Transcription Activator Swi6 Interacts with Mbp1 in MluI Cell Cycle Box-Binding Complex and Regulates Hyphal Differentiation and Virulence in Beauveria bassiana

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Abstract: Mbp1 protein acts as a DNA-binding protein in MluI cell cycle box-binding complex (MBF) and plays an essential role in filamentous myco-pathogen Beauveria bassiana. In the current study, BbSwi6 (a homologue of yeast Swi6) was functionally characterized in B. bassiana. Both BbSwi6 and BbMbp1 localize in the nucleus and display a direct interaction relationship which is indicated by a yeast two-hybrid assay. BbSwi6 significantly contributes to hyphal growth, asexual sporulation and virulence. On the aerial surface, ∆BbSwi6 grew slower on various nutrients and displayed abnormal conidia-producing structures, which hardly produced conidia. In liquid media, BbSwi6 loss led to 90% reduction in blastospore yield. Finally, the virulence of the ∆BbSwi6 mutant was modestly weakened with a reduction of 20% in median lethal time. Comparative transcriptomics revealed that BbSwi6 mediated different transcriptomes during fungal development into conidia and blastospores. Notably, under the indicated condition, the BbSwi6-mediated transcriptome significantly differed to that mediated by BbMbp1. Our results demonstrate that, in addition to their roles as the interactive components in MBF, BbSwi6 and BbMbp1 mediate divergent genetic pathways during morphological transitions in B. bassiana.

Keywords: MBF complex; development; conidiation; dimorphism; pathogenicity; insect myco-pathogen

1. Introduction
Filamentous fungi have evolved a conidiation process to produce numerous conidia for dispersal and survival in their environment [1]. Beauveria bassiana is a filamentous entomopathogenic fungus that naturally parasitizes insects, and therefore has a great potential for pest management [2]. Conidia attach to the host cuticle and germinate into invasive mycelia via mobilizing endogenous nutrients, which initiates the infection process [3,4]. Once entering into the insect hemocoel, B. bassiana develops into yeast-like cells (in vivo hyphal bodies) via dimorphic transition. Hyphal bodies assimilate the nutrients from hemolymph and propagate rapidly. At the late stage of infection, hyphal bodies transform into robust mycelia which cause the host to be mummified [5]. Finally, fungal hyphae grow outside of the cadaver and generate numerous conidia for the sequential infection cycle [6,7]. Therefore, conidiation process is critical for the consecutive infection cycle of B. bassiana. The cell-division cycle is essential for fungal proliferation [8]. Previous studies have revealed that the cell-division cycle mechanism is involved in development and morphological transition during B. bassiana lifecycle [9,10].

In Saccharomyces cerevisiae, this cycle included four phases, i.e., genome duplication (S-phase), nuclear division (M-phase) and two gap phases (G1 and G2). Two transcription-factor complexes (Swi4/Swi6 (SBF) and MluI (MBF) cell cycle box binding complex) possess comprehensive transcriptional regulation at G1/S transition (a checkpoint of cell-division cycle) [11,12]. In S. cerevisiae SBF and MBF, Swi6 acts as a transcriptional activator and recruits Swi4 and Mbp1, respectively [13,14]. However, the MBF components differ between

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yeast species. In *Candida albicans*, Swi4 and Swi6 are necessary for the phase transition of G1/S, but SBF lacks the binding sites in the genome. Hence, these two factors may be involved in MBF activity [13]. In *Schizosaccharomyces pombe*, MBF activates transcription of genes associated with G1/S transition and consists of Cdc10 (the ortholog of *S. cerevisiae* Swi6) and two DNA binding factors (i.e., Res1 and Res2) [16,17]. Furthermore, the MBF components play different roles in fungal physiology. In budding yeast, although cells without Mbp1 or Swi4 are nonviable, the Swi6 loss does not influence cell viability [11,12]. Our recent study has revealed that Mbp1 significantly contributes to the conidiation process in *B. bassiana*, promoting the understanding of the Mbp1 roles in filamentous fungi [10]. In the filamentous fungi *Magnaporthe oryzae* and *Fusarium graminearum*, Swi6 functions in growth as well as in both sexual and asexual development [18,19]. In *Metarhizium rileyi* (also an insect pathogenic fungus), the role of Swi6 in fungal conidiation has been examined [20], but the functions of the Swi6 homolog remain largely unknown in other entomopathogenic fungi. Although Mbp1 and Swi6 have some overlaps in fungal development and virulence, the question regarding their differential regulatory functions is still open.

To improve understanding of the differential roles of the different MBF components in the filamentous fungi, *B. bassiana* was used a representative organism. In this study, *B. bassiana* Swi6 (BbSwi6) was identified and verified as interacting with BbMbp1. Functional analyses indicated that BbSwi6 was essential for fungal development and virulence in *B. bassiana*. During the conidiation and blastospore production process, comparative transcriptomic analysis was used to reveal the BbSwi6-mediated transcriptome which partially overlaps with the BbMbp1-mediated one [10]. We further found that the overlapped downstream 928 genes played an important role in aerial conidiation and 520 genes played a significant role in blastospore production. Our results highlight that, in addition to acting as the MBF components, Swi6 and Mbp1 display significantly functional differentiation during fungal development.

2. Materials and Methods

2.1. Strains and Growth Conditions

Fungal strains were cultured as previously described [10]. The wild-type *B. bassiana* ARSEF2860 (WT) (USDA-ARS Collection of Entomopathogenic Fungal Cultures, Ithaca, NY, USA) was cultivated on Sabouraud dextrose agar (SDA: 4% glucose, 1% peptone and 1% yeast extract plus 1.5% agar). *Escherichia coli* DH5α (Invitrogen, Carlsbad, CA, USA) was cultured in Luria-Bertani (LB) medium for propagation of plasmids. *Agrobacterium tumefaciens* AGL-1 was cultured in YEB broth (w/v: 0.5% sucrose, 1% peptone, 0.1% yeast extract and 0.05% MgSO₄) and used in fungal transformation. Czapek-Dox agar (CZA) (3% glucose, 0.3% NaNO₃, 0.1% K₂HPO₄, 0.05% KCl, 0.05% MgSO₄ and 0.001% FeSO₄ plus 1.5% agar) was used to screen the transformants and phenotypic assay.

2.2. Bioinformatic Identification and Functional Analyses of BbSwi6

The sequence of *S. cerevisiae* Swi6 (GenBank No.: AJV59711) was used as a query to identify the homolog in the *B. bassiana* genome [21]. The resulting *B. bassiana* homolog (Locus tag: BBA_05428) was designated as BbSwi6, and then its entire ORF as well as the up- and downstream flanking sequences were identified by mapping the cDNA sequence of BbSwi6 onto the fungal genomic sequence.

Fluorescence-coupled double screening method was used in gene disruption and complementation [22,23]. All primers are given in Table S1. The up- and downstream fragments were amplified by primer P1/P2 and P3/P4, respectively, and individually cloned into the XmnI/BamHI sites and XbaI/SphI sites in plasmid p0380-GTB in which bar cassette confers ammonium glufosinate resistance. The restriction enzymes were purchased from New England BioLabs, Inc. (Beijing, China). The resultant plasmid was named p0380-BbSwi6-KO, which was transformed in the WT strain. The 8-d old conidia were used as the receptor cells in fungal transformation. The transformants were grown on CZA supplemented with phosphinothricin (200 µg/mL) (45520, Sigma, MO, USA).
The putative gene disruption mutant was screened by PCR reaction with primer P5 and P6, and was further verified under a laser scanning confocal microscope [23]. For gene complementation, the entire \textit{BbSwi6} together with the promoter and terminator regions was amplified with the primer P7 and P8. The DNA fragment was cloned into the vector p0380-sur-gateway (conferring resistance to chlorsulfuron (C11325000, Dr. Ehrenstorfer GmbH, Augsburg, Germany)), generating the plasmid p0380-sur-BbSwi6. The resultant transformants were screened by PCR with the primer pair P5/P6 and further confirmed by qRT-PCR with the primer pair P9/P10.

2.3. Sub-Cellular Localization of \textit{BbMbp1} and \textit{BbSwi6}

The intracellular locations of these two proteins were determined by fusing their nuclear location signal sequences to the fluorescent protein gene. Primers are listed in Table S1. Firstly, the promoter region of translation elongation factor 1 alpha gene (\textit{TEF1}) was amplified with primer pair P11/P12. The DNA fragment was cloned into the EcoRI/BamHI sites of p0380-sur [24], generating p0380-TEF-sur. Then, green fluorescent protein gene (GFP) fragments were amplified with the primer pair P13/P14 and cloned into the BamHI sites of p0380-BbTEF-sur, generating pBBTEF-MCS-Gfp-sur (pBMGS). Similarly, the sequence of \textit{mCherry} was amplified with the primer pair P15/P16, and cloned into the BamHI sites of p0380-BbTEF-sur, generating pBBTEF-MCS-mCherry-sur (pBMRS).

The nuclear location signal sequences were predicted by DTU Bioinformatics (http://www.cbs.dtu.dk/services/) (accessed on 6 September 2020). The first 645 bp of \textit{BbMbp1} were amplified with primer pair P17/P18, and cloned into pBMGS, generating pBMGS-BbMbp1. Similarly, the first 891 bp of \textit{BbSwi6} were amplified with primer pair P19/P20, and integrated into pBMRS, generating pBMRS-BbSwi6. The resulting constructs were individually transformed into the WT strain. The strains expressing these fused genes were grown in Sabouraud dextrose broth (SDB). Two days later, fluorescent signals in mycelia were examined with a fluorescence microscope. Nuclei were stained with 4′,6-diamidino-2′-phenylindole dihydrochloride (DAPI) (D9542, Sigma, MO, USA).

2.4. Quantification of Blastospore Production

Phenotypic assay was conducted among the WT, gene disruption and complementation strains as described previously [10]. The blastospore formation was evaluated in SDB medium under submerged condition at 25 °C. Mycelia of the indicated strain were cultured in SDB medium and the resultant spores were used as initial inocula. Blastospore suspension was inoculated into fresh SDB medium and the final concentration was adjusted to \(10^6\) spores/mL. Three days later, spore concentration in broth was quantified and blastospore yield was shown as spore number per milliliter.

2.5. Assay for Conidial Production

Conidiation on aerial surfaces was examined as described previously [10]. Inocula (in vitro blastospore) were prepared by culturing mycelia in SDB at 25 °C for 2 d. Aliquots of 100 µL blastospore suspension (\(10^7\) spores/mL) were cultured on SDA plates and incubated at 25 °C. Mycelial discs were sampled at 4, 6 and 8 d post incubation (DPI). The conidia on mycelia were quantified and calculated as conidial number per cm\(^2\). In addition, the 3- and 8-d old mycelia were dried and weighed. The dry weight was presented as milligrams per square centimeter.

2.6. Fungal Growth

To evaluate the nutrient effects on vegetative growth, the carbon and nitrogen sources in CZA were replaced as necessary. Carbon sources (at a final concentration of 3%) included glucose, sucrose, trehalose and glycerol. Nitrogen sources (at a final concentration of 0.5%) included \(\text{NH}_4\text{Cl}, \text{NH}_4\text{NO}_3\), peptone and gelatin. One microliter of blastospore suspension (\(10^6\) spores/mL) was inoculated on the modified CZA plates. After 7 days of incubation, the colony diameter was measured.
2.7. Insect Bioassay

Fungal virulence against *Galleria mellonella* larvae (bioassay insect) was evaluated with an intra-hemocoel injection method. The insects were fed with artificial diet and reared as previously described [25], and last-instar larvae (~300 mg in weight) were used. Aliquots of 5-µL of blastospore suspension (10^5 spores/mL) were injected into the larval hemocoel. The experiment included three replicates, and approximately 35 larvae were used in every replicate. Control group was injected with Tween-80 solution (0.02%). The mortality was recorded daily and subjected to Probit analysis for the median lethal time (LT50). Cadavers were kept in a moisture chamber, and mycose on cadaver was recorded at 4 d post incubation.

2.8. Yeast Two Hybrid (Y2H) Assays

Protein-protein interactions were assayed with the Matchmaker® Gold Yeast Two-Hybrid System (TaKaRa, CA, USA). All primers were listed in Table S1. The BbMbp1 ORF was amplified from cDNA template with primers P21/P22 and cloned into the pGADT7 vector. The resulting vector was confirmed by sequencing and transformed in pairs into yeast strain Y187 that was screened on selective (SD/-Leu) medium. The BbSwi6 ORF was obtained by PCR amplification with primer pair P23/P24 and cloned into the pGBK17 vector. The vector was transformed into yeast strain Y2Gold and screened on selective (SD/-Trp) medium. These two strains were mated in YPDA medium and screened on the selective dropout/-tryptophane-leucine-adenine-histidine medium (SD-Trp-Leu-His-Ade) containing 200 ng/mL aureobasidin A (AbA) (600466, TaKaRa, Dalian, China).

2.9. Analyzing the BbSwi6-Mediated Transcriptome during Sporulation

All methods for transcriptomic analysis were the same as those used in revealing the BbMbp1-mediated transcriptome [10]. Due to the presence of the transcriptome in the WT strain, only ∆BbSwi6 mutant strain was subjected to RNA-seq analysis. Blastospores were prepared as mentioned above and used as initial inocula. Aerial mycelia were cultured on SDAY plates for 3 d, and submerged mycelia were grown in SDB medium for 2 d. Total RNA was extracted from the mycelial sample (Trizol method). The mRNA molecules were purified by using magnetic oligo(dT) beads and used as template to synthesize first-strand cDNA using random hexamer primers. Second-strand cDNA synthesis was performed in reaction buffer containing deoxyribonucleotide triphosphates and DNA polymerase I. Double-strand DNA was purified and constructed into a library which was analyzed on the Illumina HiSeq X Ten platform at Vazyme Biotech Co., Ltd. (Nanjing, Jiangsu, China). The sequence data have been deposited in the NCBI Gene Expression Omnibus (accession No. GSE171375). Two independent libraries were constructed for each treatment.

All clean reads were mapped onto the Bb2860 genome using the HISAT program [26]. The resultant genes were normalized in terms of the expected number of fragments per kilobase of transcript sequence per millions base pairs sequenced (FPKM), using Cufflinks software [27]. The transcriptomic data for the WT strain were downloaded from Gene Expression Omnibus (accession No.: GSE134764) [10]. The Cuffdiff method was used to search the differentially expressed genes (DEGs) between the WT and ∆BbSwi6 mutant strains. The DEGs were considered when an absolute value of log2Ratio (fold change) > 1 at the threshold of the q-value < 0.05 was obtained (5% false discovery rate) [28]. The resulting DEGs were subjected to enrichment analysis, using the online FungiFun2 portal (https://elbe.hki-jena.de/fungifun/) (accessed on 6 September 2020). The threshold of the corrected p-value was set at 0.05 [29].

To determine whether BbMbp1 and BbSwi6 have different transcriptional influences during conidiation, we compared the BbSwi6- and BbMbp1-mediated DEGs [10].

2.10. Data Analyses

All measurements for the WT, gene disruption and complementation mutant strains, including the colony diameter and sporulation capacity were subjected to two-way ANOVA.
The measurements for LT<sub>50</sub> were subjected to one-way ANOVA. Significant differences were determined with Tukey’s honest significance test (Tukey’s HSD).

3. Results

3.1. Bioinformatic Analysis of BbSwi6 and Generation of Its Gene Disruption and Complementation Strains

Based on a BLAST search with a yeast homolog as the query, a single highly related <i>B. bassiana</i> gene, BBA_05428 (identity: 28.80%; E-value: 7 × 10<sup>−65</sup>), was identified and named BbSwi6. The open reading frame (ORF) sequence of this gene was 2433 bp long and had one intron. BbSwi6 coded for a protein with 810 amino acids.

To further examine BbSwi6 roles, the gene disruption strain was successfully constructed through the homologous recombination strategy, and the partial ORF of BbSwi6 was replaced by the phosphinothricin resistance gene (<i>bar</i>) (Figure S1A). To construct the complementation strain, the whole BbSwi6 ORF plus the promoter region was ectopically integrated into the BbSwi6 mutant strain, using the sulfonylurea resistance gene (<i>sur</i>) as the selection marker. The resulting disruption and complementation mutant strains were first screened by PCR and further verified by fluorescence detection and qRT-PCR analysis (Figure S1B–D).

3.2. BbSwi6 Interacts with BbMbp1 in Nucleus

Intracellular localization of BbSwi6 and BbMbp1 in <i>B. bassiana</i> was validated by fusing the coding sequence for nuclear targeting signal to the N-terminus of mCherry and green fluorescent protein gene (GFP), respectively. Nucleus was visualized by blue fluorescence using 4′,6-diamidino-2-phenylindole (DAPI, a nucleus-specific dye). Under a laser scanning confocal microscope, globular red and green signals were seen in the transformant expressing the fusion gene and were co-localized with the blue fluorescence well (Figure 1A). These results indicated that BbSwi6 and BbMbp1 localized in nuclei.

Figure 1. Sub-cellular analyses of the Swi6 and Mbp1 protein in <i>B. bassiana</i>. (A) The targeting signal sequences of Swi6 and Mbp1 were fused to mCherry and green fluorescent protein gene (GFP), respectively. The hybrid genes were transformed into the wild-type strain. Mycelia were stained with nucleus-specific dye DAPI. The overlapped fluorescent signals indicated that both Swi6 and Mbp1 localize in nucleus. (B) Yeast two-hybrid test indicated that Swi6 and Mbp1 have physical interaction. Only yeast cells with Swi6 and Mbp1 grew the selective media well. Negative and positive controls (NC and PC) were provided by kit. Scale bar: 10 μm.
Yeast two hybrid (Y2H) assay was used to examine the interaction of BbSwi6 and BbMbp1. All constructs were valid as indicated by the yeast growth on the selective dropout (SD)-Leu-Trp medium. The mated transformants from the pair BbSwi6/BbMbp1 grew well on the SD-Leu-Trp-His-Ade medium, which indicating that BbSwi6 interacted with BbMbp1 directly (Figure 1B).

3.3. BbSwi6 Significantly Contributes to Vegetative Growth on Nutrients

As illustrated in Figure 2A,B, ΔBbSwi6 mutant strain showed different growth trends on the minimal medium included different carbon and nitrogen sources. Colony diameters differed significantly among five strains ($F_{4,60} = 1130.0, p < 0.01$) and six nutrients ($F_{5,60} = 769.3, p < 0.01$). Compared with the WT strain, the severe growth defects of ΔBbSwi6 were observed on four carbon sources (glucose, trehalose, fructose and sucrose), and slight reduction in colony size of ΔBbSwi6 occurred on oleic acid and gelatin. In addition, the BbMbp1 loss led to more significant impairment in colony size on all tested nutrients. For example, on the medium with glucose as carbon source, the colony diameter of the WT strain was 2.43 ± 0.05 cm, and that of ΔBbMbp1 was 1.20 ± 0.0 cm, which decreased 50.68%. The colony diameter of ΔBbSwi6 was 1.77 ± 0.05 cm, which only decreased by 27.40% when compared with the WT strain.

3.4. BbSwi6 Is Required for Asexual Sporulation

On aerial surface (Figure 3A,C), the WT strains started conidiation at day 3 by formation of conidiophores and produced plenty of conidia at day 8 with a yield of $6.95 \pm 0.32 \times 10^8$ conidia/cm$^2$. In contrast, ΔBbSwi6 mutant strain, at 3 days post incubation, generated abnormal spore-producing structures. Even at day 8, only very few conidia were detected in ΔBbSwi6 mutant strain. Its conidial yield was $0.03 \pm 0.01 \times 10^8$ conidia/cm$^2$ and displayed a reduction of 99.51% when compared with the WT strain. There was significance in conidial yield among three strains ($F_{2,18} = 1160.8, p < 0.01$) and at three time points ($F_{2,18} = 371.9, p < 0.01$).

As shown in Figure 3B,D, in SDB medium, the WT and complemented strains generated numerous branch-like spore-formation structures on the mycelia. However, the blastospore-forming structures (morphology and number) were significantly changed in ΔBbSwi6 mutant strain when compared with that of the WT strain. After 2 days of culture in SDB, ΔBbSwi6 displayed a significantly reduced blastospore yield ($0.116 \pm 0.012 \times 10^8$ spores/mL), whereas the WT and complemented strains generated $1.18 \pm 0.08$ and $1.20 \pm 0.08 \times 10^8$ spores/mL, respectively.
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As shown in Figure 3B,D, in SDB medium, the WT and complemented strains generated conidiophores at 3 dpi and produced numerous conidia. However, ΔBbSwi6 mutant did not form normal conidiophores till 8 dpi. The enlarged conidiophores were shown. (B) In SDB medium, disruption of BbSwi6 caused a slight change in morphology of the blastospore-producing structures at 3 dpi. (C) Conidal production was determined at 8 dpi. “ND” means that no detectable conidia were observed. (D) Blastospore production was examined at 3 dpi. Gene loss significantly compromised the spore production on aerial surface and in submerged broth. The different lowercase letters show the significance among three strains at the indicated time point, and the capital letters indicate the significance among different time points for the indicated strain (Tukey’s HSD: $p < 0.05$). Scale bar: 10 µm.
and ΔBbSwi6 strain decreased. On the second day, the yield of ΔBbSwi6 decreased by 90.1% and on the third day, the decrease in yield was only 81.5%.

3.5. Disruption of BbSwi6 Significantly Affected Fungal Pathogenicity

Injection bioassay was performed to examine the virulence of the ΔBbSwi6 mutant against insect hosts. The accumulative mortality for ΔBbSwi6 mutant did not significantly differ with that of the WT strain, but the mortality trend has been significantly delayed (Figure 4A). Cumulative mortalities for three strains were 100%. ΔBbSwi6 mutant strain displayed an LT50 of 3.65 ± 0.036 d, with a delay of approximate 20%, when compared to those for the WT (3.07 ± 0.01 d) and complemented strain (3.14 ± 0.05 d) (Figure 4B) 

(F2,6 = 143.2, p < 0.01). In the host hemolymph, gene loss resulted in a significant reduction in vivo blastospores, although no morphological difference was found between the ΔBbSwi6 mutant and the WT or complemented strain (Figure 4C). At 3 d post infection, ΔBbSwi6 mutant strain only produced 1.50 ± 1.00 × 10⁶ spores/mL, with an approximate 88% reduction, when compared with that of the WT strain (1.30 ± 0.27 × 10⁷ spores/mL). The spore yield of the ΔBbSwi6 strain increased at 3 days post infection but still displayed an approximate 63.3% of WT (Figure 4D). After developing in moist boxes for 5 days, ΔBbSwi6 mutant strain generated few mycelia on cadavers compared with the dense mycelia of the WT and complemented strains (Figure 4E). This result indicated that fungal virulence and growth on cadaver had been significantly impaired in ΔBbSwi6 mutant.

Figure 4. Fungal virulence. Blastospores of the indicated strain were prepared as the initial inocula. Spore suspension was injected into the hemoceol of Galleria mellonella larvae, and survival percentage was recorded daily. Compared with the wild-type strain (WT), ΔBbSwi6 strain displayed a delayed survival curve (A) and an increased median lethal time (LT50) (B). Significant differences are designated the lowercase letters (Tukey’s HSD: p < 0.05). Gene loss impaired fungal dimorphism in the host hemoceol (C), and the production of in vivo blastospores was significantly reduced (D). HB: in vivo hyphal body. HC: insect hemocytes. (E) ΔBbSwi6 strain displayed a significantly compromised ability to form mycoses on cadaver under moisture condition. The different lowercase letters show the significant difference among five strains at the same time point, and the capital letters indicate the significance between two time points for the indicated strain (Tukey’s HSD: p < 0.05). Scale bar: 10 µm.
3.6. BbSwi6 Mediates the Genome-Wide Expression during Fungal Differentiation

Disruption of BbSwi6 significantly changed B. bassiana transcriptomes during conidial and blastospore formation. On aerial SDAY plates, the BbSwi6 loss led to the altered expression of 1770 genes in which 783 genes were up-regulated (~7.6% of the genome) and 987 gene were down-regulated (~9.5% of the genome) (Table S2). In SDB medium (submerged condition), the BbSwi6 transcriptome significantly differed with that of WT strain, which including 398 up-regulated genes (~3.8% of the genome) and 666 down-regulated genes (~6.4% of the genome) (Table S3).

Transcriptomic analyses indicated that BbSwi6 loss had different effects during fungal differentiation on aerial surface and under submerged conditions (Figure 5A). Under aerial condition, the down-regulated genes (DRG) were enriched in four functional categories (i.e., metabolism, cell transport, cellular rescue and interaction with environment) (Figure 5A top panel and Table S4). The set of DRG represented in metabolism included a number of genes involved in nitrogen, aromate, vitamins and secondary metabolism. Enriched DRGs associated with cellular transport included a set of genes involved in transportation of heavy metal ion, amino acid, vitamins and so on. Repressed genes involved in cell rescue included many cytochrome P450 genes (detoxification pathway). Enriched DRGs involved in interaction with environment included a number of genes critical for homeostasis of metal ions and cellular sensing to external stimulus. The up-regulated genes (URG) in the mutant were largely enriched in metabolism, energy, protein synthesis, cell transport and cellular rescue (Figure 5A top panel and Table S5). For instance, the metabolism-associated URGs included the genes critical for the metabolism of amino acids (e.g., isoleucine, valine and leucine) (Figure 5A bottom panel and Table S7).

Compared with the BbMbp1-mediated transcriptome, the BbSwi6-mediated transcriptome had 928 and 520 differentially expressed genes under aerial (Figure 5B and Table S8) and submerged conditions (Figure 5C and Table S9), respectively. However, these genes showed significantly different distributions and expression patterns, and most overlapped genes were repressed in both gene disruption mutants.

Figure 5. Functional distribution analyses of transcriptomic data. (A) Differentially expressed genes (DEGs) were individually determined in the ΔBbSwi6 mutant via comparison with the wild-type strain during conidiation and blastospore formation. The up- and down-regulated DEGs were subjected to functional distribution analyses. BbSwi6 mediates different physiological terms in two types of asexual development. Red and green indicate the up- and down-regulated DEGs, respectively. Further, BbSwi6 and BbMbp1 differentially mediate the transcriptome during conidiation on aerial surface (B) or blastospore formation in broth (C).
Under submerged condition, the DRGs were largely associated with metabolism and energy (Figure 5A bottom panel and Table S6). The metabolism-associated DEGs included genes involved in secondary metabolism (e.g., fructosyl amino acid oxidase, Cytochrome P450 and ubiquinone biosynthesis monooxygenase), biosynthesis of vitamin (e.g., torulene oxygenase and folylpolyglutamate synthase) and so on. The enriched categories for URGs involved metabolism, energy, cell cycle and so on, of which five categories (e.g., cell cycle and cellular rescue) were not enriched in the DRGs. The metabolism-associated URGs included the genes critical for the metabolism of amino acids (e.g., isoleucine, valine and leucine) (Figure 5A bottom panel and Table S7).

Compared with the BbMbp1-mediated transcriptome, the BbSwi6-mediated transcriptome had 928 and 520 differentially expressed genes under aerial (Figure 5B and Table S8) and submerged conditions (Figure 5C and Table S9), respectively. However, these genes showed significantly different distributions and expression patterns, and most overlapped genes were repressed in both gene disruption mutants.

4. Discussion

In yeast, transcriptional activator Swi6 plays a critical role in cell cycle and differentiation via recruiting Mbp1 and Swi4 to form the MBF and SBF complexes [30,31]. In filamentous fungus B. bassiana, a Swi6 ortholog was characterized. BbSwi6 contains three ANK domains. Our results demonstrate that BbSwi6 directly interacts with BbMbp1 in the nucleus, and plays important roles in growth, development and virulence in B. bassiana.

BbSwi6 is significantly involved in fungal growth, development and virulence of B. bassiana. Compared with the phenotypes in ∆BbMbp1 mutant strain [10], the BbSwi6 loss has fewer influences. Filamentous fungi develop robust mycelia and conidiation process, which is essential for fungal survival and dispersal in ecosystem [32]. After killing the hosts, B. bassiana first grows into intense mycelia on cadavers and then produces numerous conidia which initiate the following infection cycle [33]. BbSwi6 contributes to mycoses and conidiation on the dead hosts. B. bassiana conidia form and mature on the ‘zig-zag’ conidiogenous cells [34]. BbSwi6 is significantly required for formation of conidiogenous cells (Figure 3A). This result indicated that the loss of conidiation in ∆BbSwi6 mutant strain might be due to the impaired conidiophore. BbSwi6 mutation results in an approximate 99% reduction in conidial production (Figure 3C), and BbMbp1 mutant lacks conidiation. As for blastospore formation under submerged condition, BbSwi6 loss leads to 90% reduction in yield (Figure 3D), and disruption of BbMbp1 causes 95% reduction [10]. As for fungal virulence, LT50 value for BbSwi6 strain is delayed approximately 20%, and the BbMbp1 loss results in a delay of 50%. Like most host-invasion myco-pathogens [35], B. bassiana undergoes dimorphic transition to generate single-cellular hyphal bodies, which is critical for fungal virulence [36]. Therefore, significantly weakened virulence might be caused by the dramatically compromised dimorphism in the host hemolymph. However, ∆BbMbp1 mutant loses the dimorphism potential of 95% [10], and ∆BbSwi6 mutant loses 88% (Figure 4D). These findings suggest that BbMbp1 and BbSwi6 might play different roles during fungal differentiation under aerobic and submerged conditions.

To verify this hypothesis, transcriptomic analyses were used to explore their impacts on global gene expression. Similar to BbMbp1, BbSwi6 has broad influences on transcriptome during fungal development under aerial and submerged conditions; however, these two genes have different regulatory effects. During conidiation under aerobic condition, in terms of functional catalogs, BbSwi6 performed transcriptional control of metabolism, cell transport, cellular rescue and interaction with the environment, whereas BbMbp1 regulates conidial formation via transcriptional control of the cell cycle, metabolism and cell differentiation [10]. The BbSwi6- and BbMbp1-mediated transcriptomes have 39% DRGs overlapped. During conidiation, cell cycle is required for cell division. During fungal growth, cell cycle is required for septum formation and hyphal elongation [37]. BbSwi6 and BbMbp1 regulate the expression of Cdc14 which is a regulator of cyclin dependent kinase and contributes to conidiation [9]. Therefore, these two genes have positive roles in vegetative growth. In
addition, BbMbp1 contributes to the expression of a hydrophobin which localizes on the conidial surface of B. bassiana [38] and significantly contributes to conidiation process [39]. BbSwi6 and BbMbp1 are required for transcription of Ca\textsuperscript{2+}/calmodulin-dependent protein kinase (CaMK) which mediates signaling transduction in a plenty of cellular processes, and plays an important role in conidiation in filamentous fungus (e.g., Magnaporthe oryzae and Arthrobotrys oligospora) [40,41]. These findings suggest that BbSwi6 and BbMbp1 still have common effects on the conidiation-related genes, though they execute divergent transcriptional regulation on different functional categories.

During dimorphism under submerged condition, BbSwi6 also performs comprehensive transcriptional regulation. On the basis of functional categories, BbSwi6 contributes to blastospore generation by regulating the genes involved in metabolism and energy (Figure 5A). The energy metabolism has been linked with dimorphism in filamentous fungi. In Talaromyces marneffei, bioinformatic evidence indicates that energy metabolism is urgent during switching from mycelia form to yeast form [42]. In Mucor circinelloides, Arf-like proteins are involved in maintaining mitochondrial number homeostasis which is required for energy supply [43]. These results suggest that energy metabolism is the evolutionally conserved requirement for dimorphic transition in filamentous fungi. Nevertheless, to explore the mechanisms involved in transcriptional regulation, more experiments are needed. During blastospore formation, BbMbp1 mediates transcriptional control of the genes related to metabolism, cell transport and development [10]. The functional catalog of metabolism is shared by the BbSwi6- and BbMbp1-mediated transcriptomes (Figure 5B). The linkage of metabolism with dimorphism has been revealed in filamentous fungi (e.g., Penicillium marneffei and Ustilago maydis) [44,45]. In addition, these two genes are required for expression of the member of heat shock protein (Hsp) family (e.g., DnaJ protein). DnaJ protein is also known as Hsp40. B. bassiana Mas5 is homologous to DnaJ and critical for fungal blastospore production [46]. BbSwi6 and BbMbp1 contribute to the transcriptional activation of mitogen-activated protein kinase (MAPK) gene Sty1 (Table S9). MAPK cascade plays an essential role in B. bassiana development, including fungal dimorphic transition into blastospores [47]. This implies that BbSwi6 and BbMbp1 have a common regulation on the critical genes and pathways during blastospore formation, although they perform divergent roles on different functional groups of genes.

5. Conclusions

All in all, BbSwi6 interacts with BbMbp1 in nuclei and forms the transcription complex MBF. BbSwi6 plays a vital role in fungal growth, differentiation and virulence in B. bassiana. Although they work as components of MBF, BbSwi6 and BbMbp1 mediate significantly different global expression patterns during fungal differentiation into conidia and blastospores in B. bassiana.

Supplementary Materials: The following are available online at https://www.mdpi.com/article/10.3390/jof7060411/s1, Figure S1: Construction of gene disruption and complementation mutant strains; Table S1: Primers used in this study; Table S2: Differentially expressed genes (DEGs) in comparison of wild type/ΔBbSwi6 during conidiation process; Table S3: Differentially expressed genes (DEGs) in comparison of wild type/ΔBbSwi6 during blastospore formation; Table S4: FunCat analysis of the down-regulated DEGs during conidiation; Table S5: FunCat analysis of the up-regulated DEGs during conidiation; Table S6: FunCat analysis of the down-regulated DEGs during blastospore formation; Table S7: FunCat analysis of the up-regulated DEGs during blastospore formation; Table S8: DEGs in the paired comparison between the BbSwi6- and BbMbp1-mediated transcriptomes during conidiation; Table S9 DEGs in the paired comparison between the BbSwi6- and BbMbp1-mediated transcriptomes during blastospore formation.
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