Aspergillus fumigatus ffmA Encodes a C2H2-Containing Transcriptional Regulator That Modulates Azole Resistance and Is Required for Normal Growth

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ABSTRACT The production of a collection of deletion mutant strains corresponding to a large number of transcription factors from the filamentous fungal pathogen Aspergillus fumigatus has permitted rapid identification of transcriptional regulators involved in a range of different processes. Here, we characterize a gene designated ffmA (favors fermentative metabolism) as a C2H2-containing transcription factor that is required for azole drug resistance and normal growth. Loss of ffmA caused cells to exhibit significant defects in growth, either under untreated or azole-challenged conditions. Loss of FfmA caused a reduction in expression of the AbcG1 ATP-binding cassette transporter, previously shown to contribute to azole resistance. Strikingly, overproduction of the AtrR transcription factor gene restored a wild-type growth phenotype to an ffmAΔ strain. Overexpression of AtrR also suppressed the defect in AbcG1 expression caused by loss of FfmA. Replacement of the ffmA promoter with a doxycycline-repressible promoter restored nearly normal growth in the absence of doxycycline. Finally, chromatin immunoprecipitation experiments indicated that FfmA bound to its own promoter as well as to the abcG1 promoter. These data imply that FfmA and AtrR interact both with respect to abcG1 expression and also more broadly to regulate hyphal growth.

IMPORTANCE Infections associated with azole-resistant forms of the primary human pathogen Aspergillus fumigatus are associated with poor outcomes in patient populations. This makes analysis of the mechanisms underlying azole resistance of A. fumigatus a high priority. In this work, we describe characterization of a gene designated ffmA that encodes a sequence-specific transcriptional regulator. We identified ffmA in a screen of a collection of gene deletion mutant strains made in A. fumigatus. Loss of ffmA caused sensitivity to azole drugs and also a large reduction in normal growth. We found that overproduction of the AtrR transcription factor could restore growth to ffmA null cells. We provide evidence that FfmA can recognize promoters of genes involved in azole resistance as well as the ffmA promoter itself. Our data indicate that FfmA and AtrR interact to support azole resistance and normal growth.

KEYWORDS Aspergillus fumigatus, DNA-binding proteins, atrR, azole resistance, ffmA, transcription factors
consists of a compound mutation corresponding to a duplication of a 34-bp region in the {cyp51A} promoter along with a single-amino-acid substitution in the encoded enzyme (4). This allele is referred to as TR34 (34-bp promoter duplication) L98H {cyp51A}. Together, these two alterations cause a dramatic increase in voriconazole MIC. Infections associated with this {cyp51A} allele are associated with a significantly worse outcome in particular populations (5) and make understanding regulation of {cyp51A} in particular and azole resistance in general of high priority in this pathogen.

While mutations associated with the {cyp51A} gene are found with high frequency in azole-resistant {A. fumigatus} isolates, more recent studies have demonstrated that not all azole resistance in this organism can be explained by mutations at {cyp51A} (reviewed in references 6 and 7). Mutations in the gene encoding HMG coenzyme A (CoA) reductase have been described that lead to strong decreases in voriconazole susceptibility with a wild-type {cyp51A} locus present (8, 9). Additionally, an alteration in the {hapE} gene encoding a subunit of the CCAAT-binding complex also produced a strain with an elevated voriconazole MIC (10). Loss of normal {hapE} function leads to a large increase in {cyp51A} expression with associated elevation of azole resistance (11). Coupled with the well-described increase in {cyp51A} expression caused by the presence of the TR34 promoter (11, 12), these data illustrate the critical contribution of transcriptional regulation to azole resistance.

An additional non-{cyp51A} mode of azole resistance is illustrated by the overproduction of the ATP-binding cassette (ABC) transporter AbcG1 (also known as Cdr1B/abcC). Clinical isolates have been described in which elevated transcription of the gene encoding this membrane protein leads to enhanced azole resistance (13). The presence of the {abcG1} gene is required for the normal increase in voriconazole resistance seen in strains containing the TR34 L98H form of {cyp51A}, demonstrating the role of this ABC transporter even in the presence of a strongly resistant allele of {cyp51A} (14). We and others have found a transcription factor designated AtrR (ABC transporter regulator) that is required for high-level transcription of both {abcG1} and {cyp51A} (15, 16). Transcriptional control of {cyp51A} is a key contributor to azole resistance, but much remains to be uncovered concerning the range of transcriptional inputs to drug resistance in {A. fumigatus}.

We have previously described a collection of deletion mutants in genes that encode transcription factors (17). The first large-scale screening of this mutant collection employed itraconazole and led to the identification of the negative transcriptional regulator nctA-B. This mutant library also confirmed that deletion mutants in either the sterol gene regulator srbA (18) or atrR caused large increases in itraconazole susceptibility. We counterscreened these itraconazole-susceptible mutant strains for effects on expression of the AbcG1 ABC transporter using an antibody directed against this membrane protein (14). A poorly characterized gene called rleC (recently renamed {ffmA}) (19) was identified that caused a large decrease in AbcG1 expression. The {ffmA} gene encodes a protein containing a C_{2}H_{2}-containing sequence-specific DNA-binding domain (20). We also found that {ffmA} null mutants have a large defect in normal growth along with the predicted azole hypersensitivity. Interestingly, overexpression of the AtrR transcription factor was able to suppress both the growth defect of an {ffmA} null mutant and restore AbcG1 expression. These data indicate the presence of a genetic interaction between AtrR and FfmA.

RESULTS

Screening the transcription factor deletion mutant library for genes that affect azole resistance. Using a previously described transcription factor deletion mutant library (17), we identified additional mutant strains that exhibited increased susceptibility to itraconazole. These individual strains were grown from each library isolate, whole-cell protein extracts were prepared, and these were analyzed by Western blotting using an anti-AbcG1 antiserum (21). The results of this Western blot comparison are shown in Fig. 1.

We found clear and reproducible reduction in the level of AbcG1 in strains lacking either the atrR or AFUB_026340 (Afu2g10550) gene. The latter gene has recently been
characterized as a locus enhancing expression of genes involved in fermentation (favors fermentative metabolism, ffmA [19]), and we will use this designation here. The ffmA gene has also been characterized as rfeC (regulator of FLO11) on the basis of activating the FLO11 promoter when expressed as a cDNA clone in Saccharomyces cerevisiae (22). We selected ffmA for further study to examine its role as a regulator of abcG1 expression.

Phenotypes caused by loss of the ffmA gene. To confirm the effects of the ffmA gene on phenotypes, we prepared a new ffmAΔ allele in a different genetic background (AfS35). We also used the original deletion mutant library strain for comparison. Spores were prepared from the two wild-type strains and their isogenic ffmAΔ derivatives. A radial growth assay was performed for these 4 different strains (Fig. 2A).

Irrespective of the genetic background employed, the clearest effect caused by loss of ffmA is a profound growth defect. We estimate the growth rate of the ffmAΔ strain to be roughly 25% that of the wild-type cells. This was confirmed by quantitation of a radial growth assay (see Fig. S1 in the supplemental material). This defect in growth rate complicated the analysis ofazole susceptibility in the ffmAΔ strain. We also observed the accumulation of an orange/brown pigmentation upon loss of ffmA in both strain backgrounds. To compare the voriconazole susceptibility phenotypes of these strains, we used a zone-of-inhibition assay and placed a filter disk containing this azole drug in the center of the plate of spores from each wild-type and isogenic ffmAΔ strain. These plates were incubated at 37°C and then photographed (Fig. 2B).

Loss of ffmA from the A1160 wild-type strain caused a reproducible increased susceptibility to voriconazole, but this was masked by the extreme growth defect seen in the AfS35 genetic background. The ffmAΔ derivative of AfS35 appeared to be more seriously growth compromised than its A1160 ffmAΔ counterpart.
Since loss of the \( ffmA \) gene caused such a pronounced defect in growth, we examined the effect of placing \( ffmA \) under the control of a strong promoter. We hypothesized that overproduction of FfmA caused by replacement of the \( ffmA \) promoter with the corresponding region from the highly expressed \( hspA \) (AFUA_1G07440, Hsp70 family chaperone) gene might affect voriconazole resistance. Using CRISPR/cas9-mediated recombination, the \( hspA \) promoter was inserted in place of the native \( ffmA \) promoter region. Appropriate transformants were recovered and analyzed for several \( ffmA \)-dependent phenotypes.

The presence of the \( hspA-ffmA \) fusion gene reduced radial growth to \( \sim 35\% \) of normal at \( 37^\circ C \) (Fig. 3A), similar to that seen for the \( ffmA\Delta \) strain. This growth defect likely contributed to the lack of any effect seen on voriconazole susceptibility.

To confirm that use of the \( hspA \) promoter to drive \( ffmA \) expression led to overproduction of FfmA, we assessed protein levels of FfmA using Western blotting. To
To accomplish this, a rabbit polyclonal antiserum directed against bacterially expressed FfmA was prepared. Isogenic wild-type and hspA-ffmA strains were grown with or without voriconazole treatment, and levels of FfmA were analyzed by Western blotting.

As shown on the left side of Fig. 3B, treatment of wild-type cells with voriconazole led to a 2-fold increase in FfmA levels compared to untreated cells. The presence of the hspA-ffmA allele caused a 2-fold increase in FfmA levels compared to the wild-type strain, consistent with a higher transcriptional level supported by the hspA promoter as we have seen before (23). Interestingly, exposure of hspA-ffmA cells to voriconazole caused a further increase in FfmA induction to roughly 5-fold above the non-azole-treated wild-type levels.
Our finding that loss of ffmA led to a decrease in AbcG1 expression (Fig. 1) prompted us to examine the effect of hspa-ffmA on expression of both AbcG1 and the AtrR transcription factor that is key to the regulation of this ABC transporter-encoding gene (15, 16). The same protein extracts analyzed in Fig. 3B were evaluated for the level of AbcG1 and AtrR in Fig. 3C using appropriate antisera. Expression of AtrR was increased in the hspa-ffmA strain compared to the wild type by approximately 50%, with a further increase to nearly 2-fold when the hspa-ffmA cells were challenged with voriconazole. Expression of the ABC transporter AbcG1 was seen to increase by 3-fold in hspa-ffmA cells in the presence of voriconazole compared to equivalently treated wild-type cells. These data further support the view that AbcG1 expression responds to changes in ffmA expression and suggest that the atrR gene is another target of FfmA.

**Overproduction of AtrR can suppress ffmA-dependent phenotypes.** Our finding of this potential link between AtrR and FfmA led us to examine the interaction between the two genes encoding these transcriptional regulators. To test the epistasis of atrR and ffmA, we used a hypermorphic form of atrR that we characterized earlier: the hspa-atrR fusion gene. This allele of atrR was found to cause overproduction of abcg1 along with other AtrR-regulated target genes (16). We introduced the ffmAΔ null allele into a strain containing the hspa-atrR fusion at the normal atrR locus. Transformants were recovered and confirmed by PCR analyses.

The most obvious phenotype seen was the striking suppression of the growth defect of an ffmAΔ strain (Fig. 4A). While the radial growth assay demonstrated that the hspa-atrR allele restored wild-type levels of growth to the ffmAΔ strains, production of orange pigment was retained, indicating that this ffmAΔ phenotype was not suppressed. This indicates that while there is important overlap between FfmA- and AtrR-dependent phenotypes, this overlap is not universal between these factors.

To examine the ability of the hspa-atrR fusion to suppress the defect in AbcG1 expression caused by the hspaΔ lesion, we assessed levels of AbcG1 protein in these strains by Western blotting (Fig. 4B). Loss of ffmA decreased AbcG1 expression in an otherwise wild-type background, while the atrRΔ null allele reduced AbcG1 expression even further and there was no detectable immunoreactivity in an abcg1Δ strain. Importantly, the hspa-atrR allele could restore AbcG1 expression to ~80% of wild-type levels in an ffmAΔ null strain. Overexpression of AtrR from the hspa promoter was sufficient to restore near-normal growth and parental levels of AbcG1 expression to a strain lacking ffmAΔ.

**Doxycline-dependent expression of ffmA.** Both loss of ffmA and overproduction of FfmA from the hspa promoter caused a serious defect in normal growth along with reduction of both AbcG1 expression and voriconazole resistance. To develop a strain that could be grown and tested in a controllable manner, we constructed a doxycline-repressible form of ffmA. This was accomplished by integrating a doxycycline-repressible (dox-off; DO) promoter into the ffmA locus to replace the wild-type promoter region. This DO promoter was integrated either immediately upstream of the wild-type ffmA (DO-ffmA) or with a single Flag epitope fused to the N terminus of FfmA (DO-Flag-ffmA). To confirm that the presence of the DO promoter did not cause a severe growth defect in the absence of doxycycline, appropriate transformants were placed on medium containing or lacking this compound and allowed to grow at 37°C.

The presence of either DO allele of ffmA led to nearly normal growth in the absence of doxycycline (Fig. 5A). The growth rate of the DO allele-containing strains was estimated to be 80% of the wild-type strain in the absence of doxycycline. The addition of 25 mg/liter doxycycline caused a reduced growth phenotype to 25% of the wild-type level, resembling that of the ffmaΔ or hspa-ffmA strain analyzed above.

We next used the doxycycline-repressible nature of the DO-ffmA and DO-Flag-ffmA genes to determine the effect of acutely repressing FfmA production on AbcG1. Both DO promoter-driven forms of ffmA were grown for 8 h and then either left untreated or had doxycycline added to the culture. After 12 additional hours of growth, whole-cell protein extracts were prepared and analyzed by Western blotting using anti-FfmA or anti-AbcG1 antisera (Fig. 5C).
Both DO-driven forms of *ffmA* produced more FfmA protein than was observed in wild-type cells. FfmA levels were elevated to 3.6-fold higher than wild-type levels in the DO-*ffmA* fusion, while the DO-Flag-*ffmA* fusion gene produced 50% more FfmA protein than wild-type cells. Surprisingly, AbcG1 expression was lowered in the presence of both DO-*ffmA* alleles when doxycycline was absent. The DO-*ffmA*-containing strain produced only 20% of normal AbcG1 protein, while the DO-Flag-*ffmA* fusion led to production of 70% of wild-type AbcG1 levels. Both DO fusion genes were strongly repressed by the addition of doxycycline to the medium, with both FfmA and AbcG1 levels nearly undetectable. These data support the interpretation that loss of AbcG1 expression and the growth defect caused by loss of *ffmA* represent an authentic consequence of the absence of FfmA and are not an indirect effect caused by response to the growth defect elicited by FfmA loss.

**FfmA binds to the *ffmA* and *abcG1* promoters in vivo.** The presence of a C$_2$H$_2$ DNA-binding domain in the FfmA protein sequence as well as its effect on *abcG1* and *atr* expression is consistent with this protein playing a role as a transcription factor. To determine if FfmA is directly associating with target gene promoters, we carried out single-gene chromatin immunoprecipitation (ChIP) experiments using either the DO-Flag-*ffmA* fusion gene or wild-type cells. We used anti-Flag antibody to enable ChIP in the DO-Flag-*FfmA* strain and anti-FfmA in wild-type cells. Fixed chromatin samples

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**FIG 4** Epistasis of *atrR* and *ffmA*. (A) To test the relationship between *atrR* and *ffmA*, the *ffmA*Δ cassette was introduced into an *hspA-atrR* fusion-containing strain. This double mutant strain was grown along with isogenic wild-type and *ffmA*Δ strains, spotted onto minimal medium, and allowed to grow at 37°C. Plates were photographed after 3 days. (B) Whole-cell protein extracts were made from the strains described above and analyzed by Western blotting using the anti-AbcG1 antiserum. The top panel shows Ponceau S staining of the membrane prior to blotting.

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C$_2$H$_2$ Factor FfmA Regulates Voriconazole Susceptibility
from these two strains were sheared, immunoprecipitated with the appropriate antibody, and then analyzed by quantitative PCR (qPCR).

ChIP using anti-Flag antibodies in samples from the DO-Flag-ffmA-containing strain showed strong enrichment of the Flag-FfmA protein on both the \emph{abcG1} and \emph{ffmA} genes. The structure of the doxycycline-off promoter is shown by the bar at the top where \textit{bleR} is the bleomycin resistance marker, tTA represents the region expressing the doxycycline-off transactivator, and \textit{tetP} indicates the location of the doxycycline-responsive minimal promoter. When the FLAG tag is present, it is inserted immediately before the native \emph{ffmA} ATG (shown on the diagram). The \emph{ffmA} gene contains a 1,200-nucleotide-long \textit{5}'- untranslated leader that starts at the position indicated by the rightward-facing arrow. The scale of the diagram is indicated at the bottom left. (B) The conditional doxycycline-repressible (dox off, or DO) promoter was inserted just upstream of the ATG for the \emph{ffmA} open reading frame, either with or without a single FLAG epitope fused to the FfmA amino-terminal methionine residue. This makes transcription of the DO-ffmA fusion gene sensitive to the presence of exogenous doxycycline. Isogenic wild-type and DO-ffmA strains were grown and then spores placed on minimal medium containing or lacking doxycycline at the indicated concentration. Plates were developed at 37°C as before. (C) The strains used for panels A and B were grown in minimal medium with no doxycycline overnight and then diluted into fresh medium lacking (-) or containing (+) the indicated concentration of doxycycline. These cultures were allowed to grow for an additional 12 h and whole-cell protein extracts prepared. Extracts were prepared after normalizing by mycelial dry weight of each culture. Equivalent amounts of each protein extract were analyzed for total loading and levels of FfmA, AbcG1, and tubulin using appropriate antibodies. The numbers below each specific immunoreactive band indicate expression level normalized to the wild-type strain.
promoters (Fig. 6A). Note that in the case of this strain, the DO promoter was inserted between the normal ffmA promoter and the ffmA coding region, leading to retention of the wild-type ffmA promoter, although it is no longer controlling expression of ffmA.

We also examined ChIP of FfmA in unaltered AfS35 cells using anti-FfmA antibody. ChIP of FfmA in this strain showed similar enrichment of FfmA to both the fully native ffmA and abcG1 promoters. In the case of the Flag-FfmA and wild-type FfmA proteins, only marginal enrichment was seen at the atrR promoter, although it was greater than that seen on the actin promoter (act1) used as a presumptive negative control. These data are consistent with FfmA directly controlling gene expression of ffmA and abcG1 by direct promoter binding and potentially indirectly controlling atrR expression.

DISCUSSION

Our interest in ffmA emerged from the effect of this gene on itraconazole resistance as shown in the early analysis of a collection of transcription factor gene deletion mutants (17). More detailed analysis of the phenotype of a ffmAΔ strain clearly indicated that loss of this gene had a modest azole drug phenotype compared to its
pronounced defect in growth. Other screenings of this transcription factor deletion mutant library led to the detection of \textit{ffmA} as a determinant required for resistance to cell wall stress agents but also showed a very similar general growth defect (19). Based on both this and the previous study of the phenotype of the \textit{ffmA}\textit{D} strain, it seems that the primary phenotype caused by loss of \textit{ffmA} is a severe growth deficit. This is in contrast to deletion of genes such as \textit{srbA} (18) or \textit{atrR} (15) that cause striking sensitivity to azole drugs but grow relatively well in drug-free media. \textit{FfmA} is necessary for normal expression of \textit{abcG1}, but our data indicate a wider role in cell growth beyond azole resistance.

Based on the presence of a C2H2 zinc finger-containing DNA binding domain in its protein sequence, we believe that \textit{FfmA} is a sequence-specific transcription factor. Our ChIP experiments support this view, as \textit{FfmA} is found to enrich on the \textit{abcG1} promoter, which would be consistent with \textit{FfmA} acting to positively regulate expression of this gene. Additionally, we found that this same \textit{abcG1} fragment was bound by \textit{AtrR} (16), suggesting that \textit{FfmA} and \textit{AtrR} bind the same or adjacent regions of the \textit{abcG1} promoter. We also found that \textit{FfmA} showed enrichment on its own promoter, consistent with its expression being autoregulated.

The shared role for \textit{AtrR} and \textit{FfmA} in control of \textit{abcG1} expression led us to examine the interaction between these two genes. Overproduction of \textit{AtrR} using the \textit{hspA} promoter was well tolerated by the cell and led to both increased azole resistance (16) and strong suppression of the growth phenotype caused by loss of \textit{ffmA}. Owing to the difficulty in growing \textit{ffmA}\textit{D} strains, we disrupted \textit{ffmA} in the \textit{hspA-atrR} background. These transformants were readily obtained and grew normally. Furthermore, the contribution of \textit{FfmA} to \textit{AbcG1} expression was strongly suppressed by the increased levels of \textit{AtrR}. This genetic interaction led us first to suspect that \textit{FfmA} acts upstream of \textit{AtrR} in terms of function, but ChIP experiments on the \textit{atrR} promoter have not yet revealed strong enrichment for \textit{FfmA}, although we have not scanned the entire promoter. Another possibility is that both \textit{AtrR} and \textit{FfmA} stimulate expression of \textit{abcG1} transcription, and the increased levels of \textit{AtrR} are adequate to reverse the transcriptional defect caused by loss of \textit{ffmA}.

The precise role of \textit{FfmA} in the cell remains to be determined. Clearly, \textit{A. fumigatus} is exquisitely sensitive to modifications of \textit{FfmA} expression. As we show here, either loss of \textit{ffmA} or its overproduction from the \textit{hspA} promoter caused a strong defect in normal growth. Driving either a Flag-tagged or untagged \textit{FfmA} from the \textit{dox-off} promoter did allow growth that was similar to that of the wild type but remained only 80% of normal. This sensitivity to variation in expression level has been noted before in the analysis of human disease genes (24), in which genes linked to disease are rarely seen to be associated with copy number variation owing to their critical importance. These genes are often associated with developmental processes, as disturbances in the expression profiles of genes of this type lead to loss of viability. In the case of \textit{ffmA} dosage changes, cells are still viable but clearly suffer significant impacts on growth. Since this \textit{ffmA}\textit{D}-mediated growth defect can be robustly suppressed with \textit{atrR} overproduction, we suggest that the downstream functions of these transcription factors overlap. As discussed above, this is certainly true for control of \textit{abcG1} expression, but as \textit{abcG1}\textit{D} strains have no detectable growth defect in the absence of azole drugs, other target genes must be shared that impact growth under unstressed conditions. Identification of these genes will be an important future goal.

\textbf{MATERIALS AND METHODS}

\textit{A. fumigatus} strains, growth conditions, and transformation. The lab strains that were used in this study are listed in Table 1. For initial shortlisting of strains that were defective for \textit{abcG1} expression, the \textit{itraconazole}-susceptible members of the transcription factor deletion mutant were used (17). \textit{A. fumigatus} strains were typically grown at 37°C in rich medium (Sabouraud dextrose: 0.5% tryptone, 0.5% peptone, 2% dextrose [pH 5.6 \pm 0.2]). Selection of transformants was performed using minimal medium (MM: 1% glucose, nitrate salts, trace elements, 2% agar [pH 6.5]). Trace elements, vitamins, and nitrate salts are as described in the appendix of reference 25, supplemented with 1% sorbitol and either 50 mg/liter phleomycin (after adjusting the media to pH 7) or 200 mg/liter hygromycin gold (both InvivoGen).
For solid medium, 1.5% agar was added. Doxycycline promoter shut-off experiments were performed by adding 25 mg/liter doxycycline (BD Biosciences). Wild-type and doxycycline-repressible \textit{ffmA}-containing strains were grown overnight in the absence of doxycycline. Doxycycline repression was initiated and incubation continued for 12 h prior to sample preparation and analysis.

Transformation and generation of \textit{ffmA} mutants was done using \textit{in vitro}-assembled cas9-guide RNA ribonucleoproteins coupled with 50-bp microhomology repair templates (26). For generation of \textit{ffmA} deletion mutants, 2 CRISPR RNAs (5'\text{-ATCTAGGATCCATCATGAAG, corresponding to the 5' end of the gene, and 5'-ATAGTCAATGGCTCAAGGGA, corresponding to the 3' end of the gene}) were used to replace \textit{ffmA} with the hygromycin resistance marker cassette amplified from the plasmid pSP62 (16) using ultramer-grade oligonucleotides from IDT harboring 50-bp homology to the upstream and downstream junctions of the \textit{ffmA} gene. The doxycycline-repressible (DO) promoter (with or without a Flag tag) marked with the phleomycin resistance cassette was inserted upstream of the \textit{ffmA} gene using the single CRISPR RNA 5'-ATCTAGGATCCATCATGAAG. For generating the DO-\textit{ffmA} allele, a microhomology repair template containing the resistance marker cassette and DO promoter were PCR amplified from plasmid pSP114. The plasmid pSP114 was generated from pSK606-\textit{ptrA} (provided by J. Fortwendel), from which the \textit{ptrA} marker was replaced by a phleomycin resistance gene (Ble) by cloning the latter from pCH008-Phleor (also from J. Fortwendel) at the KpnI and PstI sites. To generate the Ble-DO-Flag cassette, complementary oligonucleotides corresponding to a 1\times Flag tag was annealed and ligated to the ends of Pmel-linearized pSP114. This plasmid was named pSP115 and was used as the template to generate microhomology repair constructs to generate the Ble-DO-Flag-\textit{ffmA} allele. Transformants were genotypically confirmed by diagnostic PCR of the novel upstream and downstream junction formed upon targeted integration as well as by PCR amplification to confirm lack of the DNA binding domain of \textit{ffmA} in the case of \textit{ffmA} deletion mutants. At least 3 independent, properly targeted transformants were phenotyped for all \textit{ffmA} mutants, of which a representative strain is depicted in the data presented.

**Radial growth/drug disc diffusion assay.** Fresh spores of \textit{A. fumigatus} were suspended in 1× phosphate-buffered saline (PBS) supplemented with 0.01% Tween 20 (1× PBS). The spore suspension was counted using a hemocytometer to determine the spore concentration. Spores were then appropriately diluted in 1× PBS. For the drug diffusion assay, 1× 10^6 spores were mixed with 10 ml soft agar (0.7%) and poured over 15 ml of regular agar containing (1.5%) minimal medium. A paper disk was placed on the center of the plate, and 10 μl 1 mg/liter voriconazole was spotted onto the sterile filter paper. For the radial growth assay, ~100 spores (in 4 μl) were spotted on minimal medium with or without the drug. The plates were incubated at 37°C and inspected for growth every 12 h.

**Generation of an FfmA antibody.** A region of 502 bp (corresponding to amino acids 195 to 340 based on homology to the DNA binding domain of ScAdr1) from \textit{ffmA} was PCR amplified and cloned in frame as an NdeI/HindIII fragment upstream of the C-terminal 6×His tag in pET29a+ (EMD Millipore, Inc.) to form plasmid pSP113 and was transformed into the \textit{Escherichia coli} strain BL21(DE3). Two liters of transformed bacteria was grown to log phase and induced with isopropyl-β-D-thiogalactopyranoside for 90 min. Cell lysates were prepared using a French press and subsequent protein purification accomplished using Talon metal affinity resin (Takara Bio USA, Inc.) as described by the manufacturer. Protein fractions were analyzed by staining them with Coomassie blue and by Western blotting using His-specific antibodies. The purified proteins were then lyophilized and sent to Pacific Immunology (Ramona, CA) for injection into rabbits to generate polyclonal antibodies against FfmA. Antiserum generated from these rabbits was received and tested for immunoreactivity against \textit{A. fumigatus} cell lysates. The antiserum was then purified using an AminoLink Plus coupling resin (Thermo Scientific, Inc.) according to the manufacturer’s instructions, and the affinity-purified antiserum was used to detect the FfmA protein from \textit{A. fumigatus} cell lysates.

**Western blotting.** Western blotting was performed as described in reference 14. The FfmA polyclonal antibody used here has been detailed in the reference mentioned above and was used at a 1:500 dilution. The anti-Flag M2 monoclonal antibody (F1804) was procured from Sigma and used at a 1:2,000 dilution. AbcG1 polyclonal antibody (21) was used at a dilution of 1:500.

**Chromatin immunoprecipitation.** Chromatin immunoprecipitation was done as described in reference 16, with the following modifications: 30 μl was reserved as an input control (IC) fraction for reverse cross-linking to verify sonication and control for ChiP and qPCR. The sheared chromatin was incubated

### Table 1. \textit{A. fumigatus} strains used in this study

| Strain       | Parent | Genotype          | Source or reference |
|--------------|--------|-------------------|---------------------|
| AF135        | D141   | akuAD33uraP       | FGSC                |
| A1160        | CEA10  | akuBD33pyrG       | MFIG                |
| 1G9          | A1160' | ffpMA133hph        | MFIG                |
| SPF62        | AF53   | abcG1Δhph         | 21                  |
| SPF87        | AF53   | atrRΔptrA         | 16                  |
| SPF108A      | AF53   | hspA-atrR         | This study          |
| SPF142       | AF53   | ffpMA33hph        | This study          |
| SPF144       | SPF108A| hspA-atrR ffpMAΔhph| This study          |
| SPF146       | AF53   | hspA-ffmAΔhph     | This study          |
| SPF182       | AF53   | Dex Off-ffmAΔBle   | This study          |
| SPF185       | AF53   | Dex Off-Flag-ffmAΔBle| This study          |
with either anti-Flag M2 monoclonal antibody (F1804; Sigma) at 1:250 dilution or with anti-FfmA polyclonal antibody (described above) at a dilution of 1:50 overnight (16 h) on a nutator at 4°C. This sample was further incubated with 50 μl of washed Dynabeads (Life Technologies) conjugated to either protein G (when using anti-Flag) or protein A (when using anti-FfmA) for another 8 h. Real-time PCR of ChIP DNA was also performed as described in reference 1S, with the following modifications: 0.5 μl of ChIP or input (diluted 30-fold to bring it to 1%) DNA was used in a 20-μl total volume reaction mix using SYBR green master mix (Bio-Rad) and 0.4 μM each primer. Fold enrichment was calculated to determine the enrichment of the promoter region for each gene. The oligonucleotide primer pairs used to check promoter enrichment of the promoter region for each gene. The oligonucleotide primer pairs used to check promoter enrichment consist of fma-ChIP-F (5'-GCTGAAATATGATGCTCCTC), fma-ChIP-R (5'-ACCTTCTGATCTTCGTTGGAAC), atrR-ChIP-F (5'-ACGGAGCTCCTTCTTACTC), atrR-ChIP-R (5'-CGAAAAAGAGTCTGCCTTC), abg1-ChIP-F (5'-CCTTATACTCGAATCCAC), abg1-ChIP-R (5'-TCCTTCTTGGACCGGAC), actA-ChIP-F (5'-GCCACCTAAGGTTAC), and actA-ChIP-R (5'-GCCGCTTCGTATAGGAGACC). The positions of each amplicon produced for these promoters are the following: fmaA, −2338 to −1973; atrR, −1035 to −700; abg1, −766 to −445; and actA, −916 to −580. Note the numbering is relative to the ATG for each gene.

SUPPLEMENTAL MATERIAL

Supplemental material is available online only.

FIG S1, PDF file, 0.4 MB.

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