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*Medchemcomm*. Author manuscript; available in PMC 2013 November 13.

Published in final edited form as:

*Medchemcomm*. 2012 August 1; 3(8): . doi:10.1039/C2MD20055D.

## Identification and molecular genetic analysis of the cichorine gene cluster in *Aspergillus nidulans*

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### Abstract

We recently demonstrated that the phytotoxin cichorine is produced by *Aspergillus nidulans*. Through a set of targeted deletions, we have found a cluster of seven genes that are required for its biosynthesis. Two of the deletions yielded molecules that give information about the biosynthesis of this metabolite.

### Introduction

Filamentous fungi produce a variety of low-molecular weight secondary metabolites, many of which have been proven to possess remarkable biological activities. They include the cholesterol-lowering agent lovastatin and the immunosuppressant cyclosporine.<sup>1</sup> It came as a surprise, then, that the analysis of recently sequenced fungal genomes suggested that the number of metabolites obtained in laboratory conditions is still far fewer than the number of putative secondary metabolite genes.<sup>2-5</sup>

This realization has prompted efforts to activate these silent genes through various approaches, including epigenetic modification,<sup>6</sup> promoter exchange with an inducible promoter,<sup>7</sup> controlled expression of a gene coding for a pathway-specific transcription factor,<sup>8</sup> and co-cultivation with bacteria.<sup>9</sup> A different strategy, however, is to alter the media conditions in which the fungus is cultivated, as different environmental cues may switch on heretofore unactivated pathways.<sup>10</sup> For the filamentous fungus *Aspergillus nidulans*, this approach has been key to the isolation of aspoquinolones,<sup>11</sup> aspermidine A/B,<sup>12</sup> and F9775 A and B.<sup>13</sup>

With this approach we obtained the polyketide cichorine from the cultivation of *Aspergillus nidulans* on YES (Yeast Extract Sucrose) plates at 37°C for 5 days, during the course of our work with prenylated xanthenes from this species.<sup>14</sup> Cichorine is a phytotoxin active against knapweed, corn, and soybeans.<sup>15-16</sup> It was isolated from *Aspergillus silvaticus* as well as *Alternaria cichorii*, which produces foliar blight in the important pest Russian knapweed.<sup>15</sup> It had not before been observed in *A. nidulans*. The compact, functionalized isoindolin-1-one framework was an attractive target for total synthesis.<sup>17-18</sup> Compounds featuring this framework have been shown to possess antimicrobial,<sup>19</sup> anti-HIV,<sup>20</sup> and antitumor properties.<sup>21</sup>

Because the genome of *A. nidulans* has been sequenced and relatively well-annotated,<sup>2</sup> and because a straightforward gene targeting system has been developed for this species,<sup>22-23</sup> we could proceed to efficiently investigate the genes that are responsible for the formation of cichorine. The presence of an aromatic group in cichorine indicated that it is a product of a nonreduced polyketide synthase (NR-PKS). Targeted gene deletions revealed that AN6448.4 alone failed to produce any detectable amount of cichorine. (We use the gene nomenclature of the Central Aspergillus Data Repository, CADRE, <http://www.cadre-genomes.org.uk/>, and the Aspergillus genome database, <http://www.aspgd.org/>.)

In our current investigation, we endeavored to identify other genes that are responsible for cichorine formation anticipating that intermediate compounds accumulating as a result of interruption of the pathway would help us understand how cichorine is synthesized.

## Results and discussion

Genes required for the synthesis of a particular secondary metabolite tend to cluster together in the genome in fungi. For this reason we deleted genes both upstream and downstream of the identified PKS gene, AN6448.4 (Table 1). All strains carried the deletion of *stcJ*, a gene responsible for the carcinogenic secondary metabolite sterigmatocystin.<sup>24</sup> Eliminating *stcJ* may facilitate isolation of other metabolites and also free up precursors to boost their production. The strains also carried *nkuA* to improve frequencies of correct gene targeting.<sup>22</sup>

All strains were cultivated in the same cultivation medium that yields cichorine. (Cichorine has also been generated via cultivation in Raulin-Thom media.<sup>25</sup>) All deletions were verified by diagnostic PCR (see Experimental section). The result was the elimination of cichorine in strains ranging from AN6443.4 to AN6449.4, with the exception of AN11922.4 (Figure 1). Table 2 lists the putative functions of the genes within the cluster and the ones immediately outside it. We now designate these genes as *cicA-cicH*. Aside from the PKS gene, the cluster contains a transporter (*cicA*) and transcriptional activator gene (*cicD*), and also four tailoring genes (*cicB*, *cicC*, *cicE*, and *cicH*).

Initially, we were unable to detect any obvious biosynthetic intermediates from the knockout strains. This was perhaps to be expected for the transporter, transcriptional activator, and backbone PKS genes, but it was reasonable to predict that at least some of the tailoring genes would yield intermediates upon deletion. Some polyketides feature a carboxylic acid motif or otherwise become negatively charged in aqueous media. It stands to reason, then, that these molecules will not partition into the organic layer during extraction. In these cases, it is necessary to neutralize the molecules with concentrated acid. This strategy was necessary for the acquisition of the polyketides orsellinic acid and F9775 A and B in *A. nidulans*.<sup>13</sup>

Acidifying the aqueous media allowed us to detect one compound from the *cicB* strain and an additional compound from the *cicC* strain. To determine their structures, we scaled up the cultivation and purified the metabolites using silica gel chromatography followed by preparative HPLC. Chemical structures were determined by <sup>1</sup>H and <sup>13</sup>C NMR, and the spectra were also compared with literature data.

The *cicB* intermediate is a simple polyketide, **5**, 3-methylorsellinic acid (Figure 2). Interestingly, this is the same molecule that we have recently acquired through the upregulation of the PKS gene, *cicF*. The *cicC* intermediate, nidulol, **6**, has been isolated from *Aspergillus* and *Alternaria* species, including a strain of *A. nidulans*, and it was found to be mildly cytotoxic toward human epidermoid carcinoma KB cells.<sup>26</sup>

The deletants and the intermediates allow a better understanding of cichorine biosynthesis. Because upregulation of the backbone PKS gene *cicF* and deletion of *cicB* yield the same molecule, it is reasonable to suspect that the first step in cichorine biosynthesis is the generation of **5**, followed by formation of the next (unidentified) intermediate catalyzed by the gene product of *cicB*. Interestingly, the cytochrome P450 monooxygenase gene *cicH* is homologous to *mpaD* in *Penicillium brevicompactum*, which is predicted to catalyze the lactonization of the well-known immunosuppressant mycophenolic acid. Therefore, *cicH* may play a similar ring-closing role for cichorine.

The O-methyltransferase gene, *cicE*, is plausibly the responsible gene for the methylation of one of the phenol groups. Functionalized, two-ring **6** from the *cicC* strain may be a later-stage intermediate, but lactone to lactam formation must still occur in order to generate cichorine. Other genes would be required for this conversion, and these genes are not found within this cluster. Based on this information, we have outlined a hypothetical biosynthesis as depicted in Figure 3. Although the entire set of genes pertaining to a particular fungal secondary metabolite may fit in one single cluster, it is being increasingly found in *A. nidulans* that such genes may be split into at least two distinct genomic loci, demonstrated with the prenylated xanthenes<sup>14</sup> and the terpenes<sup>27</sup> from *A. nidulans*, and possibly with F9775 A and B.<sup>13</sup>

## Conclusion

In summary we have shown that genes from a distinct cluster are required to generate cichorine in *A. nidulans*. The cluster contains the PKS gene, one regulatory and transporter gene each, and four genes involved in the tailoring of the polyketide backbone. We have also acquired two intermediates, improving our understanding of the biosynthesis of this isoindolin-1-one-based molecule.

## Experimental

### Generation of fusion PCR fragments *A. nidulans* protoplasting, and transformation

The gene deletions were performed using established gene targeting procedures.<sup>23</sup> Two ~1000 base pair fragments upstream and downstream of every targeted gene were amplified from *A. nidulans* genomic DNA using PCR. Primers used in this study are listed in Table S1. The two amplified flanking sequences and an *A. fumigatus pyrG* selectable marker cassette were fused together by PCR using nested primers. *A. nidulans* strains in this study are listed in Table 1. Protoplast generation and transformation were utilized as previously described.<sup>23</sup> The strain LO2026 carrying a deletion of the *stcJ* gene that eliminates sterigmatocystin production was used as the recipient strain. Diagnostic PCR of the deletant strains was performed employing the external primers from the first round of PCR. The difference in the size between the gene replaced by the selective marker and the native gene enabled us to determine if the transformants carried correct gene replacements. For further verification, diagnostic PCR was performed two more times, with one of the external primers and a primer located inside the marker gene, then the other external primer and an internal primer. In these cases, the deletants yielded the PCR product of the expected size whereas no product was present in non-deletants.

### Fermentation and LC/MS analysis

YES medium was prepared by combining 20 g yeast extract, 120 g sucrose, 20 g agar, and 2 mL trace element solution in 1 L H<sub>2</sub>O. For the LC/MS screening experiments, spores of LO2026 (the control strain) and three strains of each gene deletant were individually inoculated ( $1 \times 10^7$  spores) onto  $10 \times 150$  mm petri dishes which contained YES agar, and

they were cultivated for five days at 37°C. Following, the agar was split into ~2 cm<sup>2</sup> pieces, and the material was placed into 250 mL Ehrlenmeyer flasks and covered with methanol. The flasks were placed in a sonicator (Branson, Model 5510) for an hour. The methanol was then decanted. The pieces were covered with 1:1 methanol:dichloromethane, followed by another hour of sonication. The combined organic solvents were removed *in vacuo*, then partitioned between H<sub>2</sub>O (25 mL) and ethyl acetate (25 mL × 2). The combined ethyl acetate layers were evaporated. The dried, crude material was dissolved at a concentration of 20 mg/mL in DMSO, and then a sample was diluted 5-fold in methanol for LC/MS analysis.

LC/MS was performed using a ThermoFinnigan LCQ Advantage ion trap mass spectrometer with an RP C18 column (Alltech Prevail; 2.1 × 100 mm with a 3 μm particle size) at a flow rate of 125 μL/min and monitored by a UV detector at 254 nm. The solvent gradient was 95% MeCN-H<sub>2</sub>O (solvent B) in 5% MeCN-H<sub>2</sub>O (solvent A) both containing 0.05% formic acid: 0% B from 0 to 5 min, 0 to 100% B from 5 to 35 min, 100% B from 35 to 40 min, 100% B to 0% B from 40 to 45 min, and reequilibration with 0% B from 45 to 50 min.

### Isolation of metabolites

The LO2026 (*stcJ*), LO3387 (*cicB*), and LO3337 (*cicC*) strains were each cultivated in 25 × 150 mm petri dishes containing YES medium for 5 days at 37°C. As with LC/MS analysis, the agar was chopped and sonicated in methanol, then 1:1 methanol:dichloromethane. The organic material was evaporated and extracted 4x with an equal volume of ethyl acetate. For LO3387 and LO3337, the aqueous layer was acidified with concentrated HCl to a pH of 2. The crude material was subjected to silica gel column chromatography, using 98:2 dichloromethane:methanol as the eluent for **1** and **5**, and 100% dichloromethane for **6**. The materials were further separated by preparative HPLC [Phenomenex Luna 5 μm C18 (2), 250 × 21.2 mm] with a flow rate of 5.0 mL/min and measured by a UV detector at 250 nm. See Supplementary Information for more details about isolation.

**Cichorine (1)**—white powder; <sup>1</sup>H NMR and <sup>13</sup>C NMR data (DMSO-*d*<sub>6</sub>), in good agreement with the published data.<sup>17,18</sup><sup>1</sup>H NMR (DMSO-*d*<sub>6</sub>): = 2.04 (3H, s), 3.83 (3H, s), 4.39 (2H, s), 6.81 (1H, s), 7.40 (1H, br s), 8.42 (1H, s), 9.76 (1H, s); <sup>13</sup>C NMR (DMSO-*d*<sub>6</sub>): = 10.1, 43.9, 59.5, 103.7, 119.7, 123.8, 132.7, 154.2, 157.1, 170.6. For UV-Vis and ESIMS spectra, see Fig. S1. *m/z* (M+H) = 194.4; calculated = 194.1.

**3-methylorsellinic acid (5)**—white powder; <sup>1</sup>H NMR and <sup>13</sup>C NMR data (acetone-*d*<sub>6</sub>), in good agreement with the published data.<sup>29</sup><sup>1</sup>H NMR (acetone-*d*<sub>6</sub>): = 2.03 (3H, s), 2.48 (3H, s), 6.35 (1H, s); <sup>13</sup>C NMR (acetone-*d*<sub>6</sub>): = 7.4, 23.6, 107.4, 108.6, 110.6, 140.5, 160.3, 174.2. For UV-Vis and ESIMS spectra, see Fig. S1. *m/z* (M-H) = 181.2; calculated = 181.1.

**Nidulol (6)**—white powder; <sup>1</sup>H NMR and <sup>13</sup>C NMR data (acetone-*d*<sub>6</sub>), in good agreement with the published data.<sup>30</sup><sup>1</sup>H NMR (acetone-*d*<sub>6</sub>): = 2.11 (3H, s), 3.98 (3H, s), 5.15 (2H, s), 6.87 (1H, s); <sup>13</sup>C NMR (acetone-*d*<sub>6</sub>): = 8.0, 61.3, 68.5, 103.6, 108.2, 118.1, 148.8, 158.4, 163.3, 168.6. For UV-Vis and ESIMS spectra, see Fig. S1. *m/z* (M-H) = 193.6; calculated = 193.1.

### Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

## Acknowledgments

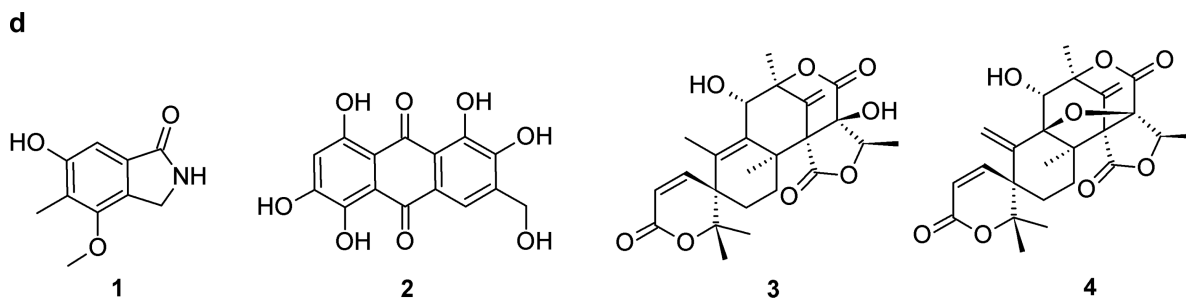
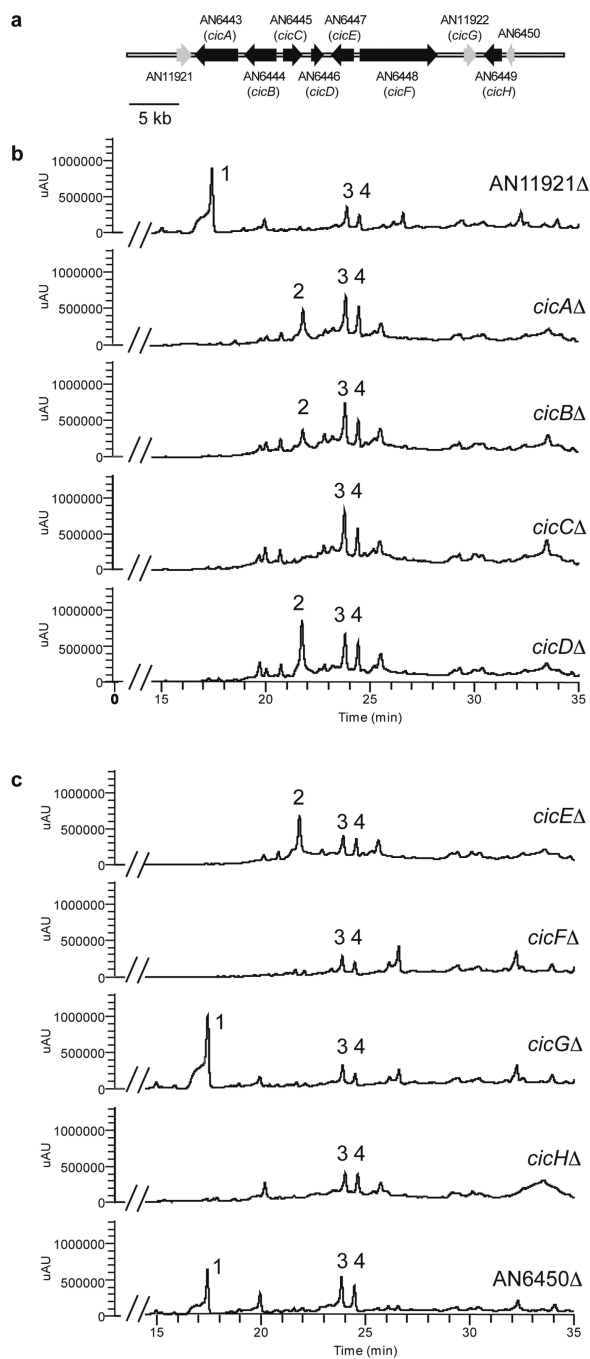
The work was supported by grant PO1GM084077 from the National Institute of General Medical Sciences to BRO and CCCW and by the Kansas University Endowment Fund to BRO.

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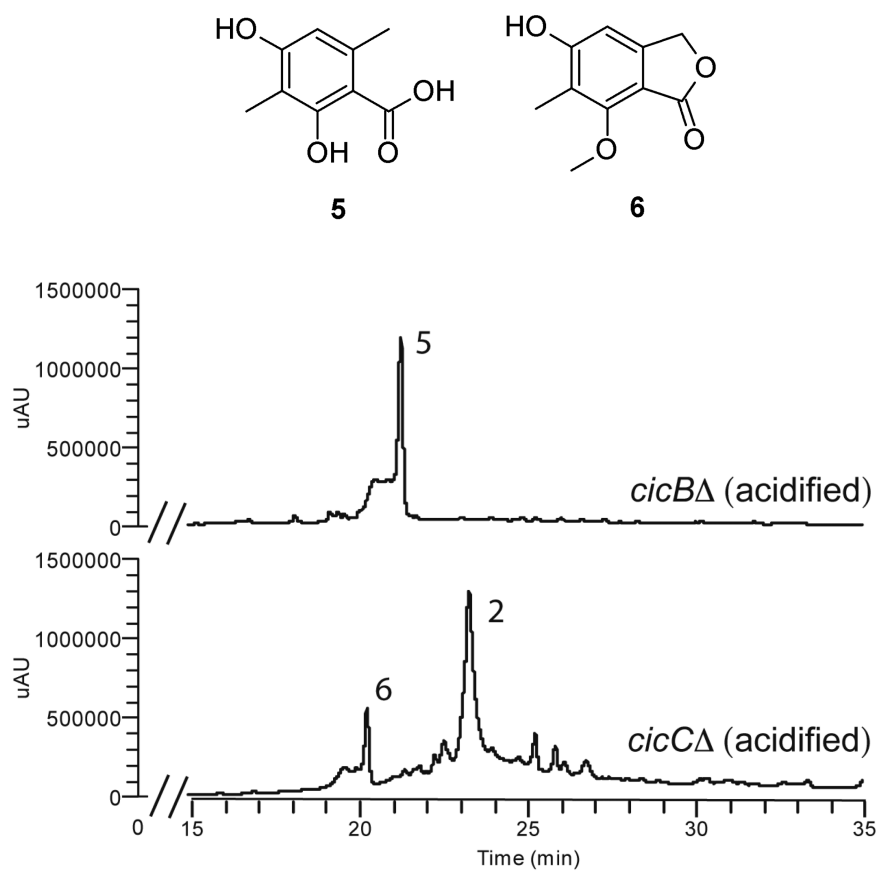




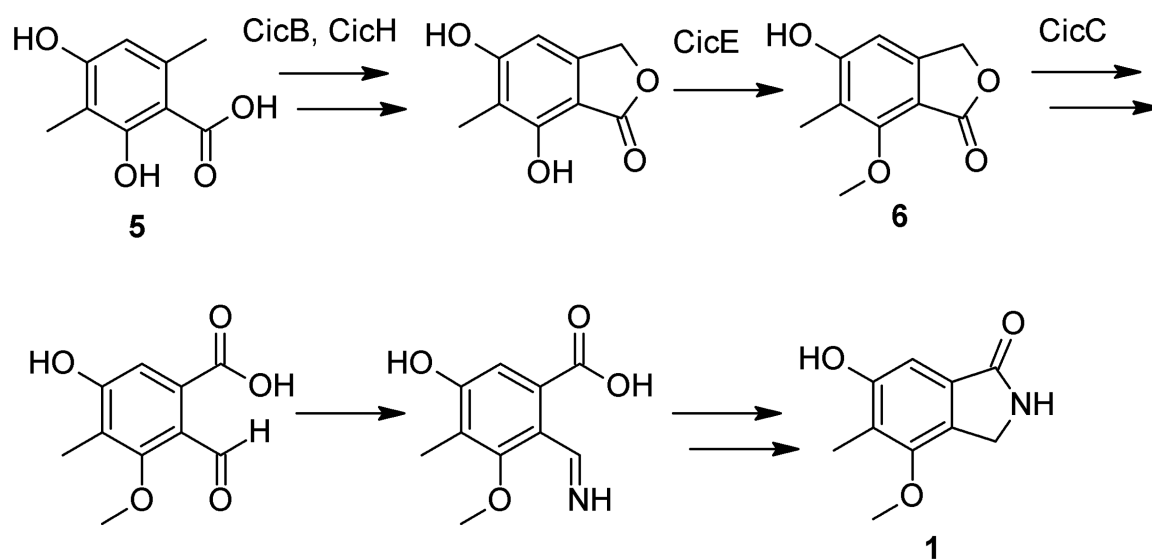
**Figure 1.**

(A) Organization of the genes surrounding the PKS *cicF* involved in cichorine biosynthesis. Genes shown in black are required for cichorine biosynthesis, whereas those in gray are not. (B) HPLC extracts of strains AN11921 to *cicD* as detected by UV absorbance at 254 nm. (C) The extracts (as in B) of *cicE* to AN6450. (D) Structures of cichorine (**1**), asperthecin (**2**), austinol (**3**), and dihydroaustinol (**4**). The biochemical origins of **2-4**, differing from cichorine, have been previously determined.<sup>27-28</sup>





**Figure 2.** 3-methylorsellinic acid (5) and nidulol (6), intermediates acquired from acidified extracts of cichorine deletant strains *cicB* and *cicC*, respectively.



**Figure 3.**  
Proposed biosynthesis of cichorine (1).

**Table 1***A. nidulans* strains used in this study

| Strain                        | Genotype  | Source     |
|-------------------------------|---|------------|
| <i>stcJ</i><br>LO2026         | <i>pyrG89; pyroA4, nkuA::argB; riboB2, stcJ::AfriboB</i>                  | 6          |
| AN11921<br>LO5000-LO5002      | <i>pyrG89; pyroA4, nkuA::argB; riboB2, stcJ::AfriboB; AN11921::AfpyrG</i> | This study |
| <i>cicA</i><br>LO4517-LO4519  | <i>pyrG89; pyroA4, nkuA::argB; riboB2, stcJ::AfriboB; cicA::AfpyrG</i>    | This study |
| <i>cicB</i><br>LO4482-LO4484  | <i>pyrG89; pyroA4, nkuA::argB; riboB2, stcJ::AfriboB; cicB::AfpyrG</i>    | This study |
| <i>cicC</i><br>LO4487-LO4488  | <i>pyrG89; pyroA4, nkuA::argB; riboB2, stcJ::AfriboB; cicC::AfpyrG</i>    | This study |
| <i>cicD</i><br>LO4492, LO4494 | <i>pyrG89; pyroA4, nkuA::argB; riboB2, stcJ::AfriboB; cicD::AfpyrG</i>    | This study |
| <i>cicE</i><br>LO4497-LO4499  | <i>pyrG89; pyroA4, nkuA::argB; riboB2, stcJ::AfriboB; cicE::AfpyrG</i>    | This study |
| <i>cicF</i><br>LO4503-LO4505  | <i>pyrG89; pyroA4, nkuA::argB; riboB2, stcJ::AfriboB; cicF::AfpyrG</i>    | This study |
| <i>cicG</i><br>LO4568-LO4569  | <i>pyrG89; pyroA4, nkuA::argB; riboB2, stcJ::AfriboB; cicG::AfpyrG</i>    | This study |
| <i>cicH</i><br>LO4507-LO4508  | <i>pyrG89; pyroA4, nkuA::argB; riboB2, stcJ::AfriboB; cicH::AfpyrG</i>    | This study |
| AN6450<br>LO4975-LO4977       | <i>pyrG89; pyroA4, nkuA::argB; riboB2, stcJ::AfriboB; AN6450::AfpyrG</i>  | This study |

**Table 2**

Putative functions of the genes within the cichorine cluster, as indicated from BLAST searches.

| <b>Gene designation</b> | <b>Putative function</b>       | <b>CADRE/AspGD annotation</b> | <b>Broad annotation</b> |
|-------------------------|--------------------------------|-------------------------------|-------------------------|
| <b>AN11921</b>          | Amino acid transporter         | AN11921.4                     | ANID_11921.1            |
| <i>cicA</i>             | ABC transporter                | AN6443.4                      | ANID_6443.1             |
| <i>cicB</i>             | Conserved hypothetical protein | AN6444.4                      | ANID_6444.1             |
| <i>cicC</i>             | Oxidoreductase                 | AN6445.4                      | ANID_6445.1             |
| <i>cicD</i>             | Regulatory protein             | AN6446.4                      | ANID_6446.1             |
| <i>cicE</i>             | O-methyltransferase            | AN6447.4                      | ANID_6447.1             |
| <i>cicF</i>             | Polyketide synthase            | AN6448.4                      | ANID_6448.1             |
| <i>cicG</i>             | Conserved hypothetical protein | AN11922.4                     | ANID_11922.1            |
| <i>cicH</i>             | Cytochrome P450                | AN6449.4                      | ANID_6449.1             |
| <b>AN6450</b>           | Ribosomal protein              | AN6450.4                      | ANID_6450.1             |