

# Characterization of a Silent Azaphilone Gene Cluster from *Aspergillus niger* ATCC 1015 Reveals a Hydroxylation-Mediated Pyran-Ring Formation

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## SUMMARY

Azaphilones are a class of fungal metabolites characterized by a highly oxygenated pyrano-quinone bicyclic core and exhibiting a broad range of bioactivities. Although widespread among various fungi, their biosynthesis has not been thoroughly elucidated. By activation of a silent (*aza*) gene cluster in *Aspergillus niger* ATCC 1015, we discovered six azaphilone compounds, azanigerones A–F (1, 3–7). Transcriptional analysis and deletion of a key polyketide synthase (PKS) gene further confirmed the involvement of the *aza* gene cluster. The biosynthetic pathway was shown to involve the convergent actions of a highly reducing PKS and a non-reducing PKS. Most significantly, *in vitro* reaction of a key flavin-dependent monooxygenase encoded in the cluster with an early benzaldehyde intermediate revealed its roles in hydroxylation and pyran-ring formation to afford the characteristic bicyclic core shared by azaphilones.

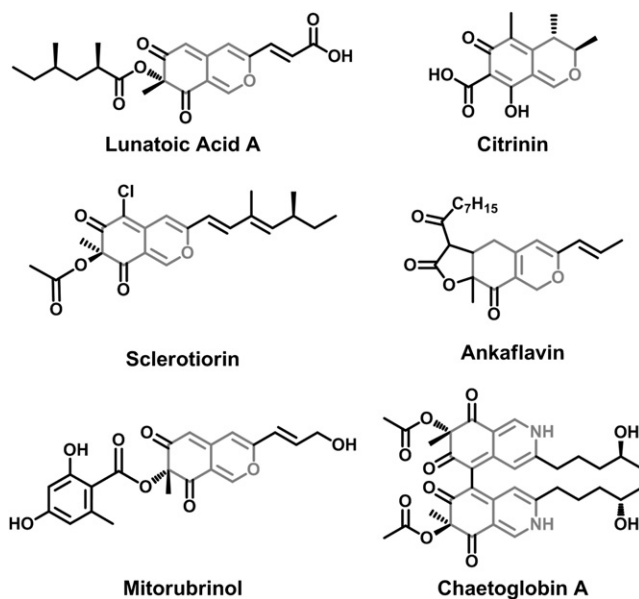
## INTRODUCTION

Filamentous fungi are known to be prolific producers of secondary metabolites, such as penicillin, lovastatin, and cyclosporine, and are an important resource for discovering small molecules of pharmaceutical and industrial value (Keller et al., 2005). In the last decade, whole-genome sequencing of various fungi has revealed that these microorganisms have immense biosynthetic potential that far surpasses the chemical diversity that we observe in laboratory culture (Sanchez et al., 2012). For example, the genome of many aspergilli are found to encode for a combined 30 to 80 polyketide synthases (PKSs), nonribosomal peptide synthetases (NRPSs), and PKS-NRPS hybrids, which far exceeds the total number of known polyketides and nonribosomal peptides (Sanchez et al., 2012). Of these, the fungal PKSs are of considerable interest due to their interesting enzymology and the polyketide structural diversity.

Fungal type I PKSs contain multiple catalytic domains and resemble the animal fatty acid synthases, where a single set of

catalytic domains is used iteratively. The chain extension by decarboxylative condensation of malonyl-CoA units is catalyzed by the minimal PKS domains, including ketosynthase (KS), malonyl-CoA:ACP transacylase (AT), and acyl carrier protein (ACP) (Cox, 2007). Non-reducing PKSs (NR-PKSs) synthesize a poly- $\beta$ -ketone backbone that is cyclized by a product template (PT) domain to yield aromatic compounds such as orsellinic acid and norsolorinic acid (Crawford et al., 2009). In contrast, highly reducing PKSs (HR-PKSs) utilize different combinations of ketoreductase (KR), dehydratase (DH), and enoyl reductase (ER) domains following each chain extension to reduce the  $\beta$ -keto positions to different extents and produce reduced polyketides such as lovastatin and fumonisins (Cox, 2007). Together with tailoring enzymes that are typically clustered in a biosynthetic pathway at the genetic level, the different fungal PKSs produce a large array of polyketides (Keller et al., 2005).

Bioinformatic analyses of different fungal genomes have revealed that it is common for two PKSs to be located in the same gene cluster (Sanchez et al., 2012). The polyketide products of several of these dual PKS-containing gene clusters are known, including hypothemycin (Reeves et al., 2008; Zhou et al., 2010), asperfuranone (Chiang et al., 2009), and lovastatin (Kennedy et al., 1999; Ma et al., 2009). The two PKSs can either work in sequence or in convergence to synthesize the polyketide product. When the two PKSs function sequentially, the polyketide chain formed by the first PKS is transferred to the second PKS to continue the chain extension process. This has been demonstrated in the biosynthesis of the resorcylic acid lactones and asperfuranone, in which the upstream HR-PKS produces a partially reduced polyketide chain that is transferred to the downstream NR-PKS to be further elongated (Chiang et al., 2009; Zhou et al., 2010). In the convergent model, the two PKSs can function independently in parallel, and the two polyketide products are ultimately connected via accessory enzymes. An example is the biosynthesis of the lovastatin, in which the nonaketide and a diketide chains produced by two different HR-PKSs are combined via the action of the acyltransferase LovD (Xie et al., 2009). With a limited number of dual PKS systems characterized so far, it is currently not possible to predict which mode of crosstalk (sequential or convergent) between the two PKSs will take place through bioinformatic means alone. Therefore, characterization of additional dual PKS-containing pathways will facilitate our understanding of the molecular and genetic basis that underlies the differences between



**Figure 1. Examples of Azaphilone Compounds Isolated from Fungi**  
The conserved bicyclic core (highlighted in gray) is the hallmark of these compounds.

the PKS-PKS partnerships and enable better prediction of the gene cluster products.

*Aspergillus niger* and closely related black aspergilli are known to produce a large number of secondary metabolites, with up to 145 compounds cataloged (Nielsen et al., 2009). Annotation of the sequenced *A. niger* genomes unveiled an impressive number of PKS genes, including 34 PKSs and 7 PKS-NRPS hybrids (Andersen et al., 2011; Pel et al., 2007). In this work, we mined the genome of *A. niger* ATCC 1015 and identified a polyketide gene cluster that contains a pair of HR-PKS and NR-PKS. Overexpression of a pathway-specific transcriptional regulator found in the gene cluster led to the overproduction of a number of previously unknown azaphilone natural products. Azaphilones possess a signature bicyclic chromophore and are known to react readily with amines to produce the vinylogous  $\gamma$ -pyridones (Osmanova et al., 2010). This family is structurally diverse and includes bioactive compounds such as the antifungal lunatoic acid A (Nukina and Marumo, 1977), the anti-inflammatory agent monascorubrin (Yasukawa et al., 1994), the lipoxygenase inhibitor sclerotiorin (Chidananda and Sattur, 2007), and the food dye ankaflavin (Manchand et al., 1973), as well as the nephrotoxic mycotoxin citrinin (Endo and Kuroda, 1976; Sakai et al., 2008) (Figure 1). We show here that the two partnering PKSs function in a convergent manner toward azaphilone biosynthesis.

## RESULTS

### Bioinformatic Analysis of *A. niger* ATCC 1015 Genome Reveals a Gene Cluster with Two PKS Genes

We scanned the sequenced genome of *A. niger* ATCC 1015 for dual PKS gene clusters and found a single gene cluster containing an HR-PKS and an NR-PKS. The NR-PKS gene

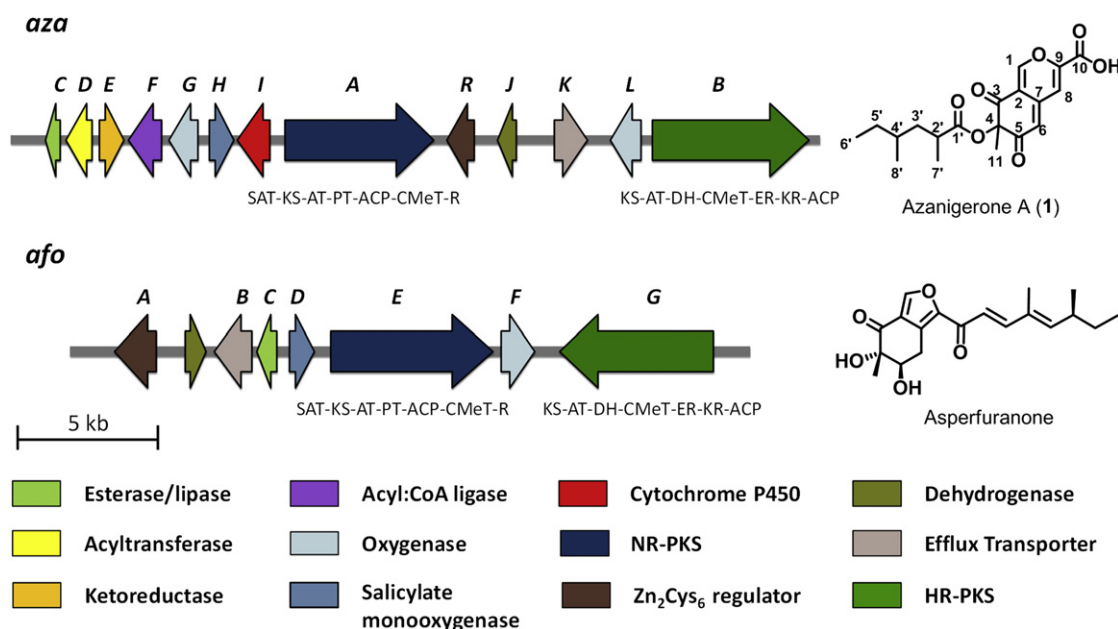
ASPNDRAFT\_56946 and the HR-PKS gene ASPNDRAFT\_188817 (herein referred to as *azaA* and *azaB*, respectively) appear to be homologs of *afoE* and *afoG* (44% and 43% protein identity) in the *A. nidulans* *afo* cluster, which is responsible for the synthesis of asperfuranone (Chiang et al., 2009) (Figure 2; Table 1). The *azaA* gene encodes for a Clade III NR-PKS according to a previous fungal PKS phylogenetic classification scheme (Kroken et al., 2003). In addition to the minimal PKS domains, it also includes an N terminus starter unit: acyl-CoA transacylase (SAT) and a C terminus didomain of C-methyltransferase (CMeT) and reductase (R). On the other hand, *AzaB* is a typical HR-PKS consisting of the minimal PKS domains, the  $\beta$ -keto processing domains (DH, ER, and KR) and CMeT domain. The domain organizations of *AzaA* and *AzaB* are parallel to that of *AfoE* and *AfoG*, respectively.

Analysis of nearby genes in the dual PKS cluster reveals the presence of ASPNDRAFT\_132962 (*azaR*) encoding for a Zn(II)<sub>2</sub>Cys<sub>6</sub> zinc finger transcription factor, which could regulate the coordinated expression of the genes in a cluster (Keller et al., 2005). Upstream and downstream of the PKSs, we find several tailoring enzymes that may modify the polyketide product. Interestingly, most of these genes have a corresponding homolog in the asperfuranone gene cluster, including *azaC* (*afoC* homolog), *azaG* (*afoF* homolog), *azaH* (*afoD* homolog), *azaJ* (AN1030.3 homolog), and *azaL* (*afoF* homolog), each with > 30% identity to their *afo* counterparts (Table 1). However, four additional genes are also found in the cluster, including an acyltransferase (*azaD*), a ketoreductase (*azaE*), an AMP-dependent CoA ligase (*azaF*), and a P450 monooxygenase (*azaI*) (Table 1). The extensive overlap between the *aza* and *afo* gene clusters suggests that the *aza* cluster may encode for the production of an asperfuranone-like compound.

### Overexpression of *azaR* Activates the *aza* Pathway

Among the polyketide metabolites cataloged for *A. niger* (Nielsen et al., 2009), none appears to be a likely candidate of the *aza* cluster. RT-PCR analysis of *azaA* and *azaB* showed that both PKS genes were weakly transcribed under our laboratory culturing conditions (Figure 3A). Ectopic integration and overexpression of pathway-specific transcriptional activator has been used as a strategy to activate the expression of silent gene cluster for production of cryptic metabolites (Bergmann et al., 2007). To interrogate the *aza* cluster, we constructed the overexpression plasmid pAZ44 by cloning *azaR* into pBARGE1 (obtained from FGSC), which contains a P<sub>gpdA</sub> for gene expression in fungi and a *bar* gene that confers resistance to glufosinate. Upon PEG-mediated protoplast transformation and selection, several transformants emerged and produced a yellow pigmentation not observed in the wild-type strain. The genomic DNA from one of the transformants (T1) was extracted and integration of the vector was confirmed via amplification of P<sub>gpdA</sub>::*azaR* present in the pAZ44 plasmid.

To determine which genes are transcriptionally activated by the regulator, RT-PCR analysis was performed on the wild-type (WT) and the *A. niger* T1 strains under identical conditions. We amplified a ~600 bp segment in each gene, particularly in those regions with introns, to distinguish between the products from spliced cDNA and potential genomic DNA carryover during RNA extraction. In addition to *azaA*, *azaB*, and *azaR*, 10 other



**Figure 2. Comparison of the *A. niger* aza Cluster with the *A. nidulans* afo Cluster and Their Respective Products**

The following PKS domain abbreviations are used: SAT, starter unit:ACP transacylase; KS, ketosynthase; AT, malonyl-CoA:ACP transacylase; PT, product template; ACP, acyl carrier protein; CMeT, C-methyltransferase; DH, dehydratase; KR, ketoreductase; ER, enoyl reductase; R, reductase. Structure of **1** is confirmed by NMR (see Table S1; Figure S5) and MS/MS analysis (Figure S2).

genes were significantly upregulated (Figure 3A). The genes immediately upstream of *azaC* and downstream of *azaB* showed no increase in transcription levels compared with the WT (Figure 3A). We therefore hypothesize that the 13 transcribed genes comprise the *aza* cluster.

LC-MS analysis of the metabolic profile of the activated *A. niger* T1 revealed the presence of several compounds not found in the WT strain (Figure 3B). The culture was scaled up to 2 l to obtain sufficient compound for structure elucidation. In addition, a time course of the metabolites produced by T1 was performed to identify possible biosynthetic intermediates of the *aza* pathway (Figure 3B).

From the day 2 and day 4 cultures of *A. niger* T1, compound **1** ( $m/z$  363 [M+H]<sup>+</sup>,  $\lambda_{\max}$  330 nm) was the most abundant product (Figure 3B; Figure S1A available online); it was purified as a yellow gum from the day 4 culture at an approximate yield of 60 mg/l. High-resolution mass of **1** is consistent with the molecular formula of C<sub>19</sub>H<sub>22</sub>O<sub>7</sub> (observed  $m/z$  363.1465 [M+H]<sup>+</sup>, calculated 363.1444). The pure compound was characterized by <sup>1</sup>H and <sup>13</sup>C NMR spectroscopy, which showed all of the 19 carbon and 22 proton signals (Table S1; Figure S5). From the 2D HSQC-INEPT135 and HMBC correlations (Figure S5), it is apparent that there are two substructures in the molecule: a fully saturated aliphatic chain and a conjugated cyclic moiety. The highly reduced chain is consistent with a 2,4-dimethyl hexanoyl structure characterized by three methyls, two methylenes, and two methines. We did not find any HMBC correlation beyond the carbonyl carbon with a chemical shift of 176.86 ppm (Figure S5D). Meanwhile, we assigned the rest of the signals to a bicyclic ring system consisting of three aromatic protons [ $\delta_{\text{H}}$  7.93 ppm (s), 7.31 ppm (s), 5.88 ppm (s)] with a carboxylic acid

moiety, which corresponded to a low-field proton signal [ $\delta_{\text{H}}$  10.2 ppm (br s)]. We identified that the only available position on the bicyclic ring system for the linkage to the 2,4-dimethyl hexanoyl moiety is the oxygen atom bonded to the methylated carbon ( $\delta_{\text{C}}$  84.06 ppm). We therefore propose the structure of **1** as shown in Figure 2, with the two substructures connected by an ester bond at C1' and C4. MS/MS analysis supports this linkage, which showed a fragment consistent with the mass of the bicyclic core ( $m/z$  237 [M+H]<sup>+</sup>) (Figure S2). Thus, **1** was determined to belong to the azaphilone family of compounds, which bears the typical pyrano-quinone bicyclic core, and was named azanigerone A. The configurations of the chiral centers in **1** are yet to be determined. The structure of **1** resembles lunatoic acid A, a previously characterized azaphilone from the plant pathogen *Cochliobolus lunatus* (Nukina and Marumo, 1977) (Figure 1). The aliphatic acyl chains of the two molecules are identical as evidenced by comparison to published values for lunatoic acid A (Nukina and Marumo, 1977). The main structural difference is the side chain on C9, where a longer acrylic acid is in place in lunatoic acid A instead of the C<sub>1</sub> carboxylic acid in **1**.

In addition to **1**, several other compounds were observed in the culture at different time points. Compound **2** ( $m/z$  251 [M+H]<sup>+</sup>,  $\lambda_{\max}$  300 nm) was produced on day 2 but disappeared on day 4 and beyond (Figure 3B; Figure S1B). This compound was purified and its structure was determined to be the tetra-substituted benzaldehyde FK17-P2a (Figure 3C; Table S1) that was previously isolated from *Aspergillus versicolor* and *Pseudobotrytis terrestris* (Arai and Sano, 1994; Yamaguchi et al., 2004). FK17-P2a should be a polyketide that originates from a methylated hexaketide that can be synthesized by a NR-PKS, which undergoes C2-C7 intramolecular cyclization and reductive release

**Table 1. Putative Functions and Homologs of the Genes in the *aza* Cluster**

Locus Tag (ASP/NIDRAFT_)	Gene Name	Putative Function	Characterized Homolog	Protein Identity/Similarity (%)
54405	N/A	Hydrolase	–	–
127101	N/A	Thioesterase	–	–
50208	<i>azaC</i>	Esterase/lipase	AN1032.3, <i>afoC</i>	44/56
189181	<i>azaD</i>	Acyltransferase	<i>F. sporotrichioides</i> Tri101	33/49
212676	<i>azaE</i>	Ketoreductase	<i>S. cerevisiae</i> GRE2	35/50
188806	<i>azaF</i>	Acyl:CoA ligase	<i>Physcomitrella patens</i> 4CL2	35/53
189194	<i>azaG</i>	FAD-dependent oxygenase	AN1035.3, <i>afoF</i>	37/64
188800	<i>azaH</i>	Salicylate monooxygenase	AN1033.3, <i>afoD</i>	41/58
43449	<i>azal</i>	Cytochrome P450	<i>A. flavus</i> CypX	30/49
56946	<i>azaA</i>	NR-PKS	AN1034.3, <i>afoE</i>	44/60
132962	<i>azaR</i>	Zn <sub>2</sub> Cys <sub>6</sub> regulator	AN1029.3, <i>afoA</i>	34/48
43447	<i>azaJ</i>	Dehydrogenase	AN1030.3	40/53
188912	<i>azaK</i>	Efflux transporter	–	–
132654	<i>azaL</i>	FAD-dependent oxygenase	AN1035.3, <i>afoF</i>	33/55
188817	<i>azaB</i>	HR-PKS	AN1036.3, <i>afoG</i>	43/60
188504	N/A	Cytochrome P450	–	–
50199	N/A	Oxidoreductase	–	–

Assignment of genes to *aza* cluster is based on transcriptional analysis on *A. niger* T1 (Figure 3A). N/A, not assigned (upstream and downstream genes).

to yield a 1,3-diketo benzaldehyde (Bailey et al., 2007). Further ketoreduction of the terminal ketone affords **2**.

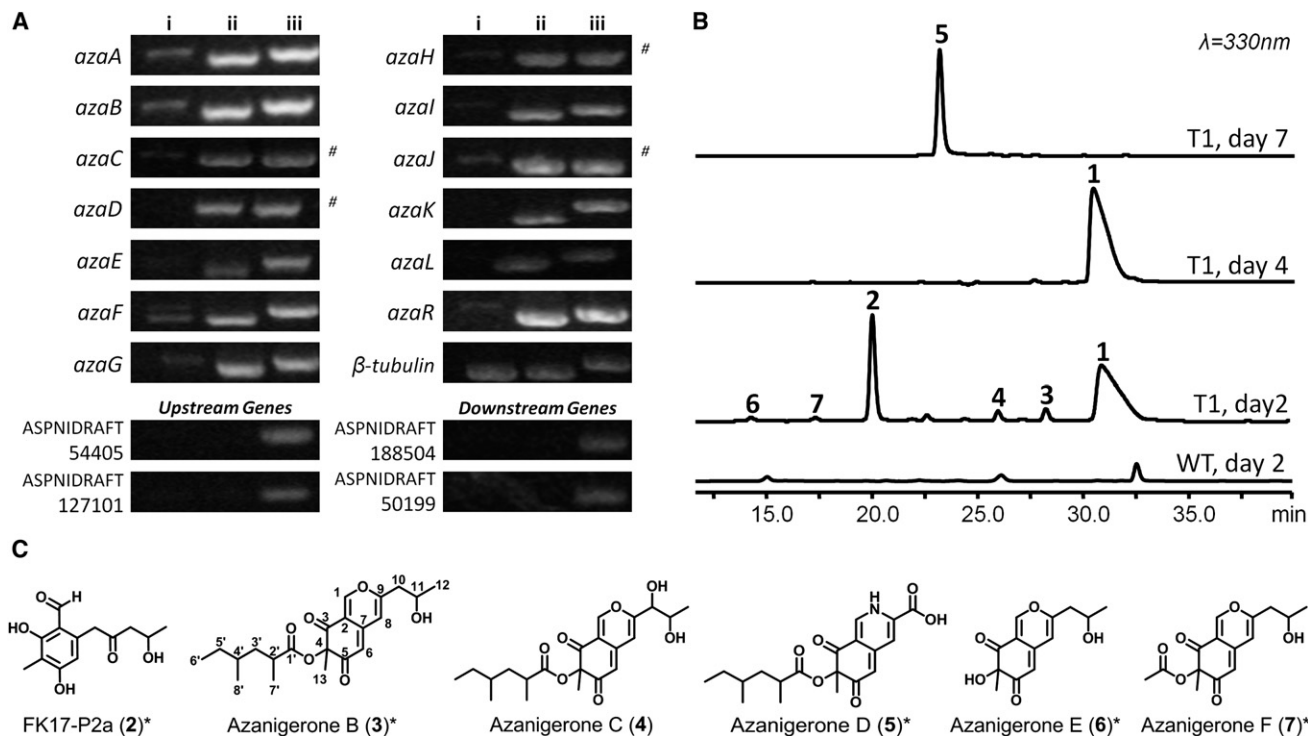
Compound **3** ( $m/z$  377 [M+H]<sup>+</sup>,  $\lambda_{\max}$  330 nm) was produced at a much lower yield during the earlier culture periods (Figure 3B; Figure S1C). The bicyclic core structure was resolved to be similar to **1**, but instead of the carboxylic acid moiety, three distinct proton signals were detected: a terminal methyl doublet ( $\delta_{\text{H}}$  1.32 ppm), a methylene doublet ( $\delta_{\text{H}}$  2.54 ppm), and a methane multiplet ( $\delta_{\text{H}}$  4.16 ppm) (Table S2; Figure S7). These signals are consistent with a 2-hydroxypropyl substitution at the C9, which is consistent with the  $m/z$  251 [M+H]<sup>+</sup> fragment from MS/MS analysis (Figure S3). The 2-hydroxypropyl moiety is reminiscent of the terminal portion of the 2-oxo-4-hydroxypentyl substituent in **2**, which suggests that **2** may be an intermediate in the synthesis of **3** (Figure 3C). Meanwhile, **4** ( $m/z$  393 [M+H]<sup>+</sup>,  $\lambda_{\max}$  330 nm) was produced at an even lower titer; its structure was not fully elucidated by NMR due to insufficient material obtained (Figure 3B; Figure S1D). By comparing the MS/MS peaks of **3** and **4**, as well as the molecular formula of **4** obtained through high-resolution MS (C<sub>21</sub>H<sub>28</sub>O<sub>7</sub>, observed  $m/z$  393.1908 [M+H]<sup>+</sup>, calculated 393.1913) (Figures S3 and S4), we propose that **4** could be a 10-hydroxy derivative of **3** (Figure 3C), and it may also be an intermediate toward the biosynthesis of **1**.

After 7 days of culturing, a new compound **5** ( $m/z$  362 [M+H]<sup>+</sup>,  $\lambda_{\max}$  357 nm) replaced **1** as the dominant product in the *A. niger* T1 culture (Figure 3B; Figure S1E). Compound **5** was purified from a 7-day culture, and its structure was confirmed via NMR to be the vinylogous  $\gamma$ -pyridone derivative of **1** (Figure 3C; Figure S8; Table S2). This compound is likely originated from **1** via a characteristic azaphilone reaction with amines present in the culture medium. Indeed, a compound with the identical retention time, mass, and  $\lambda_{\max}$  was obtained by reacting purified **1** with NH<sub>4</sub>OH (data not shown).

### Knockout of *azaB* Demonstrates a Convergent Synthesis of **1** by *AzaA* and *AzaB*

Inspection of the structure of **1** suggests that the molecule is assembled convergently from two discrete polyketide chains. We propose that the dimethylhexanoate moiety is produced by the HR-PKS *AzaB*, while the bicyclic core is cyclized from an unreduced polyketide backbone synthesized by the NR-PKS *AzaA*. This is in stark contrast to the sequential mode of collaboration between *AfoE* and *AfoG* in the highly parallel asperuranone gene cluster (Chiang et al., 2009) and underscores the difficulties in accurately predicting product structures among fungal PKSs.

Therefore, in the convergent synthesis model, disruption of *azaB* in T1 should not affect the function of *AzaA* and should lead to accumulation of the pyrano-quinone. To test this model, genetic disruption of *azaB* in *A. niger* T1 was performed using a double-crossover recombination with the zeocin-resistant gene *ble* (Figure 4A). Two desired  $\Delta$ *azaB* transformants were identified via diagnostic PCR (Figure 4A). After culturing and metabolite analysis, the production of **1** was confirmed to be abolished in the  $\Delta$ *azaB* strains. Instead, two new compounds (**6** and **7**) that have UV spectra characteristic of pyrano-quinone (identical to that of **1**) emerged (Figure 4B). Compounds **6** and **7** have molecular masses of 250 and 292, respectively (Figures S1F and S1G). Both compounds were purified and their structures were elucidated by 1D and 2D NMR (Table S3; Figures S9 and S10). Compound **6** contains the same bicyclic core and the 2-hydroxypropyl side chain as **3**, but it lacks the 2,4-dimethylhexanoyl ester as a result of *azaB* disruption (Figure 3C). Compound **7** is the O4-acetylated version of **6** (Figure 3C). In addition, the benzaldehyde **2** observed in the *A. niger* T1 strain was present in the early cultures of the  $\Delta$ *azaB* strains (Figure 4B). Upon closer examination, trace amounts of **6** and **7** were also found in *A. niger* T1 culture on day 2 (Figure 3B);



**Figure 3. Activation of the *aza* Pathway via Overexpression of *azaR***

(A) Transcriptional analysis by RT-PCR of the *aza* genes in *A. niger* WT (i) and the activated T1 strain (ii). PCR from genomic DNA (iii) are shown for comparison. #Intronless genes.

(B) Time course of *A. niger* T1 metabolites. Metabolic profiles of the activated *A. niger* T1 strain on days 2, 4 and 7 and of the WT on day 2.

(C) Compounds observed in culture of *A. niger* T1 strain (see Figure S1). Structures indicated with an asterisk are confirmed by NMR (see Tables S1–S3; Figures S6–S10). MS/MS analyses of **3** and **4** are in Figures S3 and S4.

the non-acetylated **6** is likely an intermediate in the biosynthesis of **1**.

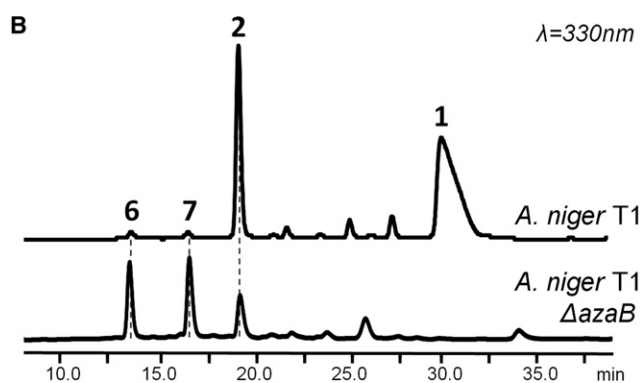
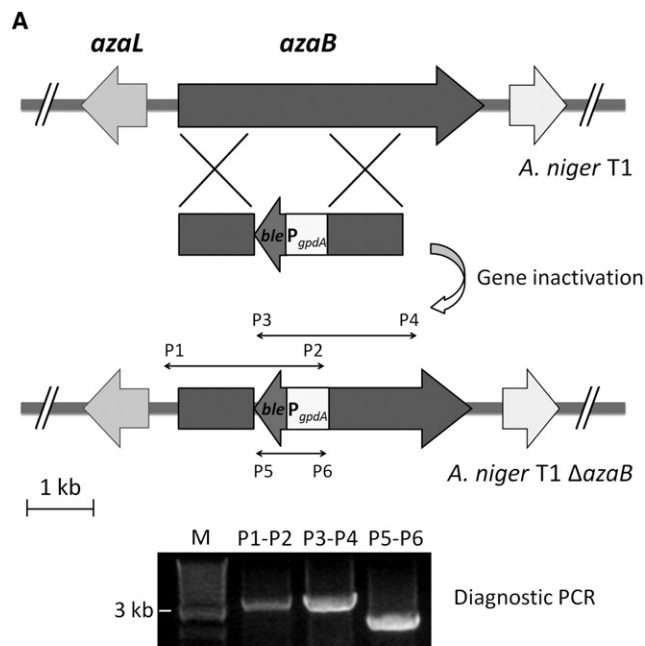
### In Vitro Reaction of AzaH with the Benzaldehyde Intermediate Confirms Its Roles in Hydroxylation and Pyran-Ring Formation

The formation of the bicyclic core of azaphilones has not been elucidated to date. It has been suggested that it proceeds from a benzaldehyde precursor in the asperuranone biosynthesis (Chiang et al., 2009), which is structurally similar to **2** isolated from *A. niger* T1 in early culturing times. Because **2** is isolated in stable form, the route to the formation of the pyran ring in the biosynthesis of **1** must be enzyme-catalyzed.

We initially proposed that AzaC is involved in the pyran-ring cyclization. Results of an NCBI conserved domain search indicate that both AzaC and its homologs (CtnB from the citrinin pathway, and AfoD) belong to the esterase-lipase family of serine hydrolases and are not oxidoreductases as assigned previously for CtnB and AfoD (Chiang et al., 2009; Sakai et al., 2008). As **1**, citrinin, and asperuranone all contain a fused heterocyclic ring system, AzaC, CtnB, and AfoB may be responsible for their corresponding heterocycle formation; a similar hypothesis has been proposed in a recent study as well (Davison et al., 2012). Accordingly, the recombinant AzaC protein (30 kDa) was purified and assayed with **2**. However, no conversion of the substrate **2** was observed in the in vitro reaction (Figure 5A). We therefore

reasoned that AzaC might act on a different substrate, possibly the C4-hydroxylated derivative of **2**.

We then searched for genes in the *aza* cluster that may hydroxylate the C4 position in **2**. AzaH, an FAD-dependent monooxygenase, was found to be homologous to a recently characterized monooxygenase TropB (43% identity, 61% similarity), which was recently shown to hydroxylate the corresponding carbon of 3-methylorcinolaldehyde during tropolone biosynthesis (Davison et al., 2012). To examine its role in the synthesis of **1**, AzaH was expressed and purified from *E. coli* BL21(DE3), and a series of in vitro assays including AzaH and AzaC/AzaH in the presence of **2** was performed. Surprisingly, incubation of both purified AzaH or AzaC/AzaH with **2** and NADPH yielded a new product that has an identical UV spectrum, retention time, and mass as **6** (Figure 5A). This indicates that AzaH alone is capable of converting **2** to **6**. However, in the absence of NADPH, which is needed for the regeneration of the reduced flavin cofactor, no substrate conversion by AzaH was observed. Therefore, the flavin-dependent hydroxylation of C-4 is likely a prerequisite for the subsequent pyran formation. Further LC/MS time-course analysis of the reaction of AzaH with **2** showed that the conversion proceeds rapidly with no detectable uncyclized C4-hydroxylated intermediate (Figure 5B). These results confirmed the critical role of AzaH in morphing the benzaldehyde intermediate into the bicyclic, pyran-containing azaphilone chromophore.



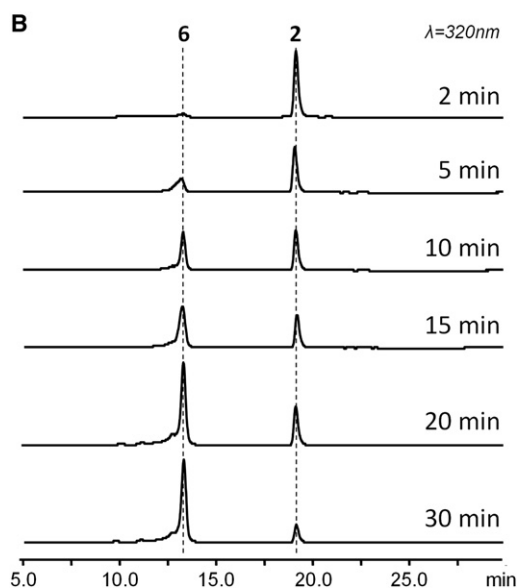
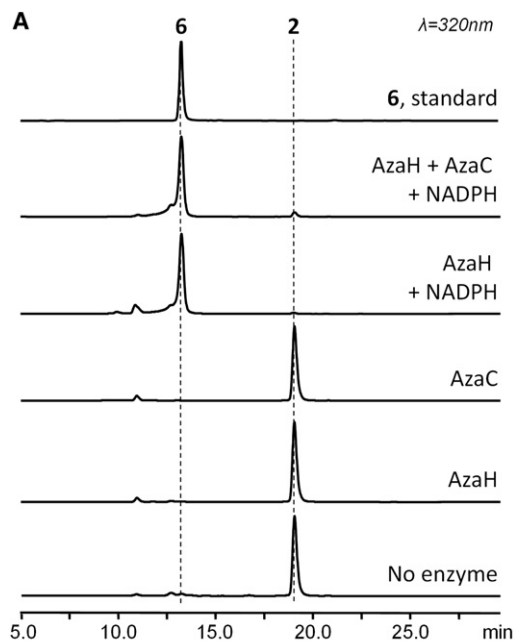
**Figure 4. Deletion of *azaB* Confirms the Involvement of the Cluster in Biosynthesis of **1****

(A) Knockout strategy employed for *azaB*. Correct integration is verified by using the three sets of primers shown.

(B) HPLC traces demonstrating the abolishment of **1** production in the  $\Delta$ *azaB* strain and the accumulation of **6** and **7** (see Table S3; Figures S9 and S10).

### Proposed Pathway for Biosynthesis of the Azanigerones

The RT-PCR transcriptional analysis indicates that there are a total of 12 genes in the *aza* cluster, which were activated upon overexpression of the pathway-specific regulator *azaR* (Figure 3A). Combining the structural characterization of the compounds from the activated *A. niger* T1 culture and T1  $\Delta$ *azaB*, as well as in vitro confirmation of the role of AzaH, a biosynthetic pathway leading to the production of **1** can be proposed (Figure 6). Based on the time course studies of the metabolic profile of *A. niger* T1 (Figure 3B), **1** is most likely to be the end product of the pathway, as it is the dominant compound in the culture after 4 days. We propose that the biosynthesis of **1** begins with the polyketide assembly by AzaA, which is a typical Clade III NR-PKS with a domain organization of SAT-KS-AT-PT-ACP-CMeT-R from the N- to C-terminal. AzaA forms the hexaketide

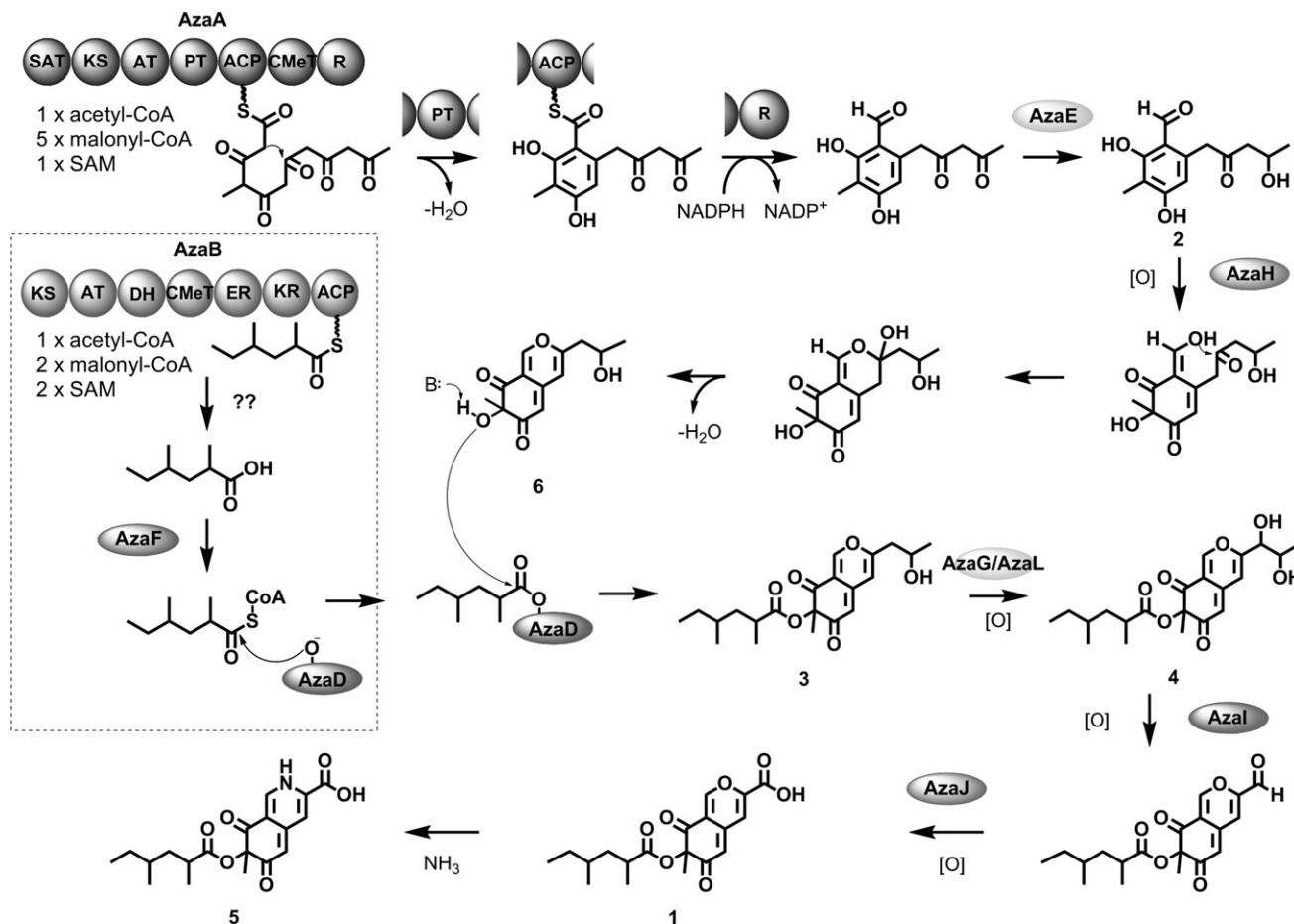


**Figure 5. In Vitro Assays Demonstrate AzaH-Mediated Pyran-Cyclization**

(A) HPLC analysis of the in vitro assays of AzaH and/or AzaC (10  $\mu$ M) with **2** (50  $\mu$ M), and comparison with the standard **6**. Conversion of **2** to **6** was observed in the presence of AzaH and NADPH (0.5 mM).

(B) Time course of AzaH-catalyzed formation of **6** from **2**. AzaH (0.5  $\mu$ M) was assayed with substrate **2** (50  $\mu$ M) and NADPH (0.5 mM). Reactions were quenched at specified time with acetonitrile (100  $\mu$ l).

precursor from successive condensations of five malonyl-CoA units, presumably with a simple acetyl-CoA starter unit. The reactive polyketide chain then undergoes a PT-mediated C2-C7 cyclization to afford the aromatic ring and is eventually released as an aldehyde through the R-domain, in a manner similar to the previously characterized 3-methylorcinaldehyde synthase in *Acremonium strictum* (Bailey et al., 2007). The



**Figure 6. Proposed Biosynthetic Pathway for Production of Azanigerones from the *aza* Cluster**

putative ketoreductase AzaE, a homolog of the methylglyoxal reductase in *Saccharomyces cerevisiae* (35% identity), is proposed to catalyze the reduction of the terminal ketone resulting in the early culture product **2**.

The monooxygenase AzaH was demonstrated to be the only enzyme required to convert **2** to **6** in vitro. We propose that AzaH first hydroxylates the benzaldehyde intermediate **2** at C4, which triggers the formation of the pyran-ring to afford **6** (Figure 6). In parallel, the 2,4-dimethylhexanoyl chain is synthesized by the HR-PKS AzaB and is proposed to be transferred to the C4-hydroxyl of **6** by the acyltransferase AzaD directly from the ACP domain of AzaB via a similar mechanism as the LovD acyltransferase (Xie et al., 2009). Alternatively, the 2,4-dimethylhexanoyl chain may be offloaded from the HR-PKS as a carboxylic acid and converted to an acyl-CoA by AzaF, which shares 35% identity to the 4-coumarate:CoA ligase (Silber et al., 2008). The resulting acyl-CoA molecule could then be taken up as a substrate by AzaD to form **3** (Figure 6). Like the homolog acetyltransferase Tri101 in trichothecene biosynthesis (33% identity to AzaD) (Garvey et al., 2008), AzaD belongs to the BAHD family of acyltransferases that catalyze transfer of various acyl-CoA substrates (D'Auria, 2006). It is interesting to note that the C4-O-acetylated **7** is also observed in the activated *A. niger* T1 and  $\Delta$ *azaB* cultures (Figure 4B). This suggests that AzaD may

have a relaxed specificity toward acyl-CoA substrates, although it is entirely possible that other endogenous acyltransferases might be involved in this reaction.

To yield the carboxylic acid substituent in **1**, the hydroxypropyl side chain of **3** would need to undergo a C-C oxidative cleavage. We propose that this C-C bond cleavage is catalyzed by a cytochrome P450 AzaI encoded in the cluster. Similar reactions have been observed in other P450s, with the CYP11A involved in the decomposition of cholesterol into pregnenolone and 4-methylpentanal as the archetypal enzyme (Burstein and Gut, 1971). Such P450 is proposed to act on a vicinal diol that leads to a C-C bond scission either through an alkoxyradical intermediate or a peroxy complex (Ortiz de Montellano and De Voss, 2005). In the biosynthesis of **1**, we propose that **3** first undergoes hydroxylation at C10, possibly catalyzed by one of the two FAD-dependent monooxygenases encoded in the cluster, AzaG or AzaL, resulting in the vicinal diol (Figure 6). A small amount of this putative intermediate **4**, with mass and fragmentation pattern consistent with such a diol, was isolated from the 2-day culture of *A. niger* T1 (Figure 3B; Figure S4). Oxidative cleavage **4** by AzaI would yield the corresponding aldehyde derivative of **1**. A homolog of AzaI, CypX, also exists in the aflatoxin biosynthesis in *A. parasiticus* (35% identity) and was found to convert averufin to hydroxyversicolorone via a still unknown oxidative cleavage

and ring rearrangement step (Wen et al., 2005). Finally, the dehydrogenase AzaJ (homologous to the AN1030.3, 40% identity) is proposed to convert the aldehyde functional group into the carboxylic acid, completing the conversion from **3** to **1**. Alternatively, the oxidation of aldehyde to carboxylic acid may be catalyzed by the same P450 enzyme AzaI via consecutive oxidation or by endogenous alcohol dehydrogenase. Interestingly, the proposed oxidative chain-shortening does not occur in the absence of the 2,4-dimethylhexanoate ester. Both **5** and **6** contain the hydroxypropyl chain and are not converted to the corresponding acids in  $\Delta$ azaB culture.

## DISCUSSION

Genome mining, in particular activation of silent secondary metabolic pathways by overexpression of a pathway-specific regulator, is becoming a useful strategy for natural product discovery in fungi (Brakhage et al., 2008; Winter and Tang, 2012). From overexpression of the cluster-specific regulator, we have uncovered novel compounds that belong to the family of azaphilones. The hallmark feature of azaphilones is the presence of a highly oxygenated pyrano-quinone bicyclic core of polyketide origin (Osmanova et al., 2010). The name derives from their tendency to react with primary amines to yield  $\gamma$ -pyridones that result from the exchange of the pyrane oxygen with nitrogen. Over 170 different azaphilone compounds have been identified from hundreds of fungal species across 23 genera from 13 families, and many of the azaphilones exhibit diverse biological activities, including antibacterial, antifungal, and antitumor activities (Osmanova et al., 2010). Several common modifications that contribute to the structural diversification of the common bicyclic chromophore include but are not limited to (1) aliphatic side chains at C9; (2) aliphatic esters at O4; (3) halogen substitution on the bicyclic core, as in sclerotiorin (Chidananda and Sattur, 2007); (4) additional fused rings, as in ankaflavin (Manchand et al., 1973); (5) aromatic esters at O4, as in mitorubrinol (Quang et al., 2005); and (6) molecular dimers, as in chaetoglobins A (Ming Ge et al., 2008) (Figure 1).

Despite their apparent ubiquity among fungi, the genetic and molecular basis for the biosynthesis of these azaphilones has not been thoroughly explored or understood. Besides labeling studies suggesting a polyketide origin for the bicyclic core (Ogihara et al., 2000), only the azaphilone-like compound asperfuranone and citrinin have been linked to their corresponding biosynthetic gene clusters (Chiang et al., 2009; Sakai et al., 2008). In this study, we have discovered a silent gene cluster in *A. niger* responsible for producing a series of azaphilones. Characterization of this *aza* gene cluster has provided important insights on the biosynthesis of this family of compounds.

In total, six azaphilones (azanigerone A–F), which were previously uncharacterized, to our knowledge, were isolated, and all share a substituted bicyclic core indicative of a common biosynthetic origin. We also observed their signature reaction with amine groups in the conversion of the major product **1** into the vinylogous  $\gamma$ -pyridone **5**. All six compounds contain a C4 hydroxyl group, which is a post-PKS tailoring modification observed in a multitude of other azaphilones. With the exception of **6**, the tertiary alcohol at C4 of azanigerones is acylated with an aliphatic acyl or an acetyl group, which is another common

feature among many azaphilones. The isolation of pyran intermediates at early stages of the *A. niger* T1 culture underscores the convergent actions of the two PKSs in the production of these molecules (Figure 3B).

A key discovery in this study is the enzyme responsible for the formation of the characteristic pyrano-quinone core common to all azaphilones. It has been proposed that the biosynthesis of azaphilones proceeds from a benzaldehyde intermediate (Chiang et al., 2009), but the enzymology of the pyran-ring formation step has not been elucidated. We have discovered that the cyclization of the benzaldehyde is mediated by the C4-hydroxylation catalyzed by the FAD-dependent monooxygenase AzaH. We propose that the hydroxylation at C4 by AzaH dearomatizes the ring and causes keto-enol tautomerization at the C1 aldehyde, leading to the condensation between the C1 enol and the C9 carbonyl to yield the pyran ring (Figure 6). Our finding that the C4-hydroxylation and pyran-ring cyclization occur very rapidly suggests that these two features are coupled. This is supported by the fact that, with a few exceptions, a majority of the azaphilones contain such geminal methyl, hydroxyl (or ester) substitution at the pyrano-quinone core (Figure 1) (Osmanova et al., 2010). For the few azaphilones that do not have this feature, such as in the case of citrinin, a different mechanism of cyclization might take place. Coincidentally, the citrinin gene cluster lacks the *azaH* homolog (Sakai et al., 2008).

It is curious to note that, with the exception of a few genes, the *aza* and the *afo* clusters are highly similar, with multiple gene homologs sharing >30% protein sequence identity (Figure 2; Table 1). However, the structures of the two compounds are considerably different. A striking disparity between the two biosynthetic pathways is how the dual PKS systems partner to form their respective products. In the case of the *afo* cluster, the HR-PKS AfoG synthesizes the tetraketide intermediate, which is passed on to the SAT domain of the NR-PKS AfoE. Such sequential mode of collaboration occurs in other HR/NR-PKS tandem systems, such as in the biosynthesis of resorcylic acid lactones (Zhou et al., 2010). We expected the two PKSs in *aza* cluster, AzaA and AzaB, to follow the same tandem mode of PKS-PKS collaboration. To our surprise, the two PKSs acted independently of each other in the convergent synthesis of the two substructures of **1**. Whereas the  $\Delta$ afoG culture completely disrupted the production of asperfuranone or any appreciable intermediate, compounds **6** and **7** lacking the reduced triketide chain were isolated in the  $\Delta$ azaB culture. Previous studies have pointed to the N terminus SAT as a specificity-conferring domain that selects the starter unit (Crawford et al., 2008). By multiple protein sequence alignment, it was noticed that the SAT domains of AzaB and AfoG have a lower identity (35% versus 43%) compared with the rest of the protein, although both contain the GXCXG motif found in most SATs. It is likely that the selectivity of the SAT and its ability to interact with HR-PKS hold the key for which mode of collaboration will take place between a pair of HR-PKS and NR-PKS. The *azaA-azaB* dual PKS system in *A. niger* is the first example of a convergent mode of collaboration between an HR and an NR-PKS; elucidation of the basis for the intriguing difference between *aza* and *afo* pathways could facilitate better prediction of polyketide products from such pathways.

Another notable difference between the structures of azanigerones and asperfuranone is the formation of a furan in the latter



pathway instead of a pyran ring that is common in azaphilones. Both pathways are proposed to go through a benzaldehyde intermediate. However, in the case of asperfuranone, it is likely that the benzylic carbon hydroxylation precedes the dearomatizing hydroxylation. It was previously proposed that AfoD catalyzes this benzylic carbon hydroxylation, which is the key branching point in the biosynthesis of asperfuranone and the azaphilone sclerotiorin (Davison et al., 2012; Somoza et al., 2012). However, it is more likely that AfoD has a similar function as the homolog AzaH characterized in this study, i.e., the formation of the tertiary alcohol. Thus AfoF, another FAD-dependent monooxygenase, may catalyze this benzylic carbon hydroxylation step in the asperfuranone pathway. Two AfoF homologs, AzaG and AzaL, are also found in the *A. niger* aza cluster (Table 1). We propose that one of them may act later in the pathway in the hydroxylation of C10 to form the precleavage vicinal diol.

Among the aza genes that share a homolog to the *afo* cluster, the function of AzaC remains enigmatic. The possibility of its involvement in the heterocycle formation has been excluded by the results in this study. In line with the results of an NCBI conserved domain search that suggest an esterase-lipase function for this enzyme, we propose that AzaC might participate in the offloading of the 2,4-dimethylhexanoyl chain from the ACP of AzaB (Figure 6). The role of the homologs AfoC and CtnB in the biosynthesis of asperfuranone and citrinin, respectively, is also unclear and should be the subject of further investigation.

## SIGNIFICANCE

**Activation of the silent aza cluster has led to the discovery of compounds named azanigerones, which have not been isolated from *A. niger* before. Characterization of the pathway has shed light on the biosynthesis of the azaphilone group of compounds, which are structurally diverse and exhibit a wide range of bioactivities. The aza pathway represents an example of a convergent mode of collaboration between an HR- and an NR-PKS, as opposed to the sequential mode of collaboration commonly observed in other dual PKS systems. More importantly, the enzymatic basis for the formation of the pyrano-quinone core common to azaphilones has been revealed for the first time, and the mechanism can likely be generalized to biosynthesis of other azaphilones. Thus, this study provides an important basis for further molecular genetics and biochemical investigations of azaphilone pathways. The sequence information will aid in identification of the gene clusters encoding for known bioactive azaphilones and discovery of new azaphilones by genome mining. The many structural diversifications on a common bicyclic core observed among this family of fungal natural products also offer a unique opportunity for combinatorial biosynthesis to expand the library of azaphilones and screen for enhanced bioactivities.**

## EXPERIMENTAL PROCEDURES

### Strain and Culture Conditions

*A. niger* ATCC 1015 was obtained from Agricultural Research Service (ARS) culture collection (as NRRL 328) and cultured at 28°C in potato dextrose agar medium. The activated *A. niger* T1 strain and the  $\Delta$ azaB mutant strains

were maintained in glucose minimal medium (GMM) containing 10 mM ammonium tartrate as nitrogen source (Chooi et al., 2010; Li et al., 2011) or yeast glucose medium (YG) (Szewczyk et al., 2006). *E. coli* strains XL1 (Stratagene) and TOP010 (Invitrogen) were used for routine cloning. *Saccharomyces cerevisiae* BJ5464 was used for in vivo yeast DNA recombination cloning. *E. coli* BL21(DE3) was used for protein expression.

### Molecular Genetic Manipulations

Polymerase chain reactions were performed using Phusion high-fidelity DNA polymerase (New England Biolabs), Platinum Pfx DNA polymerase (Invitrogen), and GoTaq Green Master Mix (Promega). PCR products were cloned into a PCR-Blunt vector (Invitrogen) for DNA sequencing and subcloning. Restriction enzymes (New England Biolabs) and T4 ligase (Invitrogen) were used, respectively, for the digestion and ligation of DNA fragments. Primers used for amplification were synthesized by Integrated DNA Technologies and are listed in Table S4.

Genomic DNA of *A. niger* ATCC 1015 was extracted using the ZYMO ZR fungal/bacterial DNA kit according to manufacturer's protocols. The *azaR* gene (including introns) was amplified from the genomic DNA and inserted into the fungal shuttle vector pBARGPE1 (digested with BglII and EcoRI) to yield pAZ44. The pBARGPE1 vector was obtained from Fungal Genetic Stock Center (FGSC) (Pall and Brunelli, 1993). The intronless *azaH* and *azaC* were also amplified from genomic DNA and inserted into the p8HIS expression vector (Jez et al., 2000) digested with EcoRI and NotI to yield pAZ81 and pAZ83, respectively.

The knockout cassette for *azaB* was assembled in yeast using the yeast recombination method (Colot et al., 2006). The two homologous regions were amplified from *A. niger* genomic DNA using primers containing overlapping regions with the yeast vector and the zeocin-resistance cassette. The zeocin-resistance cassette containing the *ble* gene under the *gpdA* promoter was amplified from pAN8-1 (Punt et al., 1987) using *gpdAfor* and *blerev*. The three DNA fragments were cotransformed with the linearized vector backbone derived from YEplac195 (Gietz and Sugino, 1988) into the *S. cerevisiae* BJ5464 (Jones, 1991) using S.c. EasyComp Transformation Kit (Invitrogen) and selected on uracil-dropout semisynthetic media. The resulting transformants were screened by colony-direct PCR, and the plasmid in the correct transformant was rescued using the Zymoprep Yeast Plasmid Miniprep Kit (Zymo Research) and transformed into *E. coli* for propagation and sequencing verification. The resulting plasmid was designated AZ61. This plasmid was used as template to obtain the linear *azaB* knockout cassette for transformation of *A. niger* T1.

### Fungal Transformation

The preparation of *A. niger* protoplasts and the PEG-mediated transformations were carried out as previously described (Li et al., 2011). The *A. niger* transformed with pAZ44 for *azaR* overexpression was selected on approximately 8 mg/ml of glufosinate prepared as described previously (Chooi et al., 2010). For genetic manipulation in the activated *A. niger* strain T1, a modified protocol was derived from the one developed for *Aspergillus nidulans* (Szewczyk et al., 2006). The following modifications were implemented: (1) the 2× protoplasting solution was prepared with 3 g of VinoTaste Pro; (2) the PEG solution used was composed of 25% PEG 4000, 10 mM CaCl<sub>2</sub>, and 10 mM Tris-HCl; (3) the transforming protoplasts were first plated on nonselective media for 5 hr before overlaying with soft agar (8 g/l) with 200 µg/ml of Zeocin; and (4) the protoplasts were stabilized with 1.2 M sorbitol.

### Expression Analysis by RT-PCR

The total RNA of *A. niger* mutant and wild-type strains were extracted using the Ambion RNA extraction kit. The first-strand cDNA was synthesized using the Oligo-dT primer and Improm-II reverse transcription system (Promega) according to the manufacturer's instructions. cDNA was then amplified with GoTaq Green Master Mix using gene-specific primers (Table S4).

### Expression and Purification of AzaC and AzaH

pAZ81 and pAZ83 were used to transform the BL21(DE3) cells for expression of AzaC and AzaH, respectively. The cells were cultured at 37°C and 250 rpm in 500 ml of LB medium supplemented with 35 µg/ml kanamycin to a final OD<sub>600</sub> of 0.4–0.6, followed by addition of 0.1 M isopropylthio-β-D-galactoside

(IPTG) to induce protein expression. The proteins were purified by Ni<sup>2+</sup> affinity chromatography as described previously (Li et al., 2011). Purified AzaC and AzaH were concentrated and exchanged into PBS (50 mM, pH 7.4) with the centrprep filters (Amicon). Protein concentration was determined with the Bradford assay using BSA as a standard.

#### In Vitro Assays with AzaC and AzaH

The reactions were performed at 50  $\mu$ l scale containing 100 mM phosphate buffer (pH 7.4) in the presence of 50  $\mu$ M of **2**, 0.5 mM NADPH, and 10  $\mu$ M of AzaH and/or AzaC. After 2 hr, the reactions were quenched by precipitating the enzyme/s with 100  $\mu$ l of acetonitrile. The acetonitrile/aqueous solution was concentrated 5-fold and directly analyzed on the LC-MS (Figure 5A). The time-course analysis was performed with the same assay condition, except that 0.5  $\mu$ M AzaH was used and the reactions were quenched at different time points (Figure 5B).

#### LC/MS Analysis

For small-scale analysis, the *A. niger* wild-type and transformants were grown in 10 ml GMMT under shaking conditions at 28°C. The cultures were extracted with equal volume of ethyl acetate (EA) with 0.1% trifluoroacetic acid (TFA) and evaporated to dryness. The dried extracts were dissolved in methanol for LC/MS analysis using a Shimadzu 2010 EV liquid chromatography mass spectrometer with positive and negative electrospray ionization and Phenomenex Luna 5  $\mu$ l 2.0  $\times$  10 mm C18 reverse-phase column. The samples were resolved on a linear gradient from 5% to 95% with CH<sub>3</sub>CN/H<sub>2</sub>O + 0.05% formic acid solvent system.

#### Isolation of Compounds

Most compounds were purified from shake flask cultures, except for **2**, which was obtained from 2-day static culturing. Compounds **1**, **3**, and **4** were extracted with EA after 4 days of culturing, **5** after 7 days, and the *azaB* knockout compounds, **6** and **7**, after 2 days. The following scheme was generally used to purify each compound from the crude extract: (1) chloroform-water partitioning; (2) hexane-90% methanol partitioning; (3) Sephadex LH-20 (GE) chromatography; and (4) preparative high pressure liquid chromatography with Phenomenex Luna 5  $\mu$ l 250  $\times$  1,000 mm C18 reverse-phase column using CH<sub>3</sub>CN/H<sub>2</sub>O + 0.1% TFA or +0.2% formic acid solvent system. DRX500 or ARX500 instruments were used to perform the NMR spectroscopy of the different compounds.

#### SUPPLEMENTAL INFORMATION

Supplemental Information includes four tables and ten figures and can be found with this article online at <http://dx.doi.org/10.1016/j.chembiol.2012.07.004>.

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