Molecular cloning, characterization and differential expression of a *Sporothrix schenckii* STE20-like protein kinase SsSte20

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**Abstract.** Dimorphic switching requires fungal cells to undergo changes in polarized growth in response to environmental stimuli. The Ste20-related kinases are involved in signaling through mitogen-activated protein kinase pathways and in morphogenesis through the regulation of cytokinesis and actin-dependent polarized growth. In this study, we isolated and characterized an Ste20 homologue gene, designated SsSte20, from yeast-form *Sporothrix schenckii* (*S. schenckii*). The full length SsSte20 cDNA sequence is 2846 bp in size, and contains an open reading frame of 2505 bp encoding 835 amino acids. The predicted molecular mass of SsSte20 is 91.31 kDa with an estimated theoretical isoelectric point of 5.76. SsSte20 kinase domain shows 63% identity with that of Don3, a germinal centre kinase (GCK) from *Ustilago maydis*. Two exons and one intron are identified within the 2578 bp SsSte20 genomic DNA sequence of *S. schenckii*. Differential expression of the SsSte20 was demonstrated by real-time RT-PCR. The expression of SsSte20 was much higher in the yeast stage compared with that in the mycelial stage, which indicated that the SsSte20 may be involved in the pathogenesis of the yeast phase of *S. schenckii*.

**Introduction**

*Sporothrix schenckii* (*S. schenckii*), the etiologic agent of sporotrichosis, is a dimorphic fungus that produces lymphocutaneous lesions. The mechanisms involved in the morphological transitions of *S. schenckii* remain unknown, although knowledge on the experimental control of these processes has increased in the past years (1,2). Inhibition of the differentiation of *S. schenckii* mycelial into the parasitic yeast forms may be a noteworthy alternative for the design of antifungal drugs to be used in sporotrichosis. Therefore, understanding the morphological transition process and, consequently, how the morphological transition can be inhibited, may aid in the development of novel strategies to effectively manage this increasingly important clinical problem.

Dimorphic switching requires fungal cells to undergo changes in polarized growth in response to environmental stimuli. Polarity establishment allows the development of a wide variety of cell morphologies and the differentiation of distinct cell types (3). The family of Ste20-related kinases is conserved from yeast to mammals and includes the p21-activated kinases (PAKs) and germinal centre kinases (GCKs). These kinases have been shown to be involved in signaling through mitogen activated protein kinase (MAPK) pathways and in morphogenesis through the regulation of cytokinesis and actin-dependent polarized growth (4). *Saccharomyces cerevisiae* (*S. cerevisiae*) Ste20 is required for efficient activity of the pheromone response pathway and an effective Hog1-mediated hypertonic stress response. It appears to play a role in the activation of myosin, adhesion of mating partners and vegetative functions relating to budding (5). Homologues of *S. cerevisiae* Ste20 have been described in a number of other fungal species. These include *Candida albicans* (*C. albicans*) Ste20 (Q92212), *Cryptococcus neoformans* (*C. neoformans*) Ste20 (AF542531.1), *Ustilago maydis* Ste20 (AF299352.1) and *Pneumocystis carinii* (*P. carinii*) Ste20 (AF332388). These Ste20 proteins were proved to play important roles in mediating the regulation of various aspects of morphogenesis, conjugation, mating and pathogenicity (6). We previously reported differentially expressed proteins between the mold and early yeast stage of *S. schenckii* using 2DE (7). The protein spot J is homologous to the Ste20-like protein kinase from *Paracoccidioides brasiliensis* and increases in the early yeast form of *S. schenckii*.

In this study, we described the molecular cloning of the *S. schenckii* Ste20 homologue, designated SsSte20. We performed necessary function analysis of the SsSte20 gene...
and detected the differential gene expression in dimorphic switch of *S. schenckii*. These will establish the primary foundation of understanding the function of the SsSte20 gene from *S. schenckii*. The cloning and characterization of the SsSte20 gene from *S. schenckii* is reported for the first time.

**Materials and methods**

**Fungal strain, media and growth conditions.** The strain of *S. schenckii* used, ATCC10268, was maintained at the Research Center for Pathogenic Fungi, Dalian Medical University, China. To obtain a mycelial culture, the ATCC10268 isolate was inoculated on Sabouraud dextrose agar (SDA) medium and incubated at 25°C. The mycelial colonies thus obtained were inoculated in Sabouraud’s fluid medium and cultured with shaking at 100 rpm at 25°C for 96 h. To achieve the switch of *S. schenckii* from the mycelial phase to the yeast phase, mycelial colonies were transferred to brain heart infusion (BHI) liquid medium at 37°C and shaken at 100 rpm for 96 h. Mycelial and yeast pellets were collected by centrifugation and stored at -80°C.

**Total RNA, genomic DNA isolation and gene cloning.** Approximately 100 mg samples of *S. schenckii* mycelial and yeast were separately pulverized under liquid nitrogen with a mortar and pestle. Total RNA isolation was carried out according to the manufacturer’s protocol using the TRIzol Reagent Kit (Invitrogen, Carlsbad, CA, USA) and treated with the RNase-free DNase I kit from Takara to eliminate DNA contamination. Genomic DNA was isolated from yeast phase colonies following the manufacturer’s protocol using the InstaGene™ Matrix kit (Bio-Rad, Hercules, CA, USA). cDNA was synthesized from 500 μg of total RNA of ATCC10268 by murine leukemia virus reverse transcriptase (MLV-RT) (Takara, Tokyo, Japan) primed with oligo(dT) following the manufacturer’s instructions, and used as template for PCR. Degenerate primers, SsSte20-F1 and SsSte20-R1, were designed based on multiple alignments of the high conserved Ste20 domains of *C. albicans* Ste20 (Q92212), *C. neoformans* GCK-F1, and *Pneumocystis carinii* GCK-R1, and *Ustilago maydis* Ste20 (AF332388) amino acid sequences. PCR product of expected size was cloned into pMD18 vector (Takara) and sequenced. The degenerate primers yielded a 302 bp fragment homologous to known Ste20. To determine the nucleotide sequence of the genomic DNA corresponding to the SsSte20, PCR was performed using the primers SsSte20-P1 and SsSte20-B3 and genomic DNA as template. The PCR products were then sequenced. The sequences of all the primers used in this study are listed in Table I.

<table>
<thead>
<tr>
<th>Primer</th>
<th>Sequence (5’-3’)</th>
</tr>
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<tbody>
<tr>
<td>SsGCK-F1</td>
<td>CAYAARYTGGTGAHTGTATAGGA</td>
</tr>
<tr>
<td>SsGCK-R1</td>
<td>TGYTGATDNAYCTCNGNGGCCATCCA</td>
</tr>
<tr>
<td>CTE869F</td>
<td>AGAAGCTCCACCGGATATCGGACC</td>
</tr>
<tr>
<td>CTE869R</td>
<td>CGCCCACATGACCAAGAAAACCT</td>
</tr>
<tr>
<td>R132-1(GSP1)</td>
<td>TGGAGGCCTAAACAGACA</td>
</tr>
<tr>
<td>R132-2(GSP2)</td>
<td>TTCTTGTGGCTGAGGAGGTAG</td>
</tr>
<tr>
<td>R132-3(GSP3)</td>
<td>CAATGATGGCAATGATGTCCTCG</td>
</tr>
<tr>
<td>P1</td>
<td>ATGGGGGACGAGGATAGCC</td>
</tr>
<tr>
<td>B3</td>
<td>CACGGCTCTTGTGTGAG</td>
</tr>
<tr>
<td>F</td>
<td>GCAACACGACAACAATTAGC</td>
</tr>
<tr>
<td>R</td>
<td>GACACGTTCCTTCTCCATG</td>
</tr>
<tr>
<td>24T</td>
<td>(FAM) CCAACACGCAACTACACACGCAACAG</td>
</tr>
</tbody>
</table>

Degenerate primers designed based on multiple alignments of the high conserved STE20 domains for gene cloning: GCK-F1 and GCK-R1. Primers of 3’-RACE: CTE869 F3, CTE869 F4; 5’-RACE: R132-1, R132-2 and R132-3. To determine the nucleotide sequence of the genomic DNA corresponding to the SsGCK PCR was performed using the primers P1 and B3. Primers and a TaqMan probe of real-time RT-PCR: 8F, 58R and 24T.

**Table I. Sequence of primers in this study.**

**Bioinformatic and phylogenetic analysis of SsSte20.** Nucleotide sequences and deduced amino acid sequences of the cloned SsSte20 gene were analyzed. The nucleotide sequences were analyzed as previously described by us (8) using Sequence software (Seqencer, USA) and the BLAST network service of the National Center for Biotechnology Information (NCBI) (http://www.ncbi.nlm.nih.gov/blast). The open reading frame (ORF) was found by the ORF Finder (http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?db=nr&id=). For the exact localization of the exon/intron boundaries, the mRNA-to-genomic alignment program Spider (http://www.ncbi.nlm.nih.gov/IEB/Research/Ostell/Spidey/index.html) was used. The deduced amino acid sequence was analyzed with the Expert Protein Analysis System (http://www.expasy.org/) and the protein domain features of SsSte20 were determined by using Simple Modular Architecture Research Tool (http://hits.isb-sib.ch/cgi-bin/PFSCAN). Isoelectric point and molecular weight prediction were carried out at (http://cn.expasy.org/tools/pi_tool.html). Multiple alignments of SsSte20 were performed with the ClustalW Multiple Alignment Program (http://www.ebi.ac.uk/clus talw/).

**Differential expression of SsSte20 in two stages during dimorphic switch.** The expression of SsSte20 transcript in different stages (mycelial, yeast) was measured by real-time RT-PCR. Primers and a TaqMan probe for target genes were designed with Primer Select in DNASTAR software (Lasergene) and are listed in Table I (24T, 8F and 58R). Fifty nanograms of total RNA were assayed from two stages of *S. schenckii* in triplicate using the PrimeScript RT-PCR Kit (Takara). The minus-reverse transcriptase control was also performed in triplicate. The amplification conditions were optimized for the ABI-PRISM-7500 instrument (Applied Biosystems). The cycling conditions using TaqMan probe detection were 95°C for 2 min followed by 40 cycles at 95°C for 10 sec, 61°C for 10 sec, 72°C for 40 sec. 18srDNA was selected as the endog-
enous control. Relative quantification of target gene expression was evaluated using the comparative cycle threshold (C\textsubscript{T}) method as previously described by Livak and Schmittgen (9). The ΔC\textsubscript{T} value was determined by subtracting the target C\textsubscript{T} of each sample from its respective 18s rDNA C\textsubscript{T} value. Calculation of ΔΔC\textsubscript{T} involved using the mycelial sample ΔC\textsubscript{T} value as an arbitrary constant to subtract from yeast sample ΔC\textsubscript{T} values. Differences in expression of target genes were determined by 2^{-ΔΔC\textsubscript{T}}. Data are expressed as arithmetic means ± SD unless otherwise indicated. Comparison between mycelial and yeast samples was performed using the Student's t-test. Differences with a P-value of <0.05 were considered statistically significant.

### Results

**Cloning and genomic structure of SsSte20.** A full-length SsSte20 cDNA (2,846 bp) including an ORF of 2505 bp, encoding 835 amino residues, was flanked by a 41 bp 5'-untranslated region (5'-UTR) and a 297 bp 3'-UTR. The SsSte20 genomic DNA is 2578 bp in length. The aligned results revealed that there is one intron between the sequences of the genomic DNA and the cDNA. Its 5' and 3' ends conformed to the basic consensus, GT/AG, for the eukaryotic splice donor and acceptor site. Based on the sequence of cDNA, the molecular weight of the predicted amino acid is ~91.31 kDa, the theoretical pl is 5.76.

Motif searches and sequence comparison showed that SsSte20 consists of a 251-amino acid N-terminal kinase domain (residues 21-271) (Fig. 1). In the N-terminal extremity of the catalytic domain there is a glycine-rich stretch of residues in the vicinity of a lysine residue (K), which has been proved to be involved in ATP binding. In the central part of the catalytic domain there is a conserved aspartic acid residue (N) which is important for the catalytic activity of the enzyme (Fig. 2). One proline/glutamic acid/serine/threonine (PEST) motif was involved using themycelial C\textsubscript{T} as a calibrator to set the baseline for comparing mean differences in the ΔC\textsubscript{T} values of yeast form. 2^{-ΔΔC\textsubscript{T}}, normalized target amount relative to the mycelial form.

### Expression of SsSte20 in two stages of S. schenckii.** The mRNA expression of SsSte20 in different stages was analyzed by real-time RT-PCR normalized against 18s rDNA levels. Expression was determined as fold increased 2^{-ΔΔC\textsubscript{T}} levels relative to the stage with lowest expression (mycelial) set to 1. The SsSte20 gene was expressed in two stages of S. schenckii, with higher mRNA levels observed in yeast (15.14-fold). There were significant differences between the mycelial and the yeast form (Table II).

### Discussion

The Ste20-like protein kinase family can be divided into two subfamilies based on their domain structure and regulation. The PAK/Ste20 subfamily contains a C-terminal catalytic domain and an N-terminal binding site for the small G proteins Cdc42 and Rac1, which are considered to be key regulatory molecules linking surface receptors to the organization of the actin cytoskeleton (11,12). By contrast, the kinases of the second subfamily, the GCK subfamily, have an N-terminal kinase domain and a C-terminal regulatory domain that does not have a recognizable GTPase binding site (13). Furthermore, GCKs are subdivided into two broad groups based on their structure. Group I GCKs have a highly conserved C-terminal regulatory domain, which does not contain any identifiable sequence motifs.

### Accession number

The full length of cDNA sequence and the genomic DNA sequence of the SsSte20 gene were submitted to the GenBank database under the accession numbers JX312329 and JX857536, respectively.
amino acids. Group II GCKs share similarity to the catalytic domain of group I; however, the C-terminal domain differs significantly. In present study, SsSte20 was labelled as a GCK based on catalytic domain located in the N-terminal site, which was also confirmed by phylogenetic relationship analysis. We further identified SsSte20 as a group II GCK for its one PEST motif, no SH3 domain and no conserved sequence in C-terminal domain.
Ste20 ortholog was proved to be involved in both the morphogenesis and the pathogenesis of consequence of an inability to grow in the pathogenic form. In *C. albicans*, filamentous growth allows adherence and penetration of tissues during infection. The cst20 (Ste20) mutants are unable to form hyphal filaments *in vitro* and consequently show reduced kidney colonization and virulence in a mouse model for systemic candidiasis (14). While in *Penicillium marneffei*
(P. marneffei), infection occurs by the inhalation of conidia that are phagocytosed by host pulmonary alveolar macrophages where they germinate into yeast cells and divide by fission. PakA (Ste20) plays an essential role in conidial germination of P. marneffei at the host temperature of 37˚C such that ΔPakA mutant conidia fail to germinate during macrophage infection (15). Although GCKs currently have no described role in virulence, this may be due to a lack of characterization rather than a lack of involvement. The Kic1 (GCK) mutants in C. albicans and C. neoformans probably have virulence defects as the C. albicans Kic1 mutant is unable to grow filamentously and the C. neoformans mutant is unable to grow at 37˚C, the temperature of the human body, both of which are requirements for virulence in these organisms (16,17). In S. schenckii, the germination of conidia is a key pathogenicity determinant since conidia are the infectious propagules. Does SsSte20, a GCK, have the same function in the virulence of S. schenckii as Ste20 ortholog in other fungi? The mRNA expression of SsSte20 in yeast cells of both ATCC10268 and a clinical S. schenckii isolated from a patient with fixed sporotrichosis (data not shown) were higher than the mycelial ones, which suggested that SsSte20 may be involved in the morphogenesis or pathogenesis of S. schenckii.

What is the environmental signal that SsSte20 responds to to participate in morphogenesis and pathogenesis? In P. marneffei, PakB (Ste20) is essential for the formation of yeast cells within macrophages and the ΔPakB mutant grows as septate hyphae in vivo. PakB is not expressed at 37˚C during in vitro yeast growth but expression is induced during macrophage infection (18). This suggests that unlike PakA, which responds to temperature, PakB responds to host cell inductive signals. In the present study, the mycelial cells of S. schenckii switched to yeast cells with obvious increased expression of SsSte20 when they were incubated in BHI liquid medium at 37˚C in vitro, which suggests SsSte20, similar to PakA, can detect the change of temperature and nutrient deprivation in the environment. Whether SsSte20 also responds to other host signals remains to be elucidated. It is most likely that there are other proteins that fungal cells come into contact with in the early stages of infection. We are carrying out further experiments on generating SsSte20-mutants to study its detailed functions in this critical fungal pathogen.

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References


