A Novel Major Facilitator Superfamily Transporter Gene Aokap4 near the Kojic Acid Gene Cluster Is Involved in Growth and Kojic Acid Production in *Aspergillus oryzae*

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Abstract: Kojic acid is an important secondary metabolite of industrial importance produced by *Aspergillus oryzae*. The kojic acid gene cluster plays an essential role in kojic acid production, and harbors *kojA*, *kojR* and *kojT*. The deletion of *kojT*, encoding a major facilitator superfamily (MFS) transporter, did not completely abolish kojic acid production, implying that other transporters are required for the transport of kojic acid. The aim of this study is to look for the transporters involved in kojic acid production. Here, Aokap4 encoding a novel MFS transporter was identified, which was adjacent to *kojT* in the kojic acid gene cluster. The deletion of Aokap4 contributed to the hyphal growth, conidial production and biomass of *A. oryzae*. Moreover, Aokap4 is required for heat- and cell-wall-stress tolerance but not oxidative and osmotic stress. The disruption of Aokap4 impaired kojic acid production with the reduced expression of *kojA*, *kojR* and *kojT*. Furthermore, when *kojT* was deleted in the Aokap4-disrupted strain, the yield of kojic acid declined to the same level as that of the *kojT*-deletion mutant, whereas the production of kojic acid was recovered when *kojT* was overexpressed in the Aokap4 knockout strain, suggesting that *kojT* acts downstream of Aokap4. Aokap4 was the second identified MSF transporter involved in kojic acid production after *kojT* was found a decade ago. This study contributes to a better understanding of the biological roles of the MFS transporter and lays a foundation for future studies on kojic acid synthesis in *A. oryzae*.

Keywords: *Aspergillus oryzae*; kojic acid; MSF transporter; kojic acid gene cluster

1. Introduction

Kojic acid has great application value in such fields as the food, cosmetic, pharmaceutical, and agriculture industries, and is a secondary metabolite produced by *Aspergillus oryzae* [1–3]. Some progress has been made in studying the mechanism underlying kojic acid synthesis [4–10]. The global transcriptional regulator of the secondary metabolite *laeA* negatively mediates kojic acid production through the regulation of the kojic acid synthesis genes *kojA*, *kojR* and *kojT* [4]. As the upstream of *laeA*, loss of function in the histone deacetylase gene *Aohst4* contributes to kojic acid biosynthesis, coupled with high levels of expression of *kojA*, *kojR* and *kojT* [6]. Additionally, some membrane proteins have been reported to be involved in kojic acid production, such as NrtA, AoZip2, AoKap1 and AoKap2 [5,11–13]. In *A. oryzae*, nrtA encodes a nitrate transporter, which abolishes the suppressive effect of nitrate on kojic acid production [5]. RNA interference of the ZIP transporter gene AoZip2 inhibits kojic acid biosynthesis [12]. Aokap1 and Aokap2 both encode an uncharacterized protein with transmembrane domains, and the knockout of Aokap1 or the overexpression of Aokap2 contributes to kojic acid production [11,13]. These proteins affect kojic acid production via altered expression of *kojA*, *kojR* and *kojT* [4,11–13].
The kojic acid gene cluster of *A. oryzae* was identified a decade ago, and consists of three genes: *kojA*, *kojR* and *kojT*. However, little is known about the function of genes near the kojic acid gene cluster except for the reported kojic acid synthesis genes [7]. *KojR* is located in the middle of *kojA* and *kojT* [7]. The Zn(II)2Cys6 transcription factor KojR positively regulates kojic acid synthesis through the kojic acid synthesis enzyme KojA and the major facilitator superfamily (MFS) transporter KojT, and *kojR* is induced by kojic acid [8]. The disruption of *kojA* causes the complete loss of kojic acid production, whereas the deletion of *kojT* only results in the reduced secretion of kojic acid, suggesting that other transporters are required for transporting synthesized kojic acid to the extracellular environment [7]. However, little information on other MSF transporters involved in kojic acid, except for the KojT transporter in *A. oryzae*, is available to date.

The MFS transporters are one of the largest superfamilies of membrane transporters [14–16]. They have a broad spectrum of substrates, including sugars, polyols, metabolites and even inorganic tetraoxyanions [14,15,17]. Most MFS transporters harbor either 12 or 14 putative transmembrane domains (TM) [18]. According to the Transporter Classification Database (https://tcdb.org/ (accessed on 20 August 2022)), the MFS superfamily is divided into 61 families, including the sugar porter family. Many MFS transporters of the sugar porter family have been identified and characterized [18–20], for example, high-affinity glucose transporter HXT2, galactose transporter GAL2, glycerol transporter STL1, etc. [21–25]. However, very limited knowledge on MFS transporters is available for *A. oryzae*.

In this present study, the novel MFS transporter gene *Aokap4* (the kojic acid-related proteins *Aokap1*, *Aokap2* and *Aokap3* have already been studied, and *Aokap4* (AO090113000139) is going to be studied in this work) was identified near the kojic acid gene cluster, which was adjacent to *kojT*. The *Aokap4*-deletion strain was constructed using the CRISPR/Cas9 system. The phenotypes of the wild-type and *Aokap4*-disrupted strains were compared and analyzed. The effects of the deletion of *Aokap4* on kojic acid production were investigated. Moreover, double-deletion mutants of *Aokap4* and *kojT* were constructed to analyze their influence on kojic acid production. Meanwhile, the strain where *Aokap4* was deleted and *kojT* was overexpressed was generated to explore the relationship between *Aokap4* and *kojT*. Our data reveal that the *Aokap4* gene near the kojic acid gene cluster is required for growth, and for the responses to heat and cell-wall stress, and is involved in kojic acid production acting upstream of *kojT*. AoKap4 was the second identified MSF transporter required for kojic acid production after the finding of *kojT* within the kojic acid gene cluster in 2010. This study provides a better understanding of the roles of the MFS transporter and contributes to future studies on the kojic acid gene cluster in *A. oryzae*.

2. Materials and Methods

2.1. Strains and Media

*A. oryzae* 3.042 (CICC 40092) was used as the wild-type strain (WT) and the recipient strain for generating mutants. The yeast strain BY4741 was used for yeast transformation. *A. oryzae* was cultured on Czapek-Dox (CD) medium (2% soluble starch, 0.2% NaNO3, 0.1% KH2PO4, 0.05% MgSO4, 0.05% KCl, 0.05% NaCl, 0.002% FeSO4 and pH 5.5) to characterize the colonial morphologies. The modified CD medium (10% soluble starch, 0.2% NaNO3, 0.1% K2HPO4, 0.05% MgSO4, 0.05% KCl, 0.001% FeSO4, 0.1% yeast extract and pH 5.5) was utilized to produce kojic acid. A YPD (yeast extract/peptone/dextrose) medium (1% yeast extract, 2% peptone, 2% D-glucose) was used for the culture of yeast. The induction medium (0.67% yeast nitrogen base, 2% galactose, 0.078% DO Supplement–Ura (Code No. 630416, Clontech, Dalian, China) was used to express the AoKap4 protein in yeast.

2.2. Sequence Analysis of Aokap4

The conserved motifs of the AoKap4 protein were examined using SMART (http://smart.embl-heidelberg.de/) (accessed on 20 August 2022)) and MOTIF Search (https://www.genome.jp/tools/motif/) (accessed on 20 August 2022)). The transmembrane helices of the AoKap4 protein were predicted using TMHMM Server v.2.0 (https://services.
healthtech.dtu.dk/service.php?TMHMM-2.0/ (accessed on 20 August 2022)). For phylogenetic tree analysis, the homologues of the AoKap4 protein were identified against the EnsemblFungi Database (http://fungi.ensembl.org/index.html (accessed on 20 August 2022)) with BLASTP. AoKap4 and its homologues were used to construct a phylogenetic tree using MEGA7 software. The parameters were as follows: neighbor-joining (NJ) method, 1000 replicates, Poisson model, uniform rates and pairwise deletion.

2.3. Mutagenesis of Aokap4 in A. oryzae

The U6 promoter with the target sequence for Aokap4 was amplified using the plasmid pPTRII-Cas9-AoGld3 [26] as a template with the primers PU6-F and PU6-Aokap4-R, generating the fragment PU6-Aokap4. The U6 terminator and sgRNA sequence were amplified from the vector pPTRII-Cas9-AoGld3 with the primers TU6-Aokap4-F and TU6-R, yielding the fragment Aokap4-TU6. The two fragments above were fused by the primers PU6-F and TU6-R, and inserted into the Smal site of the pPTRII-Cas9 vector [27], yielding the plasmid pPTRII-Cas9-Aokap4. The constructed plasmid was transformed into the A. oryzae 3.042 strain. Transformants were screened on the CD medium containing pyrithiamine, and mutations in Aokap4 were confirmed via nucleotide sequencing.

2.4. Phenotypic Characterization

To investigate the growth of fungal strains, fresh spores were collected from 5-day-old CD plates and adjusted to the concentration of $10^6$ spores/mL with 0.05% Triton X-100 solution. The spore suspensions were inoculated on CD agar medium for three days at 30 °C. The colonies were photographed and their mycelial diameters were measured. The spores were harvested from 6 mm cores using 2 mL of 0.05% Triton X-100 solution, and the numbers of spores were estimated using a hemocytometer. To determine the biomass of the fungal strains, the spore suspensions ($10^6$ spores/mL) were spread on CD agar plates covered by cellophane. After cultivation for three days, the mycelia were collected and dried overnight at 60 °C to weight their biomass. Three replicates were analyzed for each assay.

2.5. Analysis of the Yields of Kojic Acid

The concentration of kojic acid was estimated using the colorimetric method [28]. Firstly, kojic acid with different concentrations (0–0.4 mg/mL) was mixed with the freshly prepared FeCl$_3$-HCl solution (0.06 M FeCl$_3$ and 0.27 M HCl) and the absorbance of the complexation was detected at 500 nm using a Multiskan GO Microplate Reader (Thermo Scientific, Waltham, MA, USA) to manufacture the standard curve. Then, a conidial suspension ($10^6$ spores/mL) of the fungal strains was inoculated in the modified liquid CD media with shaking at 200 rpm and 30 °C for seven days. The supernatant was harvested to react with Fe$^{3+}$ ions (FeCl$_3$-HCl solution) to estimate the yield of kojic acid. The mycelia were collected and dried to quantify their biomass. The content of kojic acid was divided by the biomass. All experiments were repeated three times.

2.6. Gene Expression Analysis Using RT-qPCR

The total RNA was extracted from strains cultured on CD agar or liquid medium at 30 °C for four days using a Fungal Total RNA Extraction Kit (Omega, Norcross, GA, USA) according to the manufacturer’s protocol. The genomic DNA was removed using a gDNA Eraser (TaKaRa, Dalian, China). One μg of RNA was used to synthesize cDNA with a Prime Script™ RT reagent Kit (TaKaRa, Dalian, China). Quantitative real-time PCR was performed using a SYBR Green PCR Kit (TaKaRa, Dalian, China) and a CFX96 Real-Time PCR Detection System (Bio-Rad, Hercules, CA, USA) using the primers listed in Table S1. The histone H1 (AO090012000496) was used as a reference gene. The relative gene-expression levels were calculated using the method described previously [29]. All assays were carried out in triplicate.
2.7. Construction of Double-Deletion Mutant of Aokap4 and kojT

The constructed plasmid pPTRII-Cas9-Aokap4 was transformed into a kojT-deletion strain generated previously by us [27]. The transformants were subcultured on the CD medium containing pyrithiamine and verified via sequencing.

2.8. Overexpression of kojT in the Aokap4 Deletion Strain

The resulting plasmid pPTRII-Cas9-Aokap4 was introduced into the kojT-overexpression strain constructed previously by ourselves using the protoplast transformation method [30]. The candidates were selected using 0.1 µg/mL pyrithiamine and verified via DNA sequencing.

2.9. Heterologous Expression of AoKap4 in Yeast

The coding sequence of Aokap4 was amplified with the primer pYES2-Aokap4-GFP-F/R, and inserted into the BamHI site of the pYES2-GFP vector [31]. The transformation plasmids pYES2-GFP and pYES2-Aokap4-GFP were introduced into the yeast strain BY4741 using the PEG/LiAc-based method. The transformants with the pYES2-GFP vector served as the control. After galactose induction at 28 °C for 12 h using 2% galactose as a carbon source, the yeast cells were collected to observe the fluorescence signal using ZOE™ Fluorescent Cell Imager (BIO-RAD, Hercules, CA, USA).

2.10. Statistical Analysis

All the data were processed using Excel 2020 (Microsoft Corp., Redmond, WA, USA). The statistical significance was determined using GraphPad Prism 8 (GraphPad Software, Inc., San Diego, CA, USA) based on Student’s t-tests. The values are the mean ± SD (n = 3).

3. Results

3.1. Identification and Characterization of Aokap4 Gene

The A. oryzae Aokap4 gene (AO090113000139) is located adjacent to kojT within the kojic acid gene cluster. The Aokap4 gene contains an open reading frame (ORF) of 1578 bp, encoding a 505 amino acid residue. To explore the function of the AoKap4 protein, we firstly performed the functional domain analysis. The domain analysis results revealed that the AoKap4 protein harbored a domain of sugar transporter belonging to the major facilitator superfamily (MFS) (Figure 1A). The analysis of the transmembrane domain showed that AoKap4 contained twelve transmembrane domains across the entire MFS domain and the TC number of AoKap4 is 2.A.1.1, based on STL1 as the homologue of AoKap4 in S. cerevisiae (Figure 1B). To analyze the localization of AoKap4, the AoKap4-GFP fusion protein was expressed heterologously in the yeast strain BY4741. After galactose induction, the transformed yeast cells with the AoKap4-GFP fusion protein were observed at 12 h, and the green fluorescence signals of the AoKap4-GFP fusion protein were detected in the whole cell (Figure S1). To elucidate the phylogenetic relationship between AoKap4 and its orthologs, the phylogenetic tree was constructed. The phylogenetic analysis showed that AoKap4 and its orthologs were classified into three clades (I, II and III) (Figure 1C). AoKap4 and its orthologs of Aspergillus fell into clade I, while the transporter proteins derived from Saccharomyces cerevisiae clustered separately from those of Aspergillus, consisting of clade II (Figure 1C). However, the sugar transporter STL1 in S. cerevisiae was clustered with the members from Aspergillus (Figure 1C).
3.2. Aokap4 Affected the Growth of A. oryzae

To investigate AoKap4 function in A. oryzae, we mutated the Aokap4 gene via the CRISPR/Cas9 system, and successfully obtained two Aokap4-disruption mutants (ΔAokap4-1 and ΔAokap4-2) (Figure 2A). In the ΔAokap4-1 mutant, a 1 bp insertion occurred in the transcript of Aokap4, resulting in the termination of translation (Figure 2A). In the ΔAokap4-2 strain, a 15 bp deletion was found in the coding sequence of Aokap4, which led to the loss of five amino acids in the sugar transporter domain of the AoKap4 protein (Figure 2A). These mutations caused a significant decline in the expression of Aokap4 (Figure 2B). To study the effects of the disruption of Aokap4 on the growth of A. oryzae, the wild-type and Aokap4-disrupted strains were inoculated on the CD agar plates, cultured for three days at 30 °C, and their growth characterization was analyzed and compared. The growth of the Aokap4-deletion mutants increased significantly compared to the wild-type strain (Figure 2C). The Aokap4-deletion mutants formed more conidia than the wild-type strain.
is irrelevant to oxidative and osmotic stress. Aokap4Δ (Figure 2D). Meanwhile, the disruption of Aokap4 resulted in a significant increase in the biomass relative to the control wild-type strain (Figure 2E).

![Figure 2](image_url)

**Figure 2.** Effects of disruption of Aokap4 on the growth of *A. oryzae*. (A) Construction of Aokap4-deletion mutants using the CRISPR/Cas9 system. The targeted sequence for Aokap4 is shown in blue and the corresponding PAM sequence is indicated in red. The number and types of mutations are shown on the right. The Aokap4-deletion mutants ΔAokap4-1 and ΔAokap4-2 harbor a 1 bp insertion and a 15 bp deletion generated by CRISPR/Cas9, respectively. The wild-type (WT), ΔAokap4-1 and ΔAokap4-2 were grown on the Czapek-Dox (CD) agar medium for three days. (B) Transcript levels of Aokap4 in the WT and Aokap4-deletion strains. The mycelia from these fungal strains, cultivated in the modified CD liquid medium for four days at 30 °C, were collected to isolate total RNA. The relative expression levels of the Aokap4 gene were examined via qPCR based on three replicates. (C–E) Comparison of colony diameter (C), conidia formation (D) and biomass (E) in the WT and Aokap4-deletion strains. Asterisks indicate statistical significance in the differences between the WT and Aokap4-deletion strains based on Student’s *t*-tests (** *p* < 0.01). Scale bars = 1 cm.

### 3.3. Disruption of Aokap4 Impaired the Sensitivity to Stress

To further explore the role of Aokap4 in *A. oryzae*, the heat- (37 °C), cell-wall- (Congo red and SDS), oxidative- (menadione and H₂O₂) and osmotic (NaCl)-stress tolerance of the Aokap4 disruptants were analyzed. The wild-type and Aokap4-deletion strains were cultured on the agar medium with 120 µg/mL Congo red, 200 µg/mL SDS, 60 µg/mL menadione, 0.02% H₂O₂ or 1 M NaCl at 37 °C for three days to test their stress (Figure 3A). The Aokap4 disruptants showed significant insensitivity against heat and cell-wall stress, and had a similar growth response to these stressors as the control treatment (Figure 3A,B), suggesting that deletion of Aokap4 decreased the sensitivity to heat and cell-wall stress. However, under oxidative and osmotic stress, no significant differences in growth were observed between the wild-type strain and the ΔAokap4 mutants (Figure 3), indicating Aokap4 is irrelevant to oxidative and osmotic stress.
Figure 3. Phenotypical comparison of the wild-type and Aokap4-deletion strains under stress. (A) Colony growth of the wild-type and Aokap4-deletion strains on the CD agar medium with 120 µg/mL Congo red, 200 µg/mL SDS, 60 µg/mL menadione, 0.02% H2O2 and 1 M NaCl for three days at 30 °C. For heat stress, the plates were incubated at 37 °C. (B) Colony diameter of the wild-type and Aokap4-deletion strains under stress. The colony diameter of the wild-type and Aokap4-deletion strains cultivated on CD plates for three days under stress was measured. Asterisks indicate statistical significance in the differences between the WT and Aokap4-deletion strains based on Student’s t-tests (** p < 0.01; ns, no significant difference). Scale bars = 1 cm.

3.4. Disruption of Aokap4 Repressed Kojic Acid Production in A. oryzae

To investigate the influences of Aokap4 on kojic acid production in A. oryzae, the yield of kojic acid produced by the wild-type and Aokap4-disruption strains was analyzed. Firstly, the wild-type and Aokap4-disruption strains were inoculated on the modified CD agar plates supplemented with 1 mM FeCl3 that were chelated with kojic acid, showing a visible red. After cultivation for three days at 30 °C, the Aokap4-disrupted mutants displayed a less-red color intensity than the control wild-type strain (Figure 4A), suggesting that the Aokap4-deletion strains produced less kojic acid than the wild-type strain. To quantify the amount of kojic acid, the wild-type and Aokap4-disruption strains were cultured in the modified CD liquid medium for seven days at 200 rpm and 30 °C. The amount of kojic acid in the ∆Aokap4 mutants decreased by 92% compared to the wild-type strain (Figure 4B,C). To investigate whether the reduced kojic acid production of the Aokap4-deletion strains is associated with the genes of the kojic acid gene cluster, the expression levels of kojA, kojR and kojT were analyzed. In the ∆Aokap4 mutants, the transcript levels of kojA, kojR and kojT were down-regulated significantly, consistent with the declined production of kojic acid.
Aokap4 affected kojic acid production in A. oryzae. (A) Colony growth of the wild-type and Aokap4-deletion strains cultured on the CD plates with 1 mM FeCl₃ for four days. Kojic acids chelated with FeCl₃ showing a visible red. (B) Color reaction of kojic acid produced by the wild-type and Aokap4-deletion strains inoculated in the modified CD liquid medium for seven days at 200 rpm and 30 °C. (C) The yield of kojic acid in the wild-type and Aokap4-deletion strains cultivated in the modified CD liquid medium for seven days at 30 °C. (D–F) Transcript levels of kojA (D), kojR (E) and kojT (F) in the wild-type and Aokap4-deletion strains. The mycelia from the wild-type and Aokap4-deletion strains cultured in the modified CD liquid medium were harvested to extract total RNA. The gene-expression levels were detected via qPCR. All assays were performed in triplicate. Asterisks indicate statistical significance in the differences between the WT and Aokap4-deletion strains based on Student’s t-tests (** p < 0.01, * p < 0.05).

3.5. Knockout of kojT in the Aokap4-Disrupted Strain Impaired Kojic Acid Production

The data presented above indicate that the involvement of Aokap4 in kojic acid biosynthesis is closely related to the kojic acid gene cluster. In the kojic acid gene cluster, kojT encodes an MSF transporter responsible for delivering kojic acid to the extracellular environment. Aokap4 also belongs to the MSF transporter, which is adjacent to kojT. Given the reduced production of kojic acid in the Aokap4-deletion mutants, we reasoned that the knockout of kojT in the Aokap4-disrupted strain might severely impair kojic acid production. To this end, we constructed the double-deletion mutants of Aokap4 and kojT (ΔAokap4ΔkojT-1 and ΔAokap4ΔkojT-2) based on the previously constructed kojT-deletion mutant using the CRISPR/Cas9 system (Figures 5A and S2). Deletions of 32 bp and 1 bp occurred in the ΔAokap4ΔkojT-1 and ΔAokap4ΔkojT-2 double-deletion mutants, respectively (Figure 5A), which led to the premature translational termination of the AoKap4 protein. In the double-deletion mutants of Aokap4 and kojT, mutations in Aokap4 and kojT resulted in a significant decrease in the transcript levels of Aokap4 and kojT, respectively (Figure 5B,C).
The Quantification of kojic acid production revealed that the yield of kojic acid in the ΔAokap4ΔkojT-1 and ΔAokap4ΔkojT-2 double-deletion mutants had marked reductions and had declined to the same level as the ΔkojT mutant (Figure 5D,E). Additionally, the expression of Aokap4 was down-regulated in the kojT-deletion mutant (Figure 5F).

Figure 5. Disruption of kojT in Aokap4-deletion mutant inhibited kojic acid production. (A) Construction of the double-deletion mutants of kojT and Aokap4 using a CRISPR/Cas9 system. The targeted position is highlighted by the blue box. The Cas9 targeted sequence for Aokap4 is indicated in blue and the PAM sequence is in red. The number and type of mutants in the double-deletion mutants are indicated in blue and the PAM sequence is in red. (B, C) Transcriptional expression of Aokap4 (B) and kojT (C) in the wild-type and double-deletion mutants of kojT and Aokap4. Total RNA was extracted from the mycelia cultivated in the modified CD liquid medium for four days at 30 °C. (D) Kojic acid produced by the wild-type (WT), Aokap4-deletion mutants (ΔAokap4-1 and ΔAokap4-2), kojT-disrupted strain (ΔkojT) and double-deletion mutants of kojT and Aokap4 (ΔAokap4ΔkojT-1 and ΔAokap4ΔkojT-2) cultured in the modified CD liquid medium for seven days at 30 °C. (E) The amount of kojic acid generated by WT, ΔAokap4-1, ΔAokap4-2, ΔkojT, ΔAokap4ΔkojT-1 and ΔAokap4ΔkojT-2. These fungal strains were incubated in the modified CD liquid medium for seven days at 30 °C. (F) The mRNA level of Aokap4 in the wild-type and kojT-deletion strains. ** p < 0.01 represents significant differences between the wild-type and mutants, and ns indicates no significant difference.

3.6. Overexpression of kojT in Aokap4-Deletion Mutant Could Prevent the Reduced Production of Kojic Acid

The above-mentioned data indicate that the deletion of kojT in the Aokap4-disrupted mutant inhibits kojic acid production more severely than the single mutation in the Aokap4 gene. We therefore reasoned that the overexpression of kojT in the Aokap4-disrupted mutant might reverse the reduced production of kojic acid. To this end, we generated the ΔAokap4-OE-kojT strains (ΔAokap4-OE-kojT-1 and ΔAokap4-OE-kojT-2) via the knockout of Aokap4 in the kojT-overexpression strain using the CRISPR/Cas9 system (Figures 6A and S3), in which Aokap4 was deleted and kojT was overexpressed by the A. oryzae amyB promoter (Figure 6C). An 11 bp deletion in the Aokap4 gene was confirmed in the ΔAokap4-OE-kojT-1 strain (Figure 6A). The ΔAokap4-OE-kojT-2 strain harbored a 1 bp deletion in the target sequence of the Aokap4 gene (Figure 6A). Both mutations caused the decreased expression of Aokap4 (Figure 6B). To detect the yield of kojic acid in the ΔAokap4-OE-kojT strains,
the wild-type, Aokap4-deletion mutants, kojT-overexpression strain and ΔAokap4-OE-kojT strains were cultured in the modified CD liquid medium for seven days at 30 °C. The results showed the ΔAokap4-OE-kojT strains recovered kojic acid production but did not return to the level of the kojT-overexpression strain (Figure 6D,E). Meanwhile, the mRNA level of Aokap4 had no change in the kojT-overexpression strain compared with the control wild-type strain (Figure 6F).

Figure 6. Overexpression of kojT in Aokap4-disrupted strain rescued the reduced production of kojic acid. (A) Targeted mutation of Aokap4 in the kojT-overexpression strain using the CRISPR/Cas9 system, producing the ΔAokap4-OE-kojT strain where Aokap4 was deleted and kojT was overexpressed by A. oryzae amyB promoter. The targeted sequence of Aokap4 is indicated in blue and the PAM is shown in red. The mutation types and the numbers of mutated sequences are shown on the right. (B,C) Transcript levels of Aokap4 (B) and kojT (C) in the wild-type and ΔAokap4-OE-kojT strains. (D) Color reaction of kojic acid with ferric ion, which is generated by WT, ΔAokap4-1, ΔAokap4-2, OE-kojT, ΔAokap4-OE-kojT-1 and ΔAokap4-OE-kojT-2. These strains were cultivated in the modified CD liquid medium for seven days. (E) The yield of kojic acid produced by WT, ΔAokap4-1, ΔAokap4-2, OE-kojT, ΔAokap4-OE-kojT-1 and ΔAokap4-OE-kojT-2 cultured in the modified CD liquid medium for seven days. (F) The expression level of Aokap4 in the wild-type and kojT-overexpression strains. Asterisks indicate statistically significant differences between the wild-type and mutants (** p < 0.01). ns, no significant difference.

4. Discussion

It was identified, in 2010, that the kojic acid gene cluster contains three genes: kojA, kojR and kojT [7]. However, to date, there is no information on the function of the genes near the kojic acid gene cluster or whether the genes near the kojic acid gene cluster affect kojic acid production. Here, a novel MSF transporter gene, Aokap4, near the kojic acid gene cluster, was found to involve kojic acid production, which worked in cooperation with KojT. The deletion of Aokap4 contributed to mycelium growth and conidia formation, and resulted in enhanced heat- and cell-wall-stress tolerance in A. oryzae, suggesting that AoKap4 is required for growth and stress tolerance in A. oryzae.
The development of aspergilli primarily includes the growth of the vegetative mycelium and spore formation [32–34]. In this present study, the deletion of Aokap4 contributed to mycelium growth and conidia formation, together with increased biomass (Figure 2), suggesting that AoKap4 acts as a repressor of the growth of A. oryzae. Considering the reduced kojic acid production in the Aokap4-disrupted strain, it seems that AoKap4 is involved in antagonistic action between growth and kojic acid production. The antagonistic action appears to be widespread [9,13,35]. For example, the deletion of kpeA in A. oryzae inhibits conidia formation but increases kojic acid production, while overexpression of Aokap2 suppresses mycelium growth and conidia formation but contributes to kojic acid production [9,13]; the knockout of msnA in A. parasiticus and A. flavus represses colony growth but enhances kojic acid synthesis [35].

As an ortholog of Aokap4, the STL1 gene encodes a sugar transporter-like protein of S. cerevisiae, belonging to the MSF transporter [25]. Although the phylogenetic analysis revealed that AoKap4 is clustered with STL1 of S. cerevisiae (Figure 1C), the role of AoKap4 in the growth of A. oryzae may be different from that of STL1, which is supported by the finding that the disruption of STL1 does not influence the growth of S. cerevisiae [25]. Later research found that STL1 is involved in stress adaptation [24,36–38]. Likewise, the disruption of Aokap4 decreased the sensitivity to heat and cell-wall stress, but not oxidative and osmotic stress (Figure 3), indicating that Aokap4 is required for tolerance to heat and cell-wall stress and unrelated to oxidative and osmotic stress. Furthermore, in general, post-translational regulation is involved in the response to environmental stresses, such as phosphorylation, which is the most common method of post-translational regulation [39,40]. In A. nidulans, the HogA (high-osmolarity glycerol) kinase and histidine kinases participate in the response to stress by phosphorylating transcription factors [39–41]. This means that AoKap4 post-translational regulation may be closely related to HogA-dependent and His-kinase pathways. Collectively, these results suggest that AoKap4 plays multiple roles in growth, and in the heat- and cell-wall-stress responses, as well as kojic acid synthesis.

The kojic acid gene cluster is a key point in producing kojic acid, and includes kojA, kojR and kojT. The transcription factor KojR activates the FAD-dependent oxidoreductase KojA and the MFS transporter KojT to regulate kojic acid biosynthesis [8]. The deletion of kojT did not lead to the complete loss of kojic acid production, implying that there are other transporters that function in kojic acid production [7]. Here, a novel MSF transporter, AoKap4, was found to be involved in kojic acid production in A. oryzae. The disruption of Aokap4 inhibited, but did not abolish, kojic acid production with the decreased expression of kojA, kojR and kojT; this indicates that there is a redundant gene with functions the same as or similar to those of Aokap4 in kojic acid production, and the involvement of Aokap4 in kojic acid production is closely related to the kojic acid gene cluster. Almost all of the kojic acid-related genes are associated with the kojic acid gene cluster. The increased expression of kojA and kojR is responsible for enhanced kojic acid production, caused by the deletion of the transcriptional regular protein gene kpeA [9]. The expression of kojA, kojR and kojT decreases in the deleA disruption strain, which is deprived of kojic acid production [4]. The disruption of the nitrate transporter gene nrtA releases the inhibition of nitrate to kojic acid production and contributes to kojic acid production in the presence of nitrate coupled with the increased expression of kojA, kojR and kojT [5]. Knocking down the expression of the ZIP transporter gene AoZip2 inhibits the production of kojic acid with the reduced expression of kojA, kojR and kojT [12]. However, the correlation between the identified kojic acid-related genes and the kojic acid gene cluster is verified merely via transcriptional analysis, but not genetic analysis. In this study, the deletion of Aokap4 resulted in the decreased expression of kojT, while the disruption of kojT caused the declined expression of Aokap4 (Figures 4F and 5F), implying that there might be an interaction between Aokap4 and kojT to function in kojic acid production in A. oryzae. However, Aokap4 might not keep pace with kojT in enhancing kojic acid production, which is supported by the finding that the expression of Aokap4 has no change in the kojT-overexpression strain (Figure 6F). Furthermore, the amount of kojic acid production in the Aokap4-deletion mutants decreased by 92% compared with
the wild-type strain, while the production of kojic acid in the double-deletion mutant of Aokap4 and kojT was less than 5% of that in the Aokap4-deletion mutants, suggesting that Aokap4 and kojT are essential for kojic acid production in A. oryzae. Moreover, the knockout of kojT in the Aokap4-disrupted strain significantly inhibited kojic acid production of the Aokap4-disrupted strain and led to a decline in kojic acid production in the double-deletion mutants of Aokap4 and kojT to the same level as that of the kojT-deletion mutant (Figure 5D,E); meanwhile, the overexpression of kojT in the Aokap4 disruptant could recover the declined production of kojic acid (Figure 6D,E), indicating that kojT genetically acts downstream of Aokap4.

Interestingly, kojic acid production in the ΔAokap4-OE-kojT strain was reversed by overexpressing kojT, but did not reach the same level as the OE-kojT strain (Figure 6D,E), indicating that the overexpression of kojT can complement the phenotype of the ΔAokap4 mutant with reduced kojic acid production, but Aokap4 did not have the same role in kojic acid production as kojT. In A. oryzae, the MSF transporter KojT is predicted to be located on the cell membrane and the transport kojic acid to the extracellular environment [7]. Given that Aokap4 acts upstream of kojT, the novel MSF transporter AoKap4 may not be located on the cell membrane and may function in delivering an intermediate of kojic acid biosynthesis, but not kojic acid; this is supported by the result wherein the AoKap4-GFP fusion protein was expressed in the yeast. After galactose induction for 12 h, the green fluorescence signal was detected in the entire yeast cell, suggesting that AoKap4 is localized in the cytoplasm, and the localization of AoKap4 to the plasma membrane cannot be excluded (Figure S1). Consistent with this finding is the fact that STL1, the ortholog of AoKap4 in S. cerevisiae, is a dual-targeted protein localized to the plasma membrane and vacuolar lumen [24]. Further analyses for the subcellular localization of AoKap4 will be necessary to analyze the role of A. oryzae in kojic acid production in detail.

In conclusion, a novel MSF transporter gene, Aokap4, located near the kojic acid gene cluster was identified and characterized as being involved in growth and kojic acid production. Aokap4 negatively modulates mycelium growth and conidial formation and is required for heat and cell-wall stress tolerance in A. oryzae. Aokap4 is involved in kojic acid production in A. oryzae, genetically acting upstream of kojT. The study provides a better understanding of the role of the MSF transporter in kojic acid production in A. oryzae, and lays a foundation for further research into the kojic acid gene cluster of A. oryzae.

Supplementary Materials: The following supporting information can be downloaded at: https://www.mdpi.com/article/10.3390/jof8080885/s1, Figure S1: Heterologous expression of AoKap4 in yeast; Figure S2: Construction of the kojT-deletion mutant. Mutations in kojT; Figure S3: Generation of the kojT-overexpression strain; Table S1: Primers used in this study.

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