Deciphering the Regulatory Network between the SREBP Pathway and Protein Secretion in *Neurospora crassa*

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ABSTRACT  Sterol regulatory element binding proteins (SREBPs) are conserved from yeast to mammalian cells and function in the regulation of sterol homeostasis. In fungi, the SREBP pathway has been implicated in the adaptation to hypoxia and in virulence. In *Neurospora crassa* and *Trichoderma reesei*, the SREBP pathway also negatively regulates protein secretion under lignocellulolytic conditions. Here we utilized global transcriptional profiling combined with genetic and physiological analyses to address the regulatory link between the SREBP pathway and protein secretion in *N. crassa*. Our results demonstrated that the function of the SREBP pathway in ergosterol biosynthesis and adaptation to hypoxia was conserved in *N. crassa*. Under lignocellulolytic conditions, the SREBP pathway was highly activated, resulting in the reduced expression of lytic polysaccharide monooxygenases, which require molecular oxygen for catalytic activity. Additionally, activation of the SREBP pathway under lignocellulolytic conditions repressed a set of genes predicted to be involved in the endoplasmic reticulum stress response. Here we show that the inability of a *hac-1* mutant, which bears a deletion of the major regulator of the unfolded protein response (UPR), to efficiently produce cellulases and utilize cellulose was suppressed by mutations in the SREBP pathway. The analyses presented here demonstrated new SREBP pathway functions, including linkages to the UPR, and provide new clues for genetic engineering of filamentous fungi to improve their production of extracellular proteins.

IMPORTANCE  The role of SREBP transcription factors in the regulation of sterol biosynthesis is conserved from humans to yeast. In filamentous fungi, this pathway regulates the secretion of lignocellulolytic enzymes during plant biomass deconstruction. Here we show that the SREBP pathway in *Neurospora crassa* regulates the production of specific cellulases, lytic polysaccharide monooxygenases that utilize molecular oxygen. Via global transcriptional profile and genetic analyses, a relationship between the SREBP pathway and the unfolded protein response (UPR) pathway was revealed, suggesting a regulatory interplay of these two pathways in the trafficking of plant biomass-degrading enzymes. These findings have implications for our understanding of the cross talk of the SREBP and UPR pathways in other organisms and will guide the rational engineering of fungal strains to improve cellulolytic enzyme production.

Nonfood sources of plant biomass have been considered promising material for producing renewable biofuels and other bioproducts (1). However, plant biomass-derived products are not currently economically viable primarily because of the cost of deconstructing polymers in the plant cell wall to fermentable sugars by lignocellulosic enzymes (2), which are primarily produced by filamentous fungi. The model fungal species *Neurospora crassa* has the ability to degrade plant cell wall material into simple...
sugars for growth and reproduction. With this property and its well-developed genetic, molecular, and biochemical tools, \textit{N. crassa} has been used to investigate plant cell wall deconstruction by filamentous fungi (3).

To identify unknown factors associated with protein trafficking and secretion in \textit{N. crassa}, ~600 strains carrying deletions in genes predicted to function in the secretory pathway were screened for alterations in the secretion of cellulolytic enzymes (4). The \textit{dsc-2} and \textit{tul-1} (\textit{dsc-1}) mutant hypersecretion strains were identified. The homologs of \textit{dsc-2} and \textit{tul-1} in \textit{Schizosaccharomyces pombe} and \textit{Aspergillus fumigatus} are components of the Golgi apparatus E3 ligase complex critical for the activation of a homolog of the sterol regulatory element binding protein (SREBP) transcription factor, Sre1 or SreA, respectively, by proteolytic cleavage (5, 6). In mammalian cells, the SREBP pathway functions in the regulation of sterol homeostasis. Proteolytic cleavage releases an N-terminally active SREBP from the endoplasmic reticulum (ER) that is subsequently transported into the nucleus to activate the expression of genes associated with cholesterol biosynthesis and lipid metabolism (7). In \textit{N. crassa}, a strain carrying a deletion of the SREBP homolog \textit{sah-2} or mutations in another predicted component of the Golgi apparatus E3 ligase, \textit{dsc-3} or \textit{dsc-4}, show hyperproduction of cellulases (4). A cellulase hyperproduction phenotype was also observed in a \textit{Trichoderma reesei} strain bearing a deletion of a homolog of either \textit{sah-2} or \textit{dsc-1} (4), suggesting a conserved function of the SREBP pathway in the regulation of protein secretion under lignocellulolytic conditions. In \textit{S. pombe}, \textit{A. fumigatus}, and \textit{Cryptococcus neoformans}, in addition to regulation of intracellular sterol homeostasis and adaptation to hypoxic conditions (8–10), the SREBP pathway is associated with the regulation of cell polarity, drug resistance, and virulence (10, 11).

In this study, we further investigated the role of the SREBP pathway in the secretion of lignocellulolytic enzymes by \textit{N. crassa}. We demonstrated that strains carrying mutations in the homologs of SREBP cleavage-activating protein-encoding gene \textit{scp-1} and rhomboid protease-encoding gene \textit{rbd-2} also showed a cellulase hyperproduction phenotype. In addition to regulating ergosterol biosynthesis and hypoxia adaptation, the \textit{N. crassa} SREBP pathway negatively regulated the expression of genes encoding cellulobiose dehydrogenase (CDH-1) and most of the lytic polysaccharide monooxygenases (LPMOs), as well as many ER stress-responsive genes. Furthermore, deletion of \textit{sah-2} rescued the cellulase-negative defect of the \textit{Δhac-1} mutant; HAC-1 is the major regulator of the unfolded protein response (UPR). These findings extend our mechanistic understanding of the regulation of cellulase enzyme secretion and provide a link between the SREBP and UPR pathways in filamentous fungi during plant biomass deconstruction. Additionally, these studies provide new strategies for targeted engineering of filamentous fungi to improve current yields of plant cell wall-degrading enzymes.

\textbf{RESULTS}

\textbf{Identification of components of the SREBP pathway in \textit{N. crassa}.} SREBPs are synthesized in an inactive form and anchored in the ER membrane. In the ER membrane, SREBPs bind to the SREBP cleavage-activating protein Scap, which has a sterol-sensing domain. In sterol-replete cells, the SREBP-Scap complex remains in the ER, with Scap binding to the ER-resident protein INSIG. In sterol-depleted cells, Scap-INSIG binding is disrupted and Scap escorts SREBP to the Golgi apparatus (7). In \textit{S. pombe} and \textit{A. fumigatus}, a Golgi apparatus E3 ligase complex is employed for proteolytic activation of SREBP (5, 6, 12), along with a rhomboid protease, Rbd-2/RbdB (13, 14). In \textit{Aspergillus nidulans} and \textit{A. fumigatus}, a signal peptide peptidase, SppA, is also involved in the cleavage of SREBP (15).

A search of the \textit{N. crassa} genome for genes involved in the SREBP pathway in \textit{Homo sapiens}, \textit{S. pombe}, and \textit{A. fumigatus} showed that \textit{N. crassa} has two homologs of SREBP (\textit{sah-2} and NCU01871). We named NCU01871 \textit{sre-2}. Both \textit{sah-2} and \textit{sre-2} contain a basic helix-loop-helix (bHLH) DNA binding domain in their N terminus and have a tyrosine residue in the bHLH domain, which is a unique characteristic of SREBPs. SAH-2 also has
a DUF2014 domain in its C terminus, which is also found in S. pombe Sre1 and A. fumigatus SrBα, while SRE-2 lacks this C-terminal domain. N. crassa has no homolog of the INSIG protein but does have a homolog of Scap1, SppA, and rhomboid protease Rbd-2/RbdB. The homolog of Scap was named scp-1 (NCU03675); the homologs of the components in the Golgi apparatus E3 ligase complex were named tul-1 (here dsc-1; NCU03740), dsc-2 (NCU03459), dsc-3 (NCU11245), dsc-4 (NCU02676), dsc-5 (NCU01928), and dsc-6 (NCU00018). The homolog of the peptide peptidase SppA was named spp-1 (NCU00568) and the homolog of the rhomboid protease Rbd-2/RbdB was named rbd-2 (NCU02371). Among these homologs, Δscp-1, Δsre-2, Δdsc-3, and Δrbd-2 were not available in the N. crassa deletion strain collection. We therefore generated Δscp-1 and Δsre-2 deletion mutants by replacement of the resident locus with the hygromycin resistance gene hph (16) (see Materials and Methods).

Previously, we determined that deletion mutations in sah-2, dsc-1, dsc-2, and dsc-4 resulted in strains that showed hyperproduction of cellulases when exposed to cellulose (4). However, the effect of these mutations on the response to hypoxic conditions has not been determined. To determine if the SREBP pathway regulates adaptation to hypoxia in N. crassa and if the identified homologs are required, we assessed the growth phenotype of the Δsah-2, Δsre-2, Δscp-1, Δdsc-1, Δdsc-2, Δdsc-4, Δdsc-5, Δrbd-2, and Δspp-1 mutants under hypoxia and normoxia conditions. The deletion mutants showed somewhat variable growth rates under normoxia conditions, the Δdsc-1 and Δdsc-4 mutants showed a significantly lower growth rate, while the Δsah-2, Δscp-1, and Δrbd-2 mutants grew slightly more slowly (Fig. 1A). The growth rates of the Δdsc-5, Δsre-2, and Δspp-1 mutants displayed growth rates that were not statistically significantly different from that of the wild type (WT). However, under hypoxic condition (0.2% O₂ in N₂), all but the Δsre-2 and Δspp-1 mutant strains showed a severe growth defect (Fig. 1B; see Fig. S1A in the supplemental material). Additionally, the hypoxia-sensitive strains showed reduced aerial hyphal growth under normoxia conditions, while the Δsre-2 and Δspp-1 mutants did not (see Fig. S1B). These data indicate that the SREBP components scp-1, the Golgi apparatus E3 ligase complex, rbd-2, and sah-2 have a conserved role in hypoxia adaptation in N. crassa, while sre-2 and spp-1 do not.

To further confirm the hypothesis that any strain containing a loss-of-function mutation in a component of the SREBP pathway would show a cellulolytic hyperproduction phenotype, we compared the total secreted protein level and endoglucanase activity under Avicel conditions of the nine predicted SREBP pathway deletion strains with those of the WT strain. As predicted, these results showed that the total secreted protein level and endoglucanase activity of the Δsah-2, Δscp-1, Δdsc-1, Δdsc-2, Δdsc-4, Δdsc-5, and Δrbd-2 mutants were significantly higher than those of the WT strain (Fig. 1C and D). In contrast, the two mutants that showed no defect in growth under hypoxic conditions, the Δsre-2 and Δspp-1 mutants, also showed no hyperproduction of cellulases. Taken together, these data indicated that the SREBP pathway, which is important for the response of N. crassa to hypoxic conditions, also plays a role in the regulation of the production/secretion of cellulolytic enzymes.

**Ergosterol levels are reduced in SREBP-related mutants.** To determine whether the SREBP pathway plays a role in sterol homeostasis and to determine if this role is involved in the cellulase hyperproduction phenotype in N. crassa, we crossed three key SREBP pathway mutants, the Δsah-2, Δdsc-1, and Δscp-1 mutants, with a strain that constitutively expresses the major transcriptional regulator of cellulolytic genes (Pccg-1-clr-2; termed Mclr-2) (17), resulting in strains Mclcr-2 Δsah-2, Mclcr-2 Δdsc-1, and Mclcr-2 Δscp-1. We constructed strains containing this clr-2 construct (Pccg-1-clr-2), as the production of cellulases is inducer independent and the cellulase hyperproduction phenotype of the Δdsc-1 and Δdsc-2 mutants was conserved in these strains (4). Consistent with the phenotype of the single-deletion mutants, the Mclcr-2 Δsah-2, Mclcr-2 Δdsc-1, and Mclcr-2 Δscp-1 strains grew slightly more slowly than the Mclr-2 strain, showed reduced aerial hyphae when grown on sucrose under normoxic conditions, and displayed growth defects under hypoxic conditions (see Fig. S1C to F). To assess
the relationship between protein secretion and ergosterol in these mutants, the total secreted protein levels and intracellular ergosterol levels in the \( M^{\Delta s\text{ah}-2} \), \( M^{\Delta s\text{ah}-2} \), and \( M^{\Delta s\text{ah}-2} \) strains were compared to those in the \( M^{\Delta s\text{ah}-2} \) strain. As expected, the protein secretion levels in the \( M^{\Delta s\text{ah}-2} \), \( M^{\Delta s\text{ah}-2} \), and \( M^{\Delta s\text{ah}-2} \) strains were significantly increased. However, ergosterol levels were significantly lower in the SREBP-related mutants than in the \( M^{\Delta s\text{ah}-2} \) strain (Fig. 2A). Moreover, gas chromatography (GC)-mass spectrometry (MS) showed that the pattern abundances of the sterol intermediates in the SREBP-related mutants were very similar (Fig. 2B and C) to those of the \( M^{\Delta s\text{ah}-2} \) strain but with important key differences. For example, two ergosterol intermediates, 4,4-dimethyl-zymosterol and lanosterol, accumulated in the SREBP-related mutants, while 5,7,22,24(28)-ergostatetraenol and an unidentified component with a retention time of 7.08 min were markedly lower than in the \( M^{\Delta s\text{ah}-2} \) strain.

To determine whether SREBP has a role in ergosterol biosynthesis under noncellulosic conditions, we compared the levels of ergosterol and its intermediates in the \( \Delta \text{sah-2} \) mutant and WT strains under sucrose conditions (see Fig. S2). The results showed that the sterol pattern in the \( \Delta \text{sah-2} \) mutant was similar to that of the \( M^{\Delta d\text{sc}-1} \) strain, and \( M^{\Delta d\text{sc}-1} \) strains (Fig. 2; see Fig. S2). These data indicated that the SREBP pathway regulates ergosterol homeostasis in \( N. \text{crassa} \), as observed in other fungi.

Ergosterol is a component of fungal cell membranes, and the concentration of sterols within eukaryotic membranes affects many membrane-associated functions.
including vesicle formation and protein sorting (18). For example, blocking of ergosterol synthesis with the antifungal agent fluconazole results in increased heterologous recombinant protein production in the yeast *Pichia pastoris* (19). To determine whether the hypersecretion phenotype observed in SREBP-related mutants resulted from lower ergosterol content, we constructed Δ*erg-25* mutant strains by replacing the resident gene with the nourseothricin acetyltransferase gene *nat1* (see Materials and Methods). The Δ*erg-25* deletions, along with the Δ*erg-3* and Δ*erg-24* deletions (available from the *N. crassa* deletion collection), were introduced into the *Mclr-2* strain via crosses. Intracellular ergosterol levels and the total secreted protein levels of the *Mclr-2 Δerg-3*, *Mclr-2 Δerg-24*, and *Mclr-2 Δerg-25* strains under sucrose conditions were measured. As expected, the ergosterol levels in all three mutants were significantly lower than those in the *Mclr-2* strain. However, the levels of secreted proteins were not increased in any of the ergosterol mutants in the *Mclr-2* background (Fig. 2D). As an alternative approach to explore the relationship between ergosterol levels and protein secretion, we used fluconazole, a specific inhibitor of the essential protein ERG-11, to treat the *Mclr-2* strain under sucrose conditions. As shown in Fig. S2C, treatment with fluconazole also did not increase secreted protein levels. These data indicated that the hyperproduction of proteins observed in the SREBP-related mutants under cellulosic conditions did not result from disruption of the ergosterol biosynthesis pathway.

**Transcriptional profiling revealed that the SREBP pathway is highly activated under lignocellulolytic conditions.** To gain insight into the molecular basis of the

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**FIG 2** SREBP-regulated ergosterol biosynthesis pathway in *N. crassa*. (A) Total secreted protein level and intracellular ergosterol levels in a 48-h VMM culture of the strains indicated after a shift from a 16-h VMM culture. (B) GC-MS chromatogram of sterol extracts from the strains indicated. MCounts, mega counts, indicating the intensity (abundance) of the signal for each peak. (C) Quantified level of each sterol component in extracts of *Mclr-2, Mcr*-Δ*dsc-1, Mcr*-Δ*sah-2*, and *Mcr*-Δ*scp-1* cultures. The relative level of each sterol component was normalized by cholesterol internal standard. (D) Total secreted protein level and intracellular ergosterol levels of the strains indicated under the same conditions as for panel A.
relationship between the SREBP pathway and hyperproduction of cellulases, we conducted transcriptome sequencing (RNA-seq) to compare global changes in gene expression levels among the WT, Δsah-2 mutant, Mclr-2, and Mclr-2 Δsah-2 strains grown in sucrose minimal medium. Under this condition, cellulase gene expression is repressed in the WT and Δsah-2 mutant strains but induced in the Mclr-2 and Mclr-2 Δsah-2 strains. Principal-component analysis (PCA) and Euclidean distance analyses of the 12 sets of RNA-seq data showed that the biological replicate samples clustered together and that the expression pattern of Mclr-2 Δsah-2 was most similar to that of the Δsah-2 mutant (see Fig. S3). Numbers of reads per kilobase of transcript per million mapped reads (RPKM) for all genes were calculated for biological replicates, and differential gene expression analyses (≥4-fold change; P ≤ 0.01) between Δsah-2 mutant and WT, Mclr-2 and WT, Mclr-2 Δsah-2 and Mclr-2, and Mclr-2 Δsah-2 and Δsah-2 mutant samples were performed (see Fig. S4 and Data Set S1).

Hierarchical clustering of genes involved in ergosterol biosynthesis revealed that erg-1 (NCU08280), erg-11 (NCU02624), erg-3 (NCU06207), erg-25 (NCU06402), and erg-5 (NCU05278) showed significantly lower expression levels in the Δsah-2 mutant compared to the WT and in Mclr-2 and Mclr-2 Δsah-2 strains. Principal-component analysis (PCA) and Euclidean distance analyses of the 12 sets of RNA-seq data showed that the biological replicate samples clustered together and that the expression pattern of Mclr-2 Δsah-2 was most similar to that of the Δsah-2 mutant (see Fig. S3). Numbers of reads per kilobase of transcript per million mapped reads (RPKM) for all genes were calculated for biological replicates, and differential gene expression analyses (≥4-fold change; P ≤ 0.01) between Δsah-2 mutant and WT, Mclr-2 and WT, Mclr-2 Δsah-2 and Mclr-2, and Mclr-2 Δsah-2 and Δsah-2 mutant samples were performed (see Fig. S4 and Data Set S1).

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In the Mclr-2 strain, the erg-1, erg-11, erg-3, erg-25, and erg-5 expression levels were strongly increased relative to those in the WT strain, suggesting that the induction of cellulases affected SAH-2 activation and thus expression of the erg genes (Fig. 3A). Importantly, the clr-2 transcript abundance in Mclr-2 Δsah-2 was not higher than in Mclr-2, indicating that the hyperproduction of cellulases in Mclr-2 Δsah-2 was not due to an increase in clr-2 expression. To test the hypothesis that the induction of cellulases is associated with the activation of SAH-2, we performed qRT-PCR to examine the erg-3, erg-11, and erg-25 mRNA abundance in the WT, Mclr-2, Δsah-2 mutant, and Mclr-2 Δsah-2 strains.

FIG 3 Influence of sah-2 on the expression of genes involved in the ergosterol biosynthesis pathway. (A) Hierarchical clustering of genes involved in ergosterol biosynthesis. Results are displayed as heat maps with log numbers of RPKM ranging from minimum (bright blue) to maximum (bright yellow). The SREBP direct targets, as identified in A. fumigatus (30), are shown in bold at the right. (B) Ergosterol biosynthesis pathway. The genes highlighted in yellow were those whose expression levels were downregulated both in the Δsah-2 mutant versus the WT and in the Mclr-2 Δsah-2 strain versus the Mclr-2 strain. The genes highlighted in pink were those whose expression levels were upregulated both in the Δsah-2 mutant versus the WT and in the Mclr-2 Δsah-2 strain versus the Mclr-2 strain. Oxygen-requiring enzymes are marked.
strains at different time points. These data showed that the erg gene expression levels in the Mcr-2 strain were higher at 24 and 48 h (see Fig. S5B) than at the 4-h time point. Since these genes are predicted to be directly regulated by SAH-2 (see Discussion), this observation indicated that the SREBP pathway was activated during the induction and trafficking of cellulases, which are associated with deconstruction of plant biomass.

In S. pombe and A. fumigatus, the activation of orthologs of SREBP (Sre1 and SreA, respectively) is triggered by hypoxia. This observation suggested that the trafficking and secretion of cellulases might result in reduction of oxygen availability in cells, resulting in activation of the SREBP pathway. To test this hypothesis, we performed growth experiments with parallel bioreactor fermenters containing a dissolved oxygen (DO) control system (see Materials and Methods). Consistent with this hypothesis, the increased protein secretion in Mcr-2 control system (see Materials and Methods). Consistent with this hypothesis, the increased protein secretion in Mcr-2 Δsah-2 was only observed when the automatic DO control system in the fermenter was turned off. The levels of protein secretion in the Mcr-2 Δsah-2 and Mcr-2 strains were not significantly different when the automatic DO control system was turned on (see Fig. S5C). These data suggest that cellulase production leads to higher oxygen consumption, thereby lowering the oxygen level in the surrounding environment and resulting in activation of the SREBP pathway.

**CDH-1 and LPMO gene expression levels increase in the SREBP mutant.** Our previous data showed that deletion of sah-2 or dsc-1 does not significantly affect the expression level of the major cellulase genes cbh-1, cbh-2, and gh5-1 (4). To determine if the expression of genes encoding other extracellular proteins was affected in the SREBP pathway mutants, we performed hierarchical clustering of genes encoding proteins identified as extracellular when N. crassa was exposed to Avicel or Miscanthus (20). This analysis showed that the expression of ~2/3 of this gene set is CLR-2 dependent and 20 of them showed higher expression levels in the Mcr-2 Δsah-2 strain than in the Mcr-2 strain (Fig. 4A; see Table S1). Ten of these 20 genes encode LPMO enzymes. LPMOs are copper-dependent redox enzymes that require molecular oxygen and an external electron donor to cleave glycosidic bonds (21). CDH-1, the key electron donor for LPMOs (22), showed 12-fold higher expression in the Mcr-2 Δsah-2 strain than in the Mcr-2 strain. Consistent with the RNA-seq data, SDS-PAGE analysis of the supernatants of the Mcr-2 Δsah-2 and Mcr-2 strains showed variations in specific protein levels rather than an overall increase in the levels of secreted proteins in the Mcr-2 Δsah-2 strain (Fig. 4B). We thus measured the extracellular activity of cellobiohydrolase (CBH-1; cbh-1 expression was not significantly affected in the Δsah-2 mutant) and CDH-1 (cdh-1 expression increased in the Δsah-2 mutant) from the Mcr-2 Δsah-2 and Mcr-2 strains. These data showed that the activity of CDH-1 was 2.2-fold higher and that of CBH-1 was 1.4-fold higher in the Mcr-2 Δsah-2 strain than that in the Mcr-2 strain (Fig. 4C). These data suggested that the expression level of genes and thus the activity of the enzymes associated with oxygen consumption under lignocellulolytic conditions are negatively regulated by SAH-2.

There are 15 LPMO genes predicted in the N. crassa genome. To determine if SAH-2 influences the expression of the remaining five LPMOs, we assessed the expression levels of these remaining LPMO genes in the WT, Mcr-2, Δsah-2 mutant, and Mcr-2 Δsah-2 strains. While the expression of two predicted LPMO genes, gh61-8 (NCU03000) and gh61-15 (NCU07974), was not affected by the deletion of sah-2, the expression levels of two additional LPMO genes, gh61-10 (NCU01867) and gh61-12 (NCU02344), were 7.6- and 2.3-fold higher, respectively, in the Δsah-2 mutant than in the WT (Fig. 4D). Thus, deletion of sah-2 resulted in a significant increase in the expression levels of 12 out of 15 LPMO genes identified in the genome of N. crassa. These data indicate that under low-oxygen conditions, SAH-2 is required for activation of the expression of genes required for sterol synthesis and adaptation to hypoxia, but the expression of genes encoding extracellular proteins, such as LPMOs, whose function requires molecular oxygen, is repressed.

**Genes involved in ER stress responses showed increased expression levels in the SREBP mutant.** Considering the substantial cellulase hyperproduction phenotype
observed in all of the SREBP pathway mutants, we asked whether SAH-2 also has a role in the modulation of protein secretion. We thus evaluated genes with expression changes (≥4-fold changes, P ≤ 0.01) in the Δsah-2 mutant RNA-seq data compared to those of the WT, as well as in the Mclr-2 and M\(^{Δsah-2}\) strains compared to those in Mclr-2 after 48 h of growth in VMM. (C) The expression levels (RPKM) of 15 currently identified N. crassa LPMO-encoding genes in the WT, Mclr-2, the Δsah-2 mutant, and M\(^{Δsah-2}\). Asterisks indicate significant differences (*, P < 0.05; ***, P < 0.001).

To search for genes that are predicted to affect protein secretion but are not regulated directly by CLR-2, we focused on the 477 genes (236 plus 241; Fig. 5A) whose expression level increased in the M\(^{Δsah-2}\) strain relative to that in the Mclr-2 strain, while 282 genes were upregulated and 585 genes were downregulated in the M\(^{Δsah-2}\) strain compared to those in Mclr-2. In these gene sets, 244 genes in the upregulated sets overlapped, while no overlap was detected in the downregulated gene sets (Fig. 5A and B; see Data Set S1).
and destination were enriched ($P = 0.01$); and 20 genes involved in cellular export and secretion were enriched ($P = 0.02$) (see Data Set S1). FunCat analysis of the set of 241 upregulated genes showed that 71 genes involved in cellular transport, transport facilities, and transport routes were enriched ($P = 0.002$) and 14 genes involved in protein folding and stabilization were enriched ($P = 0.01$). A comparison of these two gene sets revealed that a total of 176 protein secretion-associated genes showed higher expression levels in the $M^{clr-2} \Delta sah-2$ strain than in the $M^{clr-2}$ strain (see Data Set S1). In contrast, no protein secretion-associated functional categories were enriched in the set of 571 downregulated genes.

Investigation of the 176-gene set enriched in protein secretion functional categories revealed that the expression levels of many genes encoding heat shock proteins, such as $hsp30$ (NCU09364), $hsp70-5$ (NCU08693), $hsp98$ (NCU00104), and $hsp60$ (NCU01589), were significantly higher in the $M^{clr-2} \Delta sah-2$ strain than in the $M^{clr-2}$ strain. These proteins are typically involved in the UPR; activation of the UPR is an indicator of ER stress (24). We therefore evaluated the expression level of genes encoding proteins potentially involved in the UPR, including chaperones, foldases, glycosylation enzymes, vesicle transport proteins, and ER-associated degradation proteins (25, 26), in the $M^{clr-2} \Delta sah-2$ strain compared to those in the $M^{clr-2}$ strain. As shown in Table S2, the expression levels of many ER stress response genes, such as $grp-78$ (NCU03928) and $pdi-1$ (NCU09223), were $>2$-fold higher in the $M^{clr-2} \Delta sah-2$ strain than in the $M^{clr-2}$ strain. However, the expression levels of hac-1 and ire-1, whose major function is activation of the UPR pathway (26), were not increased. A key attribute used to assess activation of the UPR is the nonspliceosomal splicing reaction of the hac-1 mRNA, which results in the removal of a 23-nucleotide unconventional intron and subsequent translational activation of HAC-1 (27). We therefore used qRT-PCR to assess the expression levels of total hac-1 transcripts, spliced hac-1 transcripts, and nonspliced hac-1 transcripts with primer sets described previously (27). As shown in Fig. 6A, the expression levels of total hac-1 transcripts were significantly lower in the $M^{clr-2}, \Delta sah-2$ mutant, and $M^{clr-2} \Delta sah-2$ strains than in the WT strain. The spliced hac-1 transcript levels were also significantly lower in both the $\Delta sah-2$ mutant and $M^{clr-2} \Delta sah-2$ strains, suggesting that the activation of genes involved in ER stress and the UPR in the $M^{clr-2} \Delta sah-2$ strain may occur by an alternative pathway.

The reducing agent dithiothreitol (DTT) induces ER stress by altering the ER oxidative status and disulfide bond formation, leading to the accumulation of misfolded proteins; exposure to DTT triggers UPR in N. crassa (26, 27). To further investigate the
The upregulated UPR in \( M^{\text{clr-2}} \Delta \text{sah-2} \) is independent of \( \text{hac-1} \) regulation. (A) Relative quantification of the amount of \( \text{hac-1} \) mRNA splice variants in the WT, \( \text{Mclr-2} \), \( \Delta \text{sah-2} \) mutant, and \( M^{\text{clr-2}} \Delta \text{sah-2} \) strains by qRT-PCR from samples grown for 24 h in VMM following a shift (16-h culture grown in VMM). (B) Fold changes in the expression levels of \( \text{hac-1} \) mRNA splice variants with DTT treatment. RNA samples were obtained from cultures grown in VMM or in VMM plus 10 mM DTT for 4 h after a shift of 16-h-old VMM cultures. (C) Fold change in the expression levels of \( \text{grp-78} \) and \( \text{pdi-1} \) in cultures treated with DTT. RNA was isolated from cultures as described for panel B. The results in panels B and C are fold induction in a 4-h culture in VMM plus 10 mM DTT over the expression state in a 4-h VMM culture. (D) Growth phenotypes of the \( \text{Mclr-2} \), \( M^{\text{clr-2}} \Delta \text{sah-2} \), \( M^{\text{clr-2}} \Delta \text{hac-1} \), and \( M^{\text{clr-2}} \Delta \text{sah-2} \Delta \text{hac-1} \) strains in VMM with 2% Avicel for 96 h. (E, F) Total secreted protein level (E) and endoglucanase activities (F) of \( \text{Mclr-2} \), \( M^{\text{clr-2}} \Delta \text{sah-2} \), \( M^{\text{clr-2}} \Delta \text{hac-1} \), and \( M^{\text{clr-2}} \Delta \text{sah-2} \Delta \text{hac-1} \) strains in the supernatant of 48-h VMM cultures after a shift from 16-h VMM cultures. Protein concentration and endoglucanase activity were normalized by total fungal cell biomass in each culture. Asterisks indicate significant differences (*) \( p < 0.05 \); (**) \( p < 0.01 \); (***) \( p < 0.001 \). ns, not significant.
role of sah-2 in the UPR pathway, 16-h-old cultures of the WT, Mclr-2, Δsah-2 mutant, and M\textsuperscript{clr-2}\ Δsah-2 strains were transferred to either fresh Vogel's minimal medium (VMM) or VMM plus 10 mM DTT for 4 h. RNA was collected from these cultures, and qRT-PCR was performed to evaluate the splicing of hac-1 mRNA and the expression levels of the UPR marker genes grp-78 (NCU03928) and pdi-1 (NCU09223). In contrast to those of strains grown in VMM, the level of hac-1 transcripts was significantly induced when cultures were exposed to DTT, but the level of spliced hac-1 was much lower in the M\textsuperscript{clr-2}\ Δsah-2 strain than in the Mclr-2 strain (Fig. 6B). Both UPR marker genes grp-78 and pdi-1 were significantly induced in all four strains after treatment with DTT, but the level of induction in M\textsuperscript{clr-2}\ Δsah-2 was significantly higher than that in the other three strains (Fig. 6C).

It has been reported that a deletion of hac-1 in N. crassa results in a strain that shows a greatly reduced ability to utilize cellulose and secrete lignocellulolytic enzymes (26, 28). This observation suggested that the inability to mount an appropriate UPR results in an inability to utilize cellulose because of a dysfunction in the proper trafficking of lignocellulolytic enzymes. Because the M\textsuperscript{clr-2}\ Δsah-2 and Δsah-2 mutant strains showed an increase in the expression levels of genes involved in protein trafficking and the UPR (see Table S2) yet showed a reduction in spliced hac-1, we hypothesized that deletion of sah-2 in a Δhac-1 mutant would restore the utilization of cellulose and cellulase secretion. We first tested whether a strain carrying the Δhac-1 mutation in the Mclr-2 background (M\textsuperscript{clr-2}\ Δhac-1) failed to secrete cellulases under minimal medium conditions. Indeed, the M\textsuperscript{clr-2}\ Δhac-1 mutant showed much lower protein secretion and cellulase activity in VMM than the Mclr-2 strain (Fig. 6E and F). These data indicated that the Δhac-1 mutant’s cellulase-deficient phenotype was independent of any inducer (Avicel). We further constructed a Δhac-1 Δsah-2 mutant in the Mclr-2 background (M\textsuperscript{clr-2}\ Δsah-2 Δhac-1). In contrast to the M\textsuperscript{clr-2}\ Δhac-1 strain, and in support of our hypothesis, the M\textsuperscript{clr-2}\ Δsah-2 Δhac-1 strain showed robust protein secretion and cellulase activity that was indistinguishable from that of the M\textsuperscript{clr-2}\ Δsah-2 strain (Fig. 6E and F). The phenotype of the M\textsuperscript{clr-2}\ Δsah-2 Δhac-1 strain was also indistinguishable from that of the Mclr-2 and M\textsuperscript{clr-2}\ Δsah-2 strains under Avicel conditions but markedly different from the cellulase-deficient phenotype of the M\textsuperscript{clr-2}\ Δhac-1 strain (Fig. 6D).

**DISCUSSION**

Extensive studies of the SREBP pathway in a variety of organisms have revealed that, in addition to regulating lipid homeostasis, the SREBP pathway plays additional roles in the regulation of cell polarity, antifungal drug resistance, and virulence (10, 11, 29). Although N. crassa has homologs of the SREBP components identified in S. pombe and A. fumigatus, its role in the biology of this organism has not been extensively studied. Phenotypic analysis of strains containing mutations in SREBP components showed that this pathway is important for both mediation of the cellular response to hypoxia and regulation of ergosterol biosynthesis (Fig. 1; see Fig. S1). As shown for A. fumigatus, it is likely that scp-1, the Golgi apparatus E3 ligase Dsc complex, and the rhomboid protease-encoding gene rbd-2 are required for the activation of SAH-2 in N. crassa. However, neither the second homolog of SREBP, sre-2, nor spp-1 was required for SREBP pathway function in N. crassa. In A. nidulans, SppA, which is an aspartyl protease involved in regulated intramembrane proteolysis, is essential for adaptation to hypoxia (15), while in A. fumigatus, SrbB, which has homology to SrbA (SREBP), is also required for the response to hypoxia and for virulence (30). Our data also support a role for SREBP in the secretion of lignocellulolytic enzymes in N. crassa.

Chromatin immunoprecipitation sequencing (ChIP-seq) analysis of SrbA in A. fumigatus (30), combined with the expression pattern of ergosterol biosynthesis genes for N. crassa (Fig. 3A), supports the hypothesis that SAH-2 directly regulates erg-1, erg-3, erg-5, erg-11, and erg-25 and that the increased expression of these direct target genes is a good indicator of SAH-2 activation. The expression of these genes was significantly higher in the Mclr-2 strain than in the WT, suggesting that...
SREBP was highly activated when cellulase trafficking and secretion were induced. However, the hypersecretion phenotype in the Mclr-2 Δsah-2 strain was eliminated when the oxygen supply was increased in bioreactors (see Fig. S5C), suggesting that the production of cellulases in N. crassa might lead to a reduction of available oxygen. Importantly, comparative analyses of transcriptional profiles revealed that gene expression levels of oxygen-dependent extracellular enzymes like LPMOs, which are involved in cellulose deconstruction, were increased in the Mclr-2 Δsah-2 strain. These data indicate that activation of SAH-2 has a negative impact on the expression of extracellular-oxygen-consuming enzymes, particularly cellulases that utilize molecular oxygen, suggesting feedback regulation of genes encoding oxygen-requiring enzymes that are not essential (Fig. 7).

In addition to genes encoding LPMOs, a set of genes associated with ER stress responses had higher expression levels in the Mclr-2 Δsah-2 strain than in the Mclr-2 strain (see Table S2). This observation led us to test the interconnections among the UPR, the SREBP pathway, and cellulase secretion. Previously, it was reported that the UPR is triggered when fungal cells are exposed to recalcitrant carbon sources; that spliced hac1 transcripts increase in T. reesei (31), A. nidulans (32), and N. crassa (26, 27) when they are grown on lignocellulose; and that a Δhac-1 mutant is deficient in the secretion of cellulases (26, 28). However, our RNA-seq data analyses showed that the increased expression of UPR-associated genes observed in the Mclr-2 Δsah-2 mutant was apparently not the result of the classic IRE-1/HAC-1-mediated UPR pathway, since spliced hac-1 levels were lower in the Mclr-2 Δsah-2 strain (Fig. 6A). Additionally, of the 176 protein secretion-associated genes that showed increased expression levels in the Mclr-2 Δsah-2 strain, only 6 (NCU08707, NCU10051, NCU09485, NCU09265, NCU03982,
and NCU08607) were identified within the HAC-1 regulon in *N. crassa* (26). Our data raise the possibility that a hac-1-independent UPR pathway exists in *N. crassa* and that this pathway is repressed (at some level) by SAH-2 (Fig. 7). The hypothesis was supported by the observation that a Δhac-1 Δsah-2 mutant showed normal secretion of cellulases and utilization of cellulose. The UPR in mammals uses at least two additional ER-resident sensors that are thought to act in parallel to induce downstream UPR targets. These include ATF-6 and the ER transmembrane kinase PERK (33, 34). Even though homologs of ATF-6 or PERK are not present in the genomes of yeast or filamentous fungi, our data suggest the presence of a second regulatory mechanism for the fungal UPR pathway that involves SAH-2. In mammalian cells, ER stress induces SREBP activation, although molecular mechanisms associated with cross talk between these two pathways may occur by several mechanisms that are not well elucidated (35). Additionally, it raises the question of the relationship of the UPR and the SREBP pathway for functions in filamentous fungi not associated with sterol homeostasis, such as virulence. Future characterization of the regulatory network of the UPR and SREBP pathways will be very informative for deciphering cellulase trafficking and lignocellulose deconstruction in filamentous fungi. This knowledge can benefit rational strain engineering to improve the production of lignocellulose-degrading enzymes and other biomass-based bioproducts.

**MATERIALS AND METHODS**

**Strains and strain construction.** *N. crassa* WT strain OR74A (FGSC2489) and available single-gene deletion mutants were obtained from the Fungal Genetics Stock Center (http://www fgsc net/). The Δscp-1 mutant was constructed by transforming WT strain OR74A with a 3.8-kbp DNA fragment containing a 1.3-kbp flanking region homologous to the sequence upstream of scp-1, a selectable marker (tpc promoter-driven hygromycin B phosphotransferase) (16), and a 1.1-kbp flanking sequence homologous to the downstream region of scp-1. Transformants were selected for resistance to hygromycin and tested for genotypes by diagnostic PCR with primers Vscp1-F (GATAGTAGCTTTGGGAGATATTGC) and hph-R (GGGGCATACACGGCTTATTGGACCT). The Δsre-2 mutant was constructed by transforming WT strain OR74A with a 3.7-kbp DNA fragment containing a 1.1-kbp flanking region homologous to the sequence upstream of sre-2, the hygromycin B-selectable marker, and a 1.2-kbp flanking region homologous to the downstream region of sre-2. Transformants were selected for resistance to hygromycin and tested for genotypes by diagnostic PCR with primers VsreB-F (CTTGACCTGCTCGACGAAAACACAC) and hph-R (GGGGCATACACGGCTTATTGGACCT). The Δerg-25 strain was constructed by transforming the Mcl-2 (his-3::Pcc-1-2 rdd-1 Δasd-1) strain (17) with a 3.4-kbp DNA fragment containing a 1.2-kbp flanking region homologous to the sequence upstream of erg-25, the tpc promoter-driven nourseothricin acetyltransferase nat-1 gene from Streptomyces noursei (36), and a 1.2-kbp flanking region homologous to the downstream region of erg-25. Transformants were selected for resistance to nourseothricin and tested for genotypes by diagnostic PCR with primers Verg25-F (GTGATGATCTAGAACGAGTTGAT) and nat-R (GGGTAAAGCCACCCTCAG). Fifty micromolar SCR7 (37), which is an inhibitor of nonhomologous end joining, was used to increase the homologous recombination ratio of all transformations. Positive transformants were backcrossed to FGSC4200 to obtain Δscp-1 mutant, Δsre-2 mutant, and M**inh**-2 Δerg-25 homokaryotic strains.

The M**inh**-2 Δhac-1 strain was created by transforming the Mcl-2 strain with a 3.4-kbp DNA fragment containing a 1.2-kbp flanking region homologous to the sequence upstream of hac-1, a tpc promoter-driven nourseothricin acetyltransferase nat-1 gene, and a 1.2-kbp flanking region homologous to the downstream region of hac-1. The M**inh**-2 Δhac-1 Δsah-2 strain was created by transforming the same fragment described above into the M**inh**-2 Δsah-2 strain. SCR7 (50 μM) (37) was used for these two transformations. Transformants were selected for resistance to nourseothricin and tested for genotypes by diagnostic PCR with primers Vhac-1-F (TGGATGCTTTGGGAGATATTGC) and nat-R (GGGTAAAGCCACCCA CTCAG). Homokaryotic M**inh**-2 Δhac-1 and M**inh**-2 Δhac-1 Δsah-2 strains were obtained by microconidial purification as described elsewhere (http://www fgsc net/fgs 37/ebbole1.html). The M**inh**-2 Δsah-2, M**inh**-2 Δsah-1, M**inh**-2 Δsah-1 Δsah-2, and M**inh**-2 Δerg-24 strains were generated by crosses.

**VMM (38) was used to grow N. crassa.** Conidia were inoculated into 100 ml of liquid medium at 10⁶ conidia/ml and grown at 25°C on a rotary shaker (200 rpm) for the times indicated in constant light. Synthetic cross medium (39) was used for crosses. For hypoxic conditions, a hypoxia incubation chamber (Invivo2; Ruskinn) was used. The chamber was maintained at 25°C and filled with N₂ with 0.5% O₂.

**Protein and enzyme assays.** The protein concentration in supernatants was determined by the Bradford method (Bio-Rad Protein Assay). Endoglucanase activity was determined by with azo-carboxymethyl cellulose (Azo-CM-Cellulose; Megazyme) in accordance with the manufacturer’s specifications. CDH-1 activity was measured by monitoring the time-dependent reduction of 300 μM 2,6-dichloroindophenol (DCIP) at a wavelength of 520 nm in 100 mM sodium acetate, pH 5.0, containing 30 mM cellobiose (22). The reaction was started by adding 180 μl of the DCIP-based assay solution to a 20-μl supernatant sample in 96-well plates and monitored in a plate reader at 30°C for 5 min.
One unit is equivalent to the conversion of 1 μmol min⁻¹. CBH-1 activity was measured with soluble 4-methylumbelliferyl-β-D-cellobiose (Sigma) as the substrate as previously described (40).

**Sterol extraction and GC-MS analysis.** Fungal biomass from cultures was filtered and dried in a 65°C oven for 2 days. One hundred milligrams of dry biomass of each sample was ground into powder for extraction of sterol, and GC-MS was used to analyze the sterol components by suspension in a solution consisting of 4 ml of methanol and 2 ml of 60% (wt/vol) KOH supplemented with 10 μg of cholesterol as a recovery internal standard. Esters were saponified by incubation at 75°C with agitation for 4 h. Nonsaponifiable sterols were extracted twice with 4 ml of petroleum ether and evaporated with a stream of nitrogen gas to dry. The dry extract was resuspended in 1 ml of toluene. A 50-ml sample was added to 50 μl of N,O-bis(trimethylsilyl)trifluoroacetamide and incubated at 70°C for 30 min. The trimethylsilyl ether derivatization generated by this step was used for GC-MS analysis (41).

GC-MS analyses were performed with a CP-3800 gas chromatograph equipped with a FactorFour capillary column (VF-5 ms; 15-m length, 0.32-mm inside diameter, 0.25-μm film thickness) and analyzed with a Varian 320-MS triple-quadrupole mass spectrometer. Electron impact MS (70 eV, scanning from 80 to 600 at 1-s intervals) was performed with He as the carrier gas (30 cm/s), a detector temperature of 160°C, and column temperatures of 200 to 330°C (200°C for 1 min, 20°C/min to 330°C, and then held for 3 min). All injections were run in a splitless mode, and the injection volume was 1 μl. Sterol species were identified by comparison with standards or with MS data in the library. Peak areas for each sterol were normalized to the internal standard cholesterol peak area to correct for recovery.

**RNA-seq and data analysis.** Mycelial biomass from 24-h VMM cultures previously transferred from a 16-h VMM culture was filtered and frozen in liquid nitrogen for RNA extraction. RNA-seq libraries were prepared in accordance with standard protocols from Illumina Inc. (San Diego, CA) and sequenced on the Illumina Genome Analyzer Ix and HiSeq 4000 platforms at the genome center of the University of California Davis. Raw RNA sequencing reads (fastq files) were first subjected to FASTX toolkit (http://hannonlab.cshl.edu/fastx_toolkit/) to remove adaptor contamination and check quality. Reads passing this filter were mapped against N. crassa predicted transcripts from the OR74A genome v 12 (Neurospora crassa Sequencing Project, Broad Institute of Harvard and MIT [http://www.broadinstitute.org/]) with Tophat v 2.0.4. Numbers of RPKM were calculated with Cufflinks v 2.2.1, and differential expression analyses were performed with DESeq2 package version 1.10.1. The cluster 3.0 software suite (http://bonsai.hgc.jp/~mdehoon/software/cluster/software.htm) was used to perform hierarchical clustering analysis. The Munich Information Center for Protein Sequences FunCat online tool (http://mips.helmholtz-muenchen.de/functcatDB/) was used for functional classification of genes of interest.

**Bioreactor fermentation.** Conidia at 10⁶/ml were inoculated into a parallel 1-liter stirred tank bioreactor (DASGIP Bioreactor system, type DGCS4; Eppendorf AG, Germany) containing 500 ml of VMM and incubated for 48 h. The pH was set at 5.5 by the addition of 1 M HCl or 1 M KOH solution, and the temperature was maintained at 25°C. The DO tension was initially calibrated at 100%, and the DO level was controlled by manipulating the airflow or the stirrer speed. For a lower DO supply, the aeration flow rate was set at 30 liters/h and the stirrer speed was set as 400 rpm. For a higher DO supply, an automatic DO control system was used, with an aeration flow rate of up to 50 liters/h and a stirrer speed of up to 800 rpm when the DO level was lower than 40% of that at the beginning. During fermentation, 3-ml samples were taken every 12 h for measurement of the secreted protein concentration and enzyme activity.

**Statistical significance tests.** Statistical significance was determined by t-test analysis by the false discovery rate (FDR) approach. Asterisks indicate significant differences (*, P < 0.05; **, P < 0.01; ***, P < 0.001). ns, not significant.

**Accession number(s).** The data discussed in this publication have been deposited in NCBI’s Gene Expression Omnibus (42) and are accessible through GEO Series GenBank accession no GSE95807.

**SUPPLEMENTAL MATERIAL**

Supplemental material for this article may be found at https://doi.org/10.1128/mBio.00233-17.

- **FIG S1,** TIF file, 2.6 MB.
- **FIG S2,** TIF file, 0.8 MB.
- **FIG S3,** TIF file, 1.9 MB.
- **FIG S4,** TIF file, 2 MB.
- **FIG S5,** TIF file, 1.6 MB.
- **TABLE S1,** DOCX file, 0.02 MB.
- **TABLE S2,** DOCX file, 0.03 MB.
- **DATA SET S1,** XLSX file, 0.6 MB.

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