

Recent advances in genome mining of secondary metabolite biosynthetic gene clusters and the development of heterologous expression systems in *Aspergillus nidulans*

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Abstract Fungi are prolific producers of secondary metabolites (SMs) that show a variety of biological activities. Recent advances in genome sequencing have shown that fungal genomes harbor far more SM gene clusters than are expressed under conventional laboratory conditions. Activation of these “silent” gene clusters is a major challenge, and many approaches have been taken to attempt to activate them and, thus, unlock the vast treasure chest of fungal SMs. This review will cover recent advances in genome mining of SMs in *Aspergillus nidulans*. We will also discuss current updates in gene annotation of *A. nidulans* and recent developments in *A. nidulans* as a molecular genetic system, both of which are essential for rapid and efficient experimental verification of SM gene clusters on a genome-wide scale. Finally, we will describe advances in the use of *A. nidulans* as a heterologous expression system to aid in the analysis of SM gene clusters from other fungal species that do not have an established molecular genetic system.

Keywords *Aspergillus* · Secondary metabolite · Polyketide synthase · Nonribosomal peptide synthetase · Gene cluster

Introduction

Filamentous fungi have played an important role in the history of drug discovery and development. The secondary metabolites (SMs) that these organisms produce have served as a source of low molecular weight molecules with a variety of biological activities. Examples of these are antibiotics such as penicillin, immunosuppressants such as cyclosporine, antifungals such as griseofulvin and the echinocandins, and antihypercholesterolemic drugs such as lovastatin [11, 24, 35]. Many of the bioactive SMs that are easily accessible under conventional laboratory conditions have already been isolated and patented for drug development. However, advances in genome sequencing [23, 32, 38, 40] revealed that fungal species harbor an abundance of SM gene clusters and these far exceed the number of known metabolites produced by the species [45]. This potential abundance of SMs may reflect their importance in nature as a chemical arsenal for niche security [41]. The carefully controlled growth conditions in laboratory culture settings prevent any competition or life-threatening circumstances that would trigger the production of SMs, thereby leaving many of the gene clusters dormant. Activating these silent gene clusters, revealing their biosynthetic pathways, and isolating the SMs produced by these pathways is a major challenge in the search for new SMs.

Various approaches have been taken in attempts to activate silent SM gene clusters [14], including fusing of regulatable promoters to a pathway-specific transcription factor [5, 17], removal of genes required for heterochromatin

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formation [7], genome-wide analysis of mutants of LaeA, a global regulator of SM [8], co-incubation with microorganisms to mimic conditions in nature [48], and the “one strain many compounds” (OSMAC) strategy [6]. Most of these approaches were developed in *A. nidulans* due to the availability of highly efficient gene-targeting systems in this model organism. The developed approaches are often subsequently applied to other filamentous fungi.

In this review, we focus on recent advances in genome mining of secondary metabolism genes in *A. nidulans*. We also describe the current status of the annotation of the products of secondary metabolism genes in *A. nidulans*. We would also like to direct readers to the accompanying review in this issue by our collaborators Nancy Keller and Philipp Wiemann on general strategies for mining fungal natural products and to other recent reviews on this subject [10, 28, 53, 56, 61].

The status of annotating secondary metabolite genes in *A. nidulans*

Among the *Aspergillus* species, *A. nidulans* has been used as a model organism, making it the most comprehensively studied and best-characterized species in the genus with the largest body of literature. Most studies of secondary metabolite biosynthesis in *A. nidulans* have used strains derived from a common reference strain, *A. nidulans* FGSC A4. *A. nidulans* FGSC A4 was initially sequenced by Cereon Genomics (Monsanto) in 1998 to three-fold genome equivalent coverage and the sequence was publicly released in 2003. Shortly thereafter, additional sequencing was completed at the Whitehead Institute/MIT Center for Genomic Research to give a total of 13 genome-equivalent coverage. The seminal paper describing the *A. nidulans* genome was published in 2005 [23]. Access to this sequenced genome has allowed investigators to use sequence similarity to known genes from other species to mine for core genes that are involved in secondary metabolism in *A. nidulans*. Algorithms such as SMURF (Secondary Metabolite Unknown Regions Finder) [27] and antiSMASH (antibiotics and Secondary Metabolite Analysis Shell) [34] are extremely useful in predicting the core SM biosynthetic genes. Taking into consideration the most recent annotation and additional analysis of available genomic data, our group’s most recent estimate is that the *A. nidulans* genome contains 56 putative secondary metabolism core genes including 27 polyketide synthase genes (PKS), two polyketide synthase-like genes (PKS-like), 11 nonribosomal peptide synthetase genes (NRPS), 15 NRPS-like genes, and one hybrid NRPS-PKS gene. Table 1 and Fig. 1 show our current understanding of the products of these genes and the products from the pathways.

Bioinformatic advances

Since the original publication of the genome sequence data [23], *A. nidulans* gene annotations have been refined repeatedly to correct incomplete or inaccurate content [3, 4, 25, 39, 57]. The *Aspergillus* Genome Database (AspGD; <http://www.aspgd.org/>) provides gene and protein sequence data that are curated based on submitted information and published literature. Although the wealth of data and the availability of the algorithms mentioned previously have provided accurate predictions of core SM biosynthetic genes, it is still not possible to predict with accuracy the boundaries of secondary metabolite gene clusters or the functions of each member of the clusters based solely on genome sequence data. This is due to the fact that many of the genes surrounding the core SM biosynthetic genes often have unknown functions, making predictions of their involvement in the biosynthetic process of the SM almost impossible. Elucidation of biosynthetic gene clusters have thus been heavily dependent on experimental verification, a laborious process that involves single gene deletion of each gene with a suspected role in SM biosynthesis, followed by identification and characterization of SMs produced by the deletion strains. Improvements in “omics”-based methods for accurate prediction of SM gene cluster members and the availability of more precise annotations are desirable for a more rapid and efficient experimental verification of novel SM gene clusters.

Andersen et al. [2] recently published a novel strategy for the accurate prediction of SM gene cluster boundaries based on the fact that expression of genes of a given SM cluster is coordinately regulated. A DNA expression microarray was used to identify genes that were co-regulated with SM gene cluster backbone enzymes. A variety of culture media were selected that, based on SM profiling experiments, would elicit expression of as many gene clusters as possible. Samples were then taken from *A. nidulans* growing on the selected culture media for transcriptional profiling, and the generated data were combined with previously published data to form a superset of a total of 44 expression conditions for analysis. Andersen et al. developed clustering scores (CSs) that reflected the degree to which each gene was co-regulated with its neighbors. They developed statistical guidelines for identifying the extent of gene clusters, which were applied to the microarray data to generate cluster predictions. Comparisons with published data demonstrated that their algorithm predicted gene clusters with high accuracy and can even predict gene clusters that are scattered across different chromosomes. Using this algorithm, a list of 58 predicted SM gene clusters was generated.

These data have been curated at AspGD and applied as a criterion for the manual annotation of computationally

Table 1 Secondary metabolism gene clusters in *A. nidulans*

No	AspGD designation	Core gene name	Gene type	Metabolites isolated from <i>A. nidulans</i> ^a	References
1	AN0016	<i>pes1</i>	NRPS		
2	AN0150	<i>mdpG</i>	NR-PKS	Emodin (1), emodin analogs (2–10), shamixanthone (11), epishamixanthone (12), variecoxanthone A (13), emericellin (14), atochryson carboxylic acid (15), 1-hydroxy-6-methyl-8-hydroxymethylxanthone (16), paeciloxanthone (17), monodictyphenone (18), 3-(2,6-dihydroxyphenyl)-4-hydroxy-6-methyl-1(3H)-isobenzofuranone (19), arugosin A, H (20,21)	[7, 16, 36, 44, 46, 51]
3	AN0523	<i>pkdA</i>	NR-PKS	2-Ethyl-4,6-dihydroxy-3,5-dimethylbenzaldehyde (22)	[1]
4	AN0607	<i>sidC</i>	NRPS	Ferrirocic	[20]
5	AN1034	<i>afoE</i>	NR-PKS	(2Z,4Z)-4,6-Dimethylocta-2,4-dienoic acid (23), asperfuranone (24), 6-[(3E,5E)-5,7-dimethyl-2-oxonona-3,5-dienyl-2,4-dihydroxy-3-methylbenzaldehyde (25)	[17, 49]
6	AN1036	<i>afoG</i>	HR-PKS	(2Z,4Z)-4,6-Dimethylocta-2,4-dienoic acid (23), asperfuranone (24), 6-[(3E,5E)-5,7-dimethyl-2-oxonona-3,5-dienyl-2,4-dihydroxy-3-methylbenzaldehyde (25)	[17, 49]
7	AN1242		NRPS	Nidulanin A (27)	[2]
8	AN1680		NRPS-like		
9	AN1784	<i>pkjA</i>	HR-PKS		
10	AN2032	<i>pkhA</i>	NR-PKS	2,4-Dihydroxy-6-[(3E,5E,7E)-2-oxonona-3,5,7-trienyl]benzaldehyde (26)	[1]
11	AN2035	<i>pkhB</i>	HR-PKS	2,4-Dihydroxy-6-[(3E,5E,7E)-2-oxonona-3,5,7-trienyl]benzaldehyde (26)	[1]
12	AN2064		NRPS-like		
13	AN2545	<i>easA</i>	NRPS	Emericellamides (28–32)	[18]
14	AN2547	<i>easB</i>	HR-PKS	Emericellamides (28–32)	[18]
15	AN2621	<i>acvA</i>	NRPS	Penicillin	[31, 51]
16	AN2924		NRPS-like		
17	AN3230	<i>pkfA</i>	NR-PKS	Orsellinaldehyde (33), 3-(2,4-dihydroxy-6-methylbenzyl)-orsellinaldehyde (34), aspernidine A-E (37–41)	[1, 47, 58],
18	AN3386	<i>pkIA</i>	NR-PKS	7-Methyl-3-nonylisoquinoline-6,8-diol (35), 6-hydroxy-7-methyl-3-nonylisoquinoline-5,8-dione (36), 2,4-dihydroxy-3-methyl-6-(2-oxoundecyl)benzaldehyde (42), 4-hydroxy-3-methyl-6-(2-oxoundecyl)-2-pyrone (43)	[1]
19	AN3396	<i>micA</i>	NRPS-like	Microperfuranone (44)	[59]
20	AN3495	<i>inpA</i>	NRPS-like		
21	AN3496	<i>inpB</i>	NRPS		
22	AN3612		HR-PKS		
23	AN4827		NRPS-like		
24	AN5318		NRPS-like		
25	AN6000	<i>aptA</i>	NR-PKS	Asperthecin (45), 2,3,6,8,9-pentahydroxy-1-oxo-3-(2-oxopropyl)-1,2,3,4-tetrahydroanthracene-2-carboxylic acid (46)	[1, 29, 52]
26	AN6236	<i>sidD</i>	NRPS		
27	AN6431		HR-PKS		
28	AN6444		NRPS-like		
29	AN6448	<i>pkbA</i>	NR-PKS	2,5-Dimethylresorcinol (47), 3-methylorsellinic acid (48), cichorine (49), nidulol (50),	[1, 43]
30	AN6791		HR-PKS		
31	AN7071	<i>pkgA</i>	NR-PKS	Alternariol (51), citreoisocoumarin (52), analogs of citreoisocoumarin (53–56)	[1]
32	AN7084		PKS-like		
33	AN7489		PKS-like		

Table 1 continued

No	AspGD designation	Core gene name	Gene type	Metabolites isolated from <i>A. nidulans</i> ^a	References
34	AN7825	<i>stcA (pksST)</i>	NR-PKS	Norsolorinic acid (57), norsolorinic acid anthrone (58), sterigmatocystin (59)	[1, 12, 54, 62]
35	AN7837+ AN7838		HR-PKS		
36	AN7884		NRPS		
37	AN7903	<i>pkeA</i>	NR-PKS	2,4-Dihydroxy-3-methyl-6-(2-oxopropyl)benzaldehyde (60)	[1]
38	AN7909	<i>orsA</i>	NR-PKS	F9775A, B (61, 62), orsellinic acid (63), diorcinol (64), gerfelin (65), 10-deoxygerfelin (66)	[7, 42, 46, 48]
39	AN8105		NRPS-like		
40	AN8209	<i>wA</i>	NR-PKS	2-Acetoacetyl T4HN (67), naphthopyrone YWA1 (68)	[1, 21, 55]
41	AN8383	<i>ausA</i>	NR-PKS	Isoaustinone (69), analogs of isoaustinone (70, 71), austinol (72), dehydroaustinol (73), protoaustinoid (74), preaustinoid A3-A5 (75–77), austinoneol A (78), neo-austinone (79), austinolide (80)	[1, 30, 36]
42	AN8412	<i>apdA</i>	Hybrid	Aspyridone A, B (81, 82)	[5]
43	AN8513	<i>tdiA</i>	NRPS-like	Terrequinone A (83)	[9, 51]
44	AN8910		HR-PKS		
45	AN9005		HR-PKS		
46	AN9129		NRPS-like		
47	AN9226	<i>nrpA</i>	NRPS		
48	AN9243		NRPS-like		
49	AN9244		NRPS		
50	AN9291		NRPS-like		
51	AN10297		NRPS-like		
52	AN10430		HR-PKS		
53	AN10486		NRPS-like		
54	AN10576	<i>ivoA</i>	NRPS		
55	AN11191	<i>pkkA</i>	HR-PKS		
56	AN12440		NR-PKS		

PKS polyketide synthase, NRPS nonribosomal peptide synthetase, Hybrid hybrid PKS-NRPS, NR-PKS nonreduced polyketide synthase, HR-PKS highly reduced polyketide synthase

^a Bold numbers correspond to chemical structures shown in Fig. 1

predicted gene clusters as a part of a continued effort to improve and refine the prediction of SM gene cluster boundaries [25]. This updated gene cluster boundary annotation also incorporates published experimental data, synteny between clustered genes among different species, functional annotation of putative gene cluster members, and increase in the distance between predicted boundary genes and genes that are directly adjacent to it but not included in the cluster. This new and improved set of comprehensive SM gene cluster predictions will aid in facilitating the future investigation of novel *Aspergillus* SMs.

Genome-wide kinase knock-outs

The molecular genetic system of *A. nidulans* is powerful and technical advances in recent years have made

genome-wide, systematic approaches more feasible. The Fungal Genetics Stock Center (FGSC) provides a systematic gene deletion construct collection, a valuable experimental resource for the *A. nidulans* research community. De Souza et al. [19] have generated a set of gene deletion constructs for 9,851 genes, which represents 93.3 % of the encoding genome. Mutant strains generated with the cassettes are deposited with the FGSC after construction.

Using this deletion construct resource, a genome-wide kinase knock-out library consisting of deletion strains of most *A. nidulans* non-essential kinase genes was generated and deposited at the FGSC [19]. The kinase deletion strains were used for genome-wide functional analysis of kinases, resulting in identification of many previously unknown functions for kinases [19]. This kinase knock-out library was screened to test the hypothesis that manipulation of kinase expression has the potential to activate

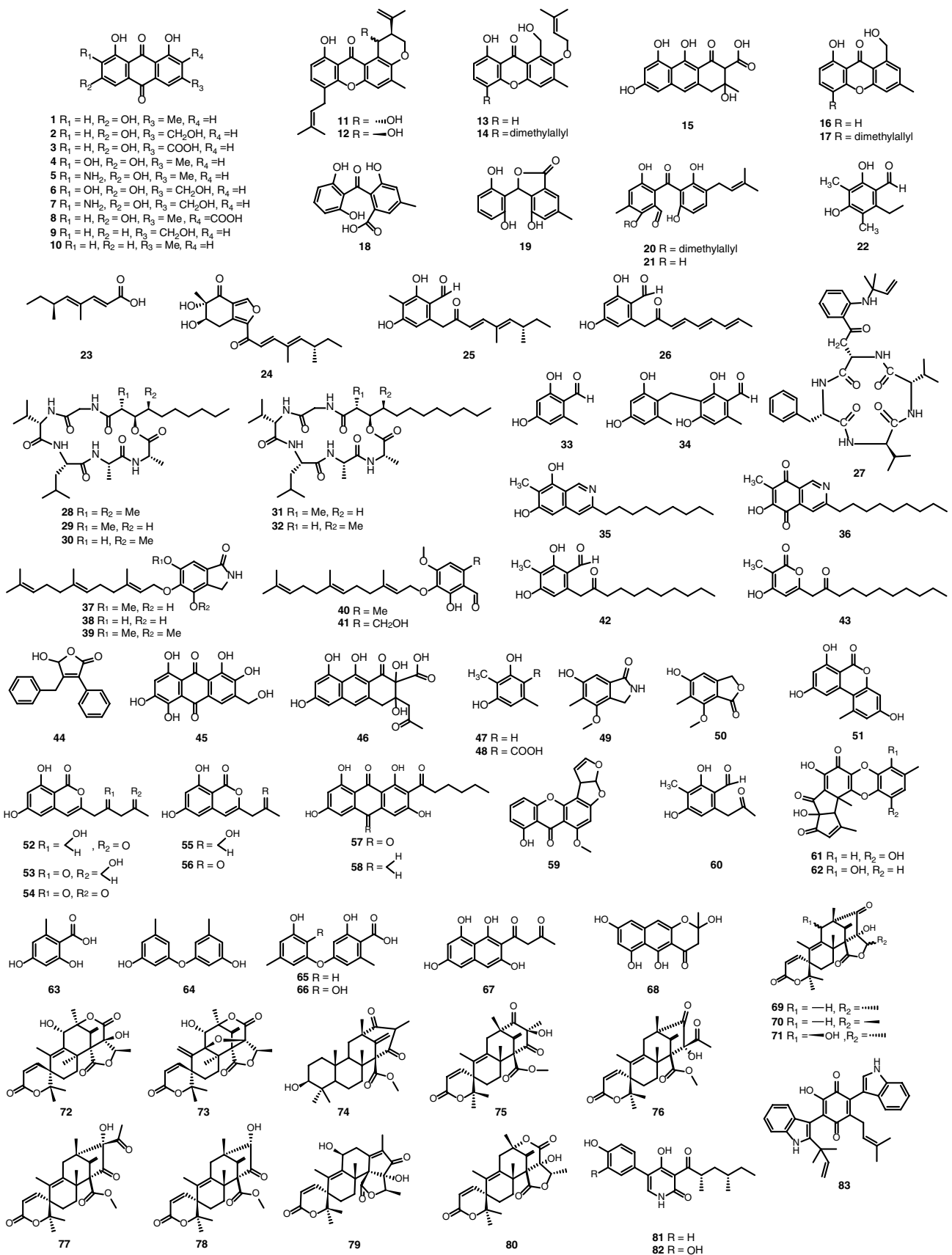


Fig. 1 Structures of compounds isolated from *A. nidulans*

silent SM gene clusters [58]. This led to the discovery of an *mpkA* knock-out strain that produced aspernidine A, a compound that had been discovered previously in *A. nidulans* [47] but the biosynthetic pathway remained unknown. The *mpkA* knock-out strain produced a sufficient amount of aspernidine A to allow the identification and analysis of the gene cluster involved in its biosynthesis. From the chemical structure of aspernidine A combined with previous data [1], it was predicted that a nonreducing polyketide synthase (NR-PKS) gene, *pkfA* (AN3230) is involved in the biosynthesis of aspernidine A. Deletion of *pkfA* confirmed this, and the boundary of the gene cluster was identified through a series of gene deletions of the surrounding genes of *pkfA*. Analysis of the SMs produced by *mpkA* deletion strains resulted in isolation and characterization of novel intermediates that aided in generating a proposed pathway for aspernidine A.

A similar deletion set of 28 protein phosphatase genes was generated and used to identify four essential phosphatases and four required for normal growth [50]. The deposited deletion constructs were also used in a study that identified multiple kinases and phosphatases involved in the sensing of carbon and energetic status, and also contributed to the understanding of the signaling cascades that result in regulation of CreA derepression and hydrolytic enzyme production [13].

Genome-wide analysis of all non-reduced polyketide synthases and NRPS-like enzymes in *A. nidulans*

Despite the success of various strategies to activate silent gene clusters, a large number of potential SM gene clusters remain untapped. To analyze clusters resistant to activation through existing approaches, a strategy was developed that completely bypasses normal regulation [1]. It takes advantage of recent advances in the construction of transforming fragments by fusion PCR and effective gene targeting to replace promoters of SM genes with the regulatable *alcA* promoter. It was applied to obtain a comprehensive understanding of the products of nonreducing polyketide synthase (NR-PKS) genes, a class of key genes of SM biosynthetic pathways [1]. The *A. nidulans* genome harbors 14 NR-PKS genes, and combined efforts by several groups over the years led to the identification of the chemical products of six of them [7, 12, 16, 17, 29, 42, 48, 52, 55, 62]. To determine the products of the remaining eight NR-PKS genes, the native promoters for each NR-PKS and other genes necessary for product formation or release were replaced with the *alcA* promoter. Induction of expression resulted in the production and release of compounds from each of the NR-PKS and allowed the completion of the determination of the products of NR-PKS genes of *A. nidulans*.

This approach can be applied to the discovery of other classes of SM biosynthetic gene clusters. This was demonstrated by systematically targeting nonribosomal peptide synthetase (NRPS)-like genes for promoter replacement, resulting in the discovery that one of the NRPS-like genes, *micA*, is the sole gene responsible for the biosynthesis of the metabolite microperfurane [59].

In another strategy carried out by Nielsen et al. [36], a genome-wide PKS deletion library was constructed by systematically deleting all 32 putative PKS genes. A reference strain was cultured on an array of culture media to find conditions that would induce production of SMs that were not previously linked to a gene cluster, and this was followed by screening of the genome-wide PKS deletion library to establish the genetic link to the SMs. This approach provided novel links between PKS genes and SMs, demonstrating its strength and the potential usefulness of the deletion library as a resource for further PKS studies.

Use of *A. nidulans* as a host for heterologous expression of SM genes from other *Aspergillus* species

The highly advanced and established molecular genetic system of *A. nidulans* can be applied to the study of SM production of other fungal species that have poor or non-existent molecular genetic systems [60]. Heterologous expression of fungal genes in other fungi has been used and with some success, but this approach is not without limitations including finding a suitable host and the difficulty of handling large genes and gene clusters. An advantage of fungal systems over bacterial for expressing fungal secondary metabolism genes is that fungi can correctly splice introns of secondary metabolism genes from other fungi resulting in successful expression [15, 22, 26]. Since many fungal SM genes are quite large and contain introns (often several introns) this is of considerable benefit.

Major advances have recently been made in establishing *A. nidulans* as a host for heterologous expression of fungal SMs. First, entire SM gene clusters have been deleted to eliminate production of unwanted *A. nidulans* SMs, resulting in reduced SM background and facilitating detection and isolation of compounds produced by the heterologously expressed genes [15].

Second, a system for transferring SM genes from other fungi while placing them under control of the *alcA* promoter has been developed [15, 33]. This system uses a strategy that involves (1) PCR amplification of each gene, (2) the use of fusion PCR to place each gene under control of the *alcA* promoter and to construct a transforming fragment, and (3) integration of the fragment into a target *A. nidulans* locus. For larger clusters several genes must be transferred into *A. nidulans* and, to avoid running out of

selectable markers for transformation, a marker recycling strategy was developed [15]. Each time a new gene is introduced into *A. nidulans* a selectable marker is evicted and this marker can be used in the subsequent transformation. This strategy allows an unlimited number of genes to be transferred into and expressed in *A. nidulans*. The use of this approach resulted in the successful expression of all six genes of the gene cluster that encodes the production of asperfuranone, a cryptic gene cluster from *A. terreus*. Furthermore, various combinations of expression genes were tested, leading to clarification of the asperfuranone biosynthetic pathway.

Another recent approach to transfer members of entire SM gene clusters is to assemble the PCR amplified individual cluster fragments into a single large transforming fragment using USER fusion, followed by insertion into the integration vector by USER cloning [37]. Using this technique, a total of 13 genes of a putative gene cluster responsible for geodin biosynthesis from *A. terreus* were transferred into *A. nidulans* in a two step process, successfully enabling geodin biosynthesis in *A. nidulans*.

Conclusions

Advances in genome sequencing in fungi have provided us with a wealth of information that suggests that the number of SM gene clusters far exceeds the number of discovered compounds. A combination of bioinformatics and experimental verification is fundamental to elucidating the SM biosynthetic pathways that these SM gene clusters encode. Among the many species of *Aspergillus*, *A. nidulans* is used as a model organism and it is the species with the most abundant literature by far and the most advanced, highly efficient molecular genetic system. Recent advances in development of prediction algorithms in *A. nidulans* and updated curation by AspGD have given us access to improved SM gene cluster predictions, which we can use as a basis for subsequent experimental verification. Advances in transforming fragment construction techniques and effective gene targeting expedite the experimental verification process. These advances, in combination, have enabled quick and systematic approaches to uncover the potential of SM production by *A. nidulans*. The application of these advances is not limited to the SMs of *A. nidulans*. Combined efforts such as the “1,000 Fungal Genomes Project (<http://1000.fungalgenomes.org/home/>)” by the DOE Joint Genome Institute (JGI) are dedicated to sequencing numerous different species of fungi and providing a database for the research community. Many of these fungi do not have good molecular genetic systems, which makes experimental verification a big challenge. Heterologous expression of fungal genes in other host fungi is one approach that is

being used, and major advances have been made to establish *A. nidulans* as a host. Newly developed methods in constructing transforming fragments and improved transformation strategies have made it possible for large or multiple genes to be transformed into *A. nidulans*. These approaches will contribute greatly to uncovering the untapped resources of SMs that the fungal genomes encode.

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