ^f
Metabolism					
CA-GA	9-13-28	3	6 E-27	Lysine synthesis/Lys2 (XP_002143711)	9 E-132
CC-GA	11-13-11	1	3 E-92	Spermine/spermidine synthase (EEH21650)	1 E-136
CC-GA	11-13-20	4	4 E-44	Acetyl CoA synthetase FacA (XP_001935668)	1 E-52
GA-GG	13-16-17	1	5 E-37	Acetolactase synthase (XP_962652)	0
GA-GG	13-16-18	1-3	5 E-34	Pap2 domain protein (XP_001931173)	1 E-72
TG-GT	8-14-12	1-3	7 E-29	Cytochrome P450 (XP_755288)	1 E-60
GT-GG	14-16-19	1-3	4 E-34	Vacuolar ATP synthase subunit B (XP_001557622)	0
GC-AG	15-4-5	1, 4	1 E-47	D-3 phosphoglycerate dehydrogenase (XP_0022382929)	4 E-162
GC-GT	15-14-33	1-2	3 E-25	Chitin synthase (XP_956331)	0
GG-AG	16-4-7	2-4	1 E-66	Protoheme IX farnesyl transferase (EER45220)	3 E-72
GG-GC	16-15-11	3-4	1 E-47	Alcohol dehydrogenase (EEH49979)	8 E-47
TG-GT	8-14-11	1-3	6 E-54	Actin polymerization protein/Bzz1 (XP_749256)	5 E-87
GG-GG	16-16-21	1-3	3 E-23	Fatty aldehyde dehydrogenase (XP_001932922)	3 E-165
GG-AG	16-4-3	1, 4	4 E-40	Caleosin domain protein (XP_001932418)	2 E-53
TG-GA	8-13-22	1	2 E-39	Elongation factor 3 (XP_001593480)	0
GT-GT	14-14-15	4	2 E-56	γ Tubulin (XP_001903940)	6 E-76
Transcription					
CG-GA	12-13-18	1	1 E-43	C2H2 zinc finger domain (XP_002482718)	6 E-32
GT-GA	14-13-9	1-2	4 E-85	C2H2 zinc finger domain (XP_002382564)	0
GT-AG	14-4-18	4	5 E-27	C2H2 transcription factor RfeC (XP_002481027)	4 E-25
GT-GT	14-14-11	2-4	7 E-65	Gal4-like transcriptional activator (AAG25917)	4 E-117
GC-GT	15-14-31	2	5 E-6	Transcriptional activator Spt7 (EEH04734)	3 E-51
GG-GG	16-16-7	2-4	3 E-38	Transcriptional elongation factor Spt6 (EEH34974)	0
GG-AG	16-4-2	1, 4	3 E-56	FlbD (EED33126)	1 E-4
GG-TG	16-8-9	2-3	4 E-32	Ring finger domain protein (XP_00125767)	7 E-130
GC-GA	15-13-10	1-3	2 E-41	CobW domain protein (XP_002384953)	3 E-170
GT-TT	14-6-8	2-4	5 E-47	Mads box transcription factor (XP_001935687)	7 E-46
GC-GG	15-16-10	1-3	6 E-51	Histone transcriptional regulator (CAE85596)	2 E-34
CA-GA	9-13-17	4	3 E-9	PHD and ring finger domain protein (EER42091)	1 E-43
Signaling					
TG-GA	8-13-19	1-2	4 E-50	Calcium permease (XP_001547054)	1 E-73
GA-GA	13-13-3	2	1 E-179	Integral membrane protein/Pth11 homolog (AAD30437)	6 E-64
CA-GA	9-13-24	4	2 E-24	Catalase peroxidase (Q8X1N3)	0
CA-GT	9-14-35	3	3 E-38	Cystolic regulator Pianissimo (XP_001938513)	0
GA-GT	13-14-14	4	1 E-59	Casein kinase I (XP_001272602)	0
CG-GA	12-13-13	1-3	9 E-39	Serine/threonine protein kinase (XP_001548642)	1 E-171
CA-GA	9-13-27	2	2 E-24	Kelch repeat protein (EEE22934)	3 E-34
Transport					
TG-GG	8-16-11	2-3	4 E-63	RAN homolog (XM_001594666)	6 E-114
GA-GA	13-13-12	2-3	3 E-35	Coatomer subunit alpha (AAC18088)	2 E-116
GT-GT	14-14-13	4	2 E-38	SAC3-GANP domain protein (EEH38661)	3 E-77
GC-GA	15-14-14	1-2	1 E-17	Importin beta-3 subunit (XP_001264986)	0
TG-GT	8-14-22	2-4	3 E-51	Stomatin family protein (XM_001594877)	2E-149
GT-GT	14-14-12	3-4	1 E-40	Integral membrane protein (XP_0022376767)	8 E-66
GT-TG	16-8-8	1, 4	3 E-64	PRELI/MSF1 domain protein (EER25643)	2 E-57
GT-TA	14-5-1	1	*	MFS monocarboxylic acid transporters (EDP52060)	5 E-07
Protein					
TG-GA	8-13-21	3	7 E-46	ER-associated proteolytic system protein DER1 (XP_747611)	3 E-77
TG-GT	8-14-21	2	9 E-39	Heat shock protein 70 (EU311400)	0
GA-GG	13-16-9	3	2 E-58	F Box domain protein (XP_001272021)	2 E-105

^a Primer combinations are given by selective nucleotides in the following 5'-3' format: AseI primer-TaqI primer. Transcription = transcription factors/DNA binding, Transport = transport/cell membrane proteins, and Protein = protein maintenance and degradation.

^b Each TDF fragment was given a number based on the primers used (first two numbers) and the fragment number within each primer combination.

^c Stages of expression are represented as follows: 1 = preconditioning, 2 = conidiophore initiation, 3 = full conidiation, and 4 = development of mature ascocarps.

^d TDFs with no match to the *Erysiphe necator* cDNA pyrosequences are scored with a * in the blastn search against the *E. necator* cDNA library. The E value given is for the highest match against the cDNA library.

^e Tblastx was conducted for each cDNA contig sequence against the GenBank database, and the annotation of the gene with highest similarity is given here. Where no contig was matched, the annotation is given on the direct search of the original TDF against the GenBank database.

^f E values of the gene with highest similarity in tblastx are listed for the direct search of the cDNA library contig or TDF sequence against the GenBank database.

match. Nine of the TDFs without matches in the *E. necator* transcriptome contigs directly matched the GenBank database, for a total of 125 TDFs (54%) with sequence similarity high enough by this approach to consider them putative homologs.

Functional classification and confirmation. Five functional classes of differentially expressed fragments were identified: metabolism, transcription, signaling, transport, and protein fate. Based on relevance of the match annotation to previously described conidiation-associated genes, select matches to the GenBank database are listed in Table 2 (a complete list of sequences is available in Supplemental Table 1). In all, 105 of the 230 sequences (46%) had no significant match at GenBank by a query of either the TDF or the sequence with highest similarity in the *E. necator* cDNA database. An additional 33 TDFs (14%) matched hypothetical fungal protein sequences of unknown function. Among the sequences that matched those of predicted or known function, the largest class, representing 13% of differential transcripts, was sequences involved in metabolism. qRT-PCR expression analysis was used to test whether differential expression of arbitrarily chosen, annotated genes was dependent on the isolate used. Expression results using RNA from isolate Fcon2 and a pairing of Fcon2 with isolate G9 confirmed expected stages of upregulation based on results for isolates 10-18 and 10-36 from the cDNA-AFLP analysis for seven of nine developmental stages (Table 4).

DISCUSSION

The cDNA-AFLP method of analyzing gene expression across developmental stages has been described as useful for comparisons across multiple treatments in non-model species (31,35) and particularly appropriate in studies of obligate biotrophs, where tissue and sequence information are limiting (40). Our results with *E. necator* demonstrated the utility of cDNA-AFLP in morphogenesis studies in this pathosystem. Using 45 primer combinations, we were able to visualize ~620 differentially expressed fragments out of 1,600 total fragments. Although this level of differential expression is high for this type of study, the result is not surprising, considering the many changes of the fungus from purely vegetative growth to asexual and ultimately sexual reproduction in this study. In addition, developmental events related to host infection occur at the same time.

Several sequences fulfilled expectations of differential expression. Analysis of those sequences that showed similarity to previously described sequences revealed several patterns in gene expression, described by gene category below, with each TDF identified as ("AFLP fragment ID": "stage"), with stage defined as 1 = preconidiation, 2 = conidiophore initiation, 3 = full conidiation, and 4 = ascocarp.

Metabolism. The largest class of identified genes was involved in cell metabolism. A number of the metabolism genes are expressed early on in the vegetative and asexual stages of growth but are switched off in late stages of infection (Table 2), which is not surprising because the fungus transitions from biotrophic colonization and conidiation to ascocarp maturation and dormancy. Structural proteins such as chitin synthase play key roles in conidiation (24), and at least one chitin synthase (15-14-33: 1,2) is upregulated in *E. necator* prior to full conidiation.

Several sequences were similar to genes involved in lipid metabolism, including links to molecules like oxylipins that have demonstrated roles in coordinating fungal development. One TDF showed a relatively strong match to a Pap2 domain protein (13-16-18; 1,2,3) linked to nutrient stress responses in yeast (22) and to the generation and degradation of lipid-signaling molecules (11). Sporulation in response to nutrient stress is common in filamentous fungi, and the Pap2 domain proteins may be responsible for the coordination of a lipid signal that stimulates this response.

Caleosins are strongly associated with the maintenance of storage lipid bodies in plant seed and fungal spores, and individual members are involved in oxylipin pathways (30). The caleosin domain sequence (16-4-3: 1,4) in *E. necator* may have a role early in infection to maintain lipid bodies upon which germinating spores rely until successful infection occurs, and again during sexual reproduction to produce storage bodies for the developing ascospores. Because conidiation is an energy-intensive process, a developmental program that halts conidiation to preserve host resources would be a distinct advantage.

Signaling. The greatest similarities between *E. necator* sequences and the model systems were noticed in signaling. In other fungi, heterotrimeric G-protein signaling has a well-established role in coordinating multiple responses, including conidiation (44). We have uncovered several elements of G-protein signaling that indicate that the same basic elements may provide the basis for developmental control in *E. necator*.

One TDF (13-13-3: 2) showed moderate similarity to homologs of Pth11. Pth11 has been predicted to encode a G-protein-coupled

TABLE 3. Results of 454-FLX pyrosequencing and *de novo* assembly of an *Erysiphe necator* normalized cDNA library

Sequencing results	Values
Total sequence	82 Mb
Reads placed in contigs	99%
Total contigs	32,405
Contigs <500 bp	18,088
Average contig length	591 bp
Range of contig length	0.05–5.8 kb
Average reads per contig	9.1
Range of reads per contig	2–800

Fig. 2. Insert sizes for the cloned, normalized cDNA library of *Erysiphe necator* used for transcriptome sequencing. Ten colonies (clones 1 to 10) were arbitrarily selected, and insert sizes were determined by agarose electrophoresis of polymerase chain reaction amplicons obtained by M13 primers. Average insert size was calculated to be 2.3 kb. M = 1-kb marker (Invitrogen, Carlsbad, CA).

receptor (GPCR) in *Magnaporthe oryzae* and may play a role in pathogenesis-related morphogenesis (29). Its specific expression at the transient stage of conidiophore initiation makes this gene an interesting candidate as a key signaling receptor immediately prior to conidiation.

In *Aspergillus* spp., the primary Gα subunit that coordinates asexual development is FadA (36). Peeters et al. (33) demonstrated in yeast that two kelch repeat proteins, Krl1p and Krl2p, can bypass the need for cAMP and directly link Gα subunits with protein kinase A. A kelch-repeat sequence in *E. necator* (9-13-27: 2) (confirmed by qRT-PCR) was expressed coincident with the putative GPCR sequence, thus suggesting that, indeed, it may have some role in transmitting signals received through the GPCR described above. In *Dictyostelium* spp., Pianissimo (9-14-35: 3) (confirmed by qRT-PCR) is required for the G-protein-linked cAMP signaling.

Transcriptional regulators. The third-largest group of discovered genes consisted of those involved in transcriptional regulation and DNA binding, with upregulation most often at the conidiophore initiation stage (Table 2), indicating large changes in gene regulation at this time point. In *Aspergillus* spp., *flbD* encodes a constitutively expressed myb-like DNA-binding protein necessary for early-stage conidiation events (42). A weak match in *E. necator* (16-4-2: 1,4) was not expressed when expected and likely possesses a different function in *E. necator*. Several C2H2 transcription factors have known roles in conidiation (e.g., *brlA* and *FlbC*) or cleistothecial development (e.g., *SteA*) (2,27,43). We identified three sequences with similarity to C2H2 transcription factors (14-13-9: 1,2, [confirmed by qRT-PCR] 12-13-8: 1, and 14-4-18: 4), none of which was expressed during conidiation. A more detailed phylogenetic analysis of the matching contigs may provide interesting insight into the conserved or novel roles of these transcription factors.

Cell membrane or transport. In *Aspergillus* spp., the velvet protein (VeA) complex plays a critical role in controlling asexual versus sexual development, acting in the nucleus to repress asexual development in the dark, and being excluded from the nucleus in light (5). Araújo-Bazán et al. (3) recently demonstrated that importin-α interacts with VeA, and importin-α mutants show impaired nuclear transport of VeA. We have found a sequence showing similarity to the importin-β subunit (15-13-14: 1,2), which joins importin-α in a heterodimer required for proper function.

Related to the function of importin is Ran, a small GTPase. The importin heterodimer releases its contents in the nucleus upon disassociation of the heterodimer, stimulated by the interaction of the β-subunit with Ran (3). A sequence showing similarity to Ran

(8-16-11: 2,3) was expressed slightly later than the importin-β subunit. Studies in *Exserohilum* spp. also showed upregulation of Ran during conidiation and not in earlier stages (19). Nuclear transport may be impaired until initiation of conidiation, with the Ran–importin interaction important for import of complexes that trigger rather than repress conidiation.

Protein maintenance or degradation. Studies in *Neurospora* spp. have indicated that genes involved in protein degradation are upregulated during asexual sporulation (25). The Cop9 signalosome, which has functions in targeted protein degradation, coordinates development in *Aspergillus* spp. (8) and plays a role in regulation of PpoA expression (39). F-box proteins, which act as substrate receptors for ubiquitin ligase, have also been proven necessary for proper cell division and sexual spore development in *Aspergillus* spp. (26).

Among the sequences related to protein degradation, two were specifically upregulated at full conidiation: an F box domain protein (13-16-9: 3) and a proteolytic system protein (8-13-21: 3), though upregulation of the former was not independently confirmed in isolate Fcon2 (Table 4). Proteolytic systems could be necessary for turnover of protein complexes responsible for the suppression of asexual development. Although no turnover points have yet been discovered, VeA contains PEST domains for rapid turnover, indicating that degradation of VeA is important for development control (10).

Novel sequences in conidiation. Several sequences identified here have not previously been found to be expressed during conidiation in other filamentous fungi and could provide novel inroads to our understanding of conidiation and of *E. necator* biology. This expectation is exemplified by the most highly upregulated gene in this study, a PRELI/MSF1 domain-containing sequence (16-8-8: 1,4) which is likely involved in intramitochondrial protein sorting (23) but has not previously been implicated in conidiation. Specialization of gene sequences and genetic regulation involved in conidiation could be expected in part due to the obligate biotrophic lifestyle of powdery mildews. Not only are host factors likely to affect fungal development but also the genomes of powdery mildews have undergone massive gene loss and gene shuffling as a result of extreme parasitism (37). In the process, regulatory pathways and structural proteins are likely to have been disrupted and new genes appropriated for the task. This will require functional analysis in powdery mildews to determine gene functions that are out of context in autotrophic fungi.

Technical considerations. Outside of the phylogenetically distinct tribe of powdery mildew fungi that contains only *B. graminis* (7), almost no sequence data have been deposited at

TABLE 4. Quantitative reverse-transcription polymerase chain reaction (qRT-PCR) analysis for seven selected genes across four time points in *Erysiphe necator* isolate Fcon2

Fragment ID ^a	Expected expression ^b	NRQ ^c	Annotation	Forward primer ^d	Reverse primer ^d
n/a	Normalizer gene	1.0	Histone H3	TAACACGCTTGGCATGGATA	TGGAACAGTTGCTCTTCGTG
14-13-9	Preconidiation	1.3	C2H2 zinc finger transcription factor	CAACGCCGAGATACTAAAT	AGCGCCAATTGATAAGACTG
	Conidiophore initiation	1.1	...		
16-8-8	Preconidiation	11.0	PRELI/MSF1 domain protein	TGCTCAAGTTTCTCCCTCAC	CTGTTGCGATCGGTAAGTCT
	Ascocarp initiation	140.6	...		
9-13-27	Conidiophore initiation	37.8	Kelch repeat protein	TCAGTCTCCTCCAAGTTTCG	CCTCAACGTAGGCTTTACGA
11-13-20	Ascocarp initiation	1.9	Acetyl CoA synthetase FacA	GTGATGCGGCAAGTAGAGAT	TCGAGAGATCCAAAGACTGG
9-14-35	Full conidiation	2.1	Pianissimo	CAGCTGCTTGTTACCCAGTT	CGTCCAAGAAACCAATCATC
13-16-9	Full conidiation	0.1	F-box domain protein	GAATCGTATGTCGCATTTC	GTCCAGATTGTCCCACTGAG
9-13-24	Ascocarp initiation	0.3	Catalase peroxidase	TTCTTGGGGAGGTGAGACAA	GCGACCAAAAGTAACTCGAA

^a Transcript-derived fragment: the first two numbers correspond to the selective primers used, and the third is the fragment number within the primer combination (Table 3 and supplemental Table 1); n/a = not applicable.

^b Developmental stage of maximum expected expression was based on fragment abundance in the cDNA amplified fragment length polymorphism (AFLP) analysis of isolates 10-18 and 10-36. For qRT-PCR, Fcon2 was paired with G9 for ascocarp initiation as an imperfect confirmation of ascocarp maturation from cDNA-AFLP analysis of isolates 10-18 and 10-36.

^c NRQ = normalized relative quantity (34), calculated as the ratio of the change in candidate gene expression to the change in the control gene Histone H3 (6), at the listed developmental stages. NRQ values >1 (highlighted in bold) reflect higher expression at the expected developmental stage.

^d Primer sequences used for qRT-PCR. All primers are shown 5'–3'.

GenBank for the Erysiphales (28). This situation, combined with the short sequences we obtained after amplification and sequencing of TDFs, resulted in only 20% of TDFs having putative homologs in GenBank. Therefore, we pyrosequenced a normalized cDNA library during a growth stage that had initiated secondary conidial infections. Although our sequencing depth provided contigs (average 591 bp) that were smaller than the estimated cDNA insert size (2.3 kb) (Table 3; Fig. 2), the cDNA library sequence nearly tripled the number of TDFs that could be functionally annotated. Still, cDNA-AFLP has the limitation of only tracking those transcripts with the restriction sites and selective nucleotides (here, 45 of 256 possible) used and only identifying previously sequenced genes. As a result, we were able to annotate only a fraction of the genes expressed in association with conidiation. Developing the resources for transcriptome expression profiling (e.g., high-throughput sequencing) would enable the discovery of all differentially regulated genes.

For preliminary assessment of the isolate-independent differential expression of genes, colonies were grown from a third isolate of *E. necator*, Fcon2, and a pairing of Fcon2 with G9 for qRT-PCR analysis of candidate genes. We decided to confirm results with additional isolates rather than with isolates 10-18 and 10-36 to estimate how many genes might have conserved roles in morphogenesis, rather than simply confirm the technical and biological reproducibility of cDNA-AFLP and qRT-PCR in this experimental system. For seven of nine of the stages for which increased expression was expected based on cDNA-AFLP, results confirmed expectations (Table 4), suggesting that many genes identified here may be differentially expressed through development independent of the isolate used. Due to technical barriers to obtaining fully mature cleistothecia from the Fcon2-G9 pairing, we replaced the mature ascocarp stage with ascocarp initiation, which may be a different enough developmental stage to have significant transcriptional differences compared with the former. When expression was expected to be highest at full conidiation, high levels of expression were also seen at ascocarp initiation. Gadoury et al. (20) demonstrated that ascocarp formation was temporally associated with cessation of conidial production. Because RNA was extracted at ascocarp initiation for qRT-PCR as opposed to fully mature ascocarps in the cDNA-AFLP work, conidiation had not yet completely shut off and some overlap between expression at full conidiation and this stage was to be expected.

Conclusions. This work represents the first investigation into differential gene expression during morphogenesis in *E. necator* and identifies candidate genes and hypotheses for further characterization in powdery mildews. Our results reveal that there may be some similarities in control of development between *E. necator* and other fungi. Particularly noteworthy were several genes with similarities to those associated with the light-dependent VeA regulatory system, G-protein signaling, transcriptional regulation, and nuclear transport in *A. nidulans*. The largest groups of sequences, however, were those with no match to known sequences (45%) and unknown function (14%). This indicates that, although control of conidiation in *E. necator* may share some basic elements with established systems, there are likely significant points of divergence as well, due in part to the extreme parasitic nature of obligate biotrophs. Through this intimate connection with the host plant, powdery mildews may rely on the host for developmental cues in ways autotrophic fungi like *Aspergillus* and *Neurospora* spp. do not. Characterization of these novel mechanisms for regulating fungal development will require functional genetic analysis to fully characterize conidiation in obligate biotrophic fungi.

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