2-1-2000

Biochemical and Genetic Conservation of Fission Yeast DSK1 and Human SRPK1

Zhaohua Irene Tang
Claremont McKenna College; Pitzer College; Scripps College

Tiffany Kuo
Jenny Shen
Ren-Jang Lin
Beckman Research Institute

Recommended Citation

This Article is brought to you for free and open access by the W.M. Keck Science Department at Scholarship @ Claremont. It has been accepted for inclusion in WM Keck Science Faculty Papers by an authorized administrator of Scholarship @ Claremont. For more information, please contact scholarship@cuc.claremont.edu.
Biochemical and Genetic Conservation of Fission Yeast Dsk1 and Human SR Protein-Specific Kinase 1

ZHAOHUA TANG, TIFFANY KUO, JENNY SHEN, AND REN-JANG LIN*

Department of Molecular Biology, Beckman Research Institute of the City of Hope, Duarte, California 91010

Received 6 August 1999/Returned for modification 28 September 1999/Accepted 25 October 1999

Arginine-serine-rich (RS) domain-containing proteins and their phosphorylation by specific protein kinases constitute control circuits to regulate pre-mRNA splicing and coordinate splicing with transcription in mammalian cells. We present here the finding that similar SR networks exist in Schizosaccharomyces pombe. We previously showed that Dsk1 protein, originally described as a mitotic regulator, displays high activity in phosphorylating S. pombe Prp2 protein (spU2AF59), a homologue of human U2AF65. We now demonstrate that Dsk1 also phosphorylates two recently identified fission yeast proteins with RS repeats, Srp1 and Srp2, in vitro. The phosphorylated proteins bear the same phosphoepitope found in mammalian SR proteins. Consistent with its substrate specificity, Dsk1 forms kinase-competent complexes with those proteins. Furthermore, dsk1Δ gene determines the phenotype of prp2Δ overexpression, providing in vivo evidence that Prp2 is a target for Dsk1. The dsk1Δ-null mutant strain became severely sick with the additional deletion of a related kinase gene. Significantly, human SR protein-specific kinase 1 (SRPK1) complements the growth defect of the double-deletion mutant. In conjunction with the resemblance of dsk1Δ and SRPK1 in sequence homology, biochemical properties, and overexpression phenotypes, the complementation result indicates that SRPK1 is a functional homologue of Dsk1. Collectively, our studies illustrate the conserved SR networks in S. pombe consisting of RS domain-containing proteins and SR protein-specific kinases and thus establish the importance of the networks in eucaryotic organisms.

Arginine-serine-rich (RS) domain-containing proteins are among the best-characterized non-snRNP proteins participating in pre-mRNA splicing (for reviews, see references 8 and 19). Members of the protein superfamily are involved in constitutive splicing and are specific modulators of alternative splicing (15, 19). Mammalian serine/arginine-rich (SR) proteins are featured by one or more RNA recognition motifs at the NH2 terminus and by an RS domain at the COOH terminus. Other RS domain-containing proteins are relatively less defined with respect to the arrangement of the two structural elements in a protein (8, 11, 19, 35).

SR proteins are heavily phosphorylated, predominantly in the RS domain (4, 5, 12, 41). Several kinases have been reported to phosphorylate RS domain-containing splicing factors (5, 12, 30, 39, 50, 53), including SR protein-specific kinase (SRPK) and Cdc28/Cdc2-like kinase (Clk/Sty). Based on studies in mammalian nuclear extracts, both phosphorylation and dephosphorylation of SR proteins are required for pre-mRNA splicing. Phosphorylation of SR proteins may promote spliceosome assembly by facilitating specific protein interactions while preventing SR proteins from binding randomly to RNA (54). Once a functional spliceosome has formed, dephosphorylation of SR proteins is necessary to allow the transesterification reaction to occur (3, 23). Recently, human type 2C Ser/Thr phosphatase PP2Cγ was reported to be required during early stages of spliceosome assembly and to be physically associated with the spliceosome in vitro (29). Therefore, the sequential phosphorylation and dephosphorylation of SR proteins may mark the transition between stages in one round of splicing reaction.

The phosphorylation state of SR proteins not only regulates their functional properties in splicing reaction but also modulates their subnuclear distribution in vivo (5, 12, 26, 50). The phosphorylation of the serine residues in the RS domain is a prerequisite for the release of splicing factors from the storage loci, nuclear speckles, to the sites of transcription and splicing, suggesting that protein phosphorylation functions as a control switch for spatially linking transcription with splicing in vivo (24). In a simplified scenario, the ability of the splicing machinery to respond to mRNA synthesis in the cell may be conferred by the differential phosphorylation of SR proteins, so that sufficient splicing factors can be recruited to the sites of transcription as gene expression is activated.

In addition to transcription, pre-mRNA splicing is closely coordinated in space and time with other nuclear events, including 5′ capping, and the 3′ processing of RNA (25). Gene expression is also synchronized with the cell division cycle, such that it is active during interphase and repressed upon entry into mitosis (9). Therefore, intrinsic interplay exists among pre-mRNA splicing, transcription, and cell cycle. RS domain-containing proteins and SR protein-specific kinases may constitute a protein relay or networks to regulate the coupling of splicing, transcription, and cell cycle in mammalian cells (6, 25). The fission yeast Schizosaccharomyces pombe, as a genetically tractable system, has been widely used to investigate cell cycle control (14, 31). S. pombe also bears resemblance to mammalian systems with respect to the high content and structure of introns in protein-encoding genes (13, 36, 48). An increasing body of evidence suggests the interplay between pre-mRNA splicing and cell cycle in fission yeast. A splicing defect is coupled with acdc phenotype at a restrictive temperature in 10 of 14 prp ts mutants identified in fission yeast, i.e., prp1, prp2, prp5 through prp8, and prp11 through prp14 (33, 45, 48, 49). Defects in nuclear division, cytokinesis, and particularly G2/M transition were observed in those 10 prp mutants. These cell cycle defects are not simply a result of malfunction in splicing since not all prp mutants impose a block on mitotic
Protein components similar to elements of the mammalian SR networks exist in S. pombe. First, several RS domain-containing proteins have been identified. The Prp2 protein, also named spU2AF59 due to its homology to the large subunit of human U2AF (35), is essential for pre-mRNA splicing in vivo (34). Another mutant allele, spSRP14-53, affects chromosome segregation and leads to minichromosome loss (45). In addition to Prp2/Mis11 protein, Srp1 and Srp2 are two proteins containing RS repeats recently found in S. pombe (11). The srp2\(^+\) gene is essential for viability, while the srp1\(^+\) gene is not. Overexpression of Srp1 protein with a mutant RS domain or the RNA-binding domain alone inhibits splicing in fission yeast, suggesting a role for Srp1 in pre-mRNA splicing (11). Second, kinases that phosphorylate RS domain-containing proteins have been discovered. Dsk1 is an S. pombe protein kinase that specifically phosphorylates Prp2 in vitro (47). Although initially described as a mitotic regulator (46), Dsk1 has also been implicated in pre-mRNA splicing according to its sequence homology to human SRPK1 (12). Another protein kinase, Prp3 (38), is reported to phosphorylate human SF2/ASF protein in vitro (10).

In further investigating the kinase activity of Dsk1 and its interaction with RS domain-containing proteins, we show here for the first time that phosphorylation of S. pombe RS domain-containing proteins by Dsk1 produces the same phosphoepitope found in mammalian SR proteins. We also obtained in vivo evidence to support the kinase-substrate relationship between Dsk1 and Prp2. The dsk1\(^{-}\)-null mutant became severely sick with additional deletion of a related kinase. Significantly, human SRPK1 protein expressed in fission yeast is capable of compensating for the loss of Dsk1 in vivo. Consistent with the notion that SRPK1 is a functional homologue of Dsk1, the kinase-substrate relationship be-

**Materials and Methods**

*S. pombe strains.* The following haploid strains of *S. pombe* were used: 1913 (h\(^-\) leu1), B6 (h\(^-\) leu1 ura4 dsk1::ura4\(^+\)) (46), 2A5 (h\(^-\) leu1 ura4 kic1::ura4\(^+\) his2), 2D4 (h\(^-\) leu1 ura4 kic1::ura4\(^+\) dsk1::ura4\(^+\) his2). Standard genetic procedures and media for growing *S. pombe* strains are described elsewhere (1, 27).

**Plasmid construction.** Fusion yeast srp1\(^+\) gene was obtained by PCR (42) from the *S. pombe* cDNA library (Clontech) by using two primers complementing the 5' and 3' sequence of the gene, respectively: 5'-GGCCGGCAGATCATGATGCGAAGGCTCCAG-3', including a BamHI site, and 5'-GGCCGATGCTGACATACATGGTTACGGG-3', including a SalI site. The BamHI-SalI fragment of ~900 bp was then inserted into pET-28a (Novagen) to generate pET-28a srp1\(^+\). To construct pET-28GST-srp1\(^+\), a BamHI-SalI fragment was produced by PCR by using pGADGH as a template with two primers: 5'-GGCCGGCAGATCATGATGCGAAGGCTCCAG-3', including a BamHI site, and 5'-GGCCGATGCTGACATACATGGTTACGGG-3', including a SalI site. Plasmids pET-28GST, pET-28SAD4k, and pET-28GST-prp2\(^+\) have been described (47). Plasmid pGADGH srp2\(^+\) DNA was isolated from the *S. pombe* cDNA library by using srp2\(^+\) gene as bait (unpublished data). An EcoRI fragment containing the coding sequence of the srp2\(^+\) gene was ligated to vector pET-28b and pET-28GST to produce pET-28bsrp2\(^+\) and pET-28GSTsrp2\(^+\). To generate pREPrp2\(^+\), a 1.4-kb NdeI-BamHI DNA fragment encoding the Prp2 protein was isolated by pREPrp2 (20). The pREPrp2 vector plus plasmid was constructed by inserting a SalI-BamHI fragment containing the open reading frame of human SRPK1 into the same sites of pREPl. The SRPK1-BamHI fragment was synthesized by PCR by using pGADH-SRPK1 (from X. D. Fu University of California at San Diego) as template and two oligonucleotides (5'-AGGTGCGGTTCTCCTCGACATTTACAAAGACCAGAC T3' and 5'-TTGTGGGATCCCTGCCTGACTCAGGTTGTGTCG-3') as primers.

**Production of recombinant proteins.** Recombinant proteins GST-Srp1, Srp1, GST-Srp2, Srp2, GST-SF2, SF2/ASF, and Dsk1 were expressed in *Escherichia coli* BL21(DE3)PlyS as described earlier (47). Iodopropyl-\(\beta\)-d-thiogalac-
topyranoside (IPTG) was added to a final concentration of 1 mM, instead of 0.4 mM, to assure a full induction of a T7lac promoter (Novagen) in bacteria. Bacterial lysate preparations and histidine-tagged Dsk1 protein purification have been described (47). The relative amounts of recombinant proteins in lysates were estimated based on Coomassie blue-stained gels by using a standard protein albumin as a standard, or the intensity of protein bands was visualized on immunoblots.

**GST pulldown assay.** Bacterial lysates containing glutathione S-transferase (GST) or GST fusion proteins were incubated with or without various non-GST-tagged proteins in lysates to allow complex formation at 23°C for 30 min. The mixture was then incubated with glutathione beads at 4°C for 1 h with constant agitation. After pulldown by microcentrifugation at 7,000 rpm for 1 min at room temperature, the beads were washed in TBS (10 mM Tris-HCl, pH 7.4; 150 mM NaCl) two to three times with 0.1% NP-40 and four times without NP-40. The beads were resuspended in TBS as a 10% suspension, aliquoted, frozen in liquid nitrogen, and stored at −80°C until use. All steps were performed in the presence of protease inhibitors: 5 μg of pepstatin, 5 μg of chymostatin, and 5 μg of leupeptin per ml plus 1 μM phenylmethylsulfonyl fluoride.

**Kinase assay.** Purified or bead-bound Dsk1 was incubated at 23°C for 30 min with in vitro translated bacteriophage T7-tagged or bound to glutathione beads in a total volume of 20 to 60 μl in a kinase buffer (50 mM Tris-HCl, pH 7.4; 10 mM MgCl\(_2\); 1 mM dithiothreitol) with 50 μM ATP and 0.1 μM of [γ-\(^{32}\)P]ATP per μl. When a bead-bound protein was present in a kinase reaction, the beads were added to the sample during incubation. The kinase reaction was terminated by boiling in sodium dodecyl sulfate (SDS) sample buffer, and the samples were resolved on a SDS–10% polyacrylamide gel. Protein phosphorylation was detected by autoradiography. For Western blot analysis, the kinase reaction was performed by employing an ATP-regenerating system (10 mM creatine phosphate, 1 mM ATP, and 0.1 μg of creatine phosphokinase per ml) without radioisotopes. Immunoblotting in most experiments was performed as previously described (47). When 3C monocalonal antibody was used, 25 mM NaF and 1 mM NaVO\(_3\) were present as phosphatase inhibitors to prevent dephosphorylation.

**Antibodies.** The anti-Dsk1 polyclonal antibodies were generated and affinity purified as described earlier (47). Monoclonal antibody (MAb) 3C5 was obtained from mouse ascites and was used in a 5000 dilution. Anti-GST polyclonal antibodies were from Santa Cruz Biotechnology. Anti-T7-Tag monoclonal antibody was purchased from Novagen.

**Transformation of *S. pombe*.** Transformation of fission yeast was accomplished by using the lithium acetate method (1) with modifications. A 3- to 5-mI culture was placed in YES medium (27) and grown at 33°C for about 5 h with shaking at 225 rpm. A 100- to 200-mI culture was then started by adding a calculated amount of cells from the small culture so that cell density would reach 0.5 \(\times\) 10\(^7\) to 1.5 \(\times\) 10\(^7\) cells/ml overnight. Cells were harvested and resuspended at a density of approximately 10\(^7\) cells/ml in 0.1 M lithium acetate in TE buffer. After 1 h of incubation at 30°C with shaking at 170 to 200 rpm, 1 μg of plasmid DNA in 15 μl of TE was mixed with 100 μl of the cell resuspension, followed by the addition of 290 μl of 50% glycerol. Samples were incubated for 1 h with occasional vortexing. After heat shock at 42°C for 15 min, cells were incubated at room temperature for 10 min. Cells were then collected, washed, and resuspended in 200 μl of EMME2 (minimal medium) (1). Finally those transformed cells were spread on EMM plates in the presence of 2 μM thiamine and incubated at 33°C until colonies appeared.

**Expression of fission yeast prp2\(^+\) gene and human SRPK1 gene in *S. pombe*.** The plasmid pREPrp2\(^+\) was transformed into fission yeast wild-type (strain 1913), dsk1-null (Δdsk1), and kic1-null (Δkic1) strains. The human SRPK1 gene was introduced as plasmid pREPlSRPK1 into wild-type (strain 1913) and dsk1 kic1 double-null (Δdsk1Δkic1) strains of *S. pombe*. The expression of prp2\(^+\) gene and human SRPK1 gene under the control of nmt\(^{-}\) promoter was induced according to procedures described elsewhere (46).

**DAPI staining.** Methods for DAPI (4',6'-diamidino-2-phenylindole) staining were modified from Alfa et al. (1) and the Fission Yeast Handbook (www.bio.uw.edu/pombe/handbook). Cells were fixed on a slide at 70°C for 1 min on a hot plate. Then, 3 to 4 μl of a freshly diluted 1:1 DAPI solution (1 μg of DAPI per ml, 1 mg of antifade per ml, 45% glycerol) was added to the fixed cells. Slides were kept in the dark to prevent fading before they were observed under a microscope.

**Results**

Dsk1-mediated phosphorylation of fission yeast Srp1, Srp2, and Prp2 proteins generates the same phosphoepitope as in mammalian SR proteins. We showed previously that Dsk1 protein specifically phosphorlates fission yeast Prp2/Mis11, a U2A65 homologue, in vitro (47). To extend our studies of the SR networks in *S. pombe* we investigated whether Srp1 and Srp2 proteins are also substrates for Dsk1 in vitro. Full-length...
Srp1 and Srp2 proteins fused at the NH2-terminus to GST, designated GST-Srp1 and GST-Srp2, were isolated on glutathione-agarose beads and incubated with or without purified Dsk1 in the presence of [γ-32P]ATP. As shown in Fig. 1, 32P-labeled proteins with apparent molecular sizes of ~56 kDa (lane 4) and ~66 kDa (lane 6) were detected, matching the predicted sizes of GST-Srp1 and GST-Srp2 proteins, respectively. These bands were not detected in the samples without Dsk1 protein (Fig. 1, lanes 3 and 5). The lower-molecular-size band observed in lane 4 of Fig. 1 was probably a degradation product of GST-Srp1. The GST portion of the fusion proteins did not contribute to the phosphorylation by Dsk1, since GST alone was not phosphorylated by Dsk1 (Fig. 1, lane 2). Therefore, in addition to Prp2, Dsk1 phosphorylates Srp1 and Srp2 proteins in vitro.

To assess the specificity of phosphorylation, we probed the Dsk1-phosphorylated proteins with SR protein-specific MAbs. Mammalian SR proteins share common phosphoepitopes, which specifically react to two MAbs, MAb 104 (40) and MAb 3C5 (2). Since MAB 3C5 is more sensitive and specific for this epitope, regardless of whether the substrate was fused with GST or not. Purified GST-Prp2 and GST-SF2/ASF were analyzed similarly (Fig. 2, top panel, lanes 10 to 13). Both proteins were detected by anti-GST polyclonal antibodies (Fig. 2, top panel, lanes 10 to 13), and after phosphorylation by Dsk1, they were recognized by MAB 3C5 (bottom panel, lanes 11 and 13). In these samples Dsk1 protein was monitored by anti-Dsk1 polyclonal antibodies (top panel, lanes 11 and 13). As shown in this experiment, all three RS domain-containing proteins, Srp1, Srp2, and Prp2, were recognized by MAB 3C5 after Dsk1 action, reflecting a general feature of Dsk1-mediated phosphorylation. The weaker signal of Prp2 is likely due to the lower amount of Prp2 protein present in the reaction mixture as well as the presence of fewer RS repeats in Prp2 than in the other proteins. The results provide the first biochemical evidence that fission yeast RS domain-containing proteins phosphorylated by Dsk1 share the same phosphoepitope with the mammalian SR proteins. Therefore, Dsk1 behaves similarly to its mammalian counterparts at the molecular level. The conserved phosphorylation of RS domain-containing proteins from distinctive organisms implicates its importance in eucaryotic systems.

**Dsk1 forms kinase-competent complexes with RS domain-containing proteins.** As kinase-substrate pairs, physical association of Dsk1 with the RS domain-containing proteins must take place during the phosphorylation process. If these interactions take place during the phosphorylation process.
actions are stable, Dsk1 protein should coisolate with GST fusion substrates in a GST pulldown assay. Lysates containing similar amounts of GST fusion proteins were incubated with a lysate containing Dsk1 protein. Glutathione-agarose beads were then added to bind the GST fusions. Portions of the mixed lysates, unbound fractions, and bound fractions from each sample were analyzed by SDS-polyacrylamide gel electrophoresis. Some samples were processed for immunoblotting by using anti-T7-Tag MAb (lanes 1 to 9), which detects GST, GST-Srp1, GST-Srp2, and Dsk1. Other samples were processed for immunoblotting first with anti-GST and subsequently with anti-Dsk1 polyclonal antibodies (lanes 10 to 12). Dsk1 protein was pulled down by each of the four RS domain-containing proteins (lanes 6, 9, and 12) but not by GST protein (lane 3). Numbers 1 to 5 on the left of the protein bands in the bound fraction of each sample represent Dsk1, GST-Srp1, GST-Srp2, and GST-Prp2, respectively, as indicated on the right side of the figure.

FIG. 3. Srp1, Srp2, and Prp2 proteins individually form a complex with Dsk1 in vitro. A bacterial lysate containing Dsk1 protein was incubated with a lysate containing GST (lanes 1 to 3) or GST fusion (lanes 4 to 12) proteins as indicated at the top of each lane to allow complex formation at 23°C for 30 min. Glutathione beads were then added to pulldown bound proteins at 4°C as described in Materials and Methods. Portions of mixed lysates, unbound fractions, and bound fractions from each sample were analyzed by SDS-polyacrylamide gel electrophoresis. Some samples were processed for immunoblotting by using anti-T7-Tag MAb (lanes 1 to 9), which detects GST, GST-Srp1, GST-Srp2, and Dsk1. Other samples were processed for immunoblotting first with anti-GST and subsequently with anti-Dsk1 polyclonal antibodies (lanes 10 to 12). Dsk1 protein was pulled down by each of the four RS domain-containing proteins (lanes 6, 9, and 12) but not by GST protein (lane 3). Numbers 1 to 5 on the left of the protein bands in the bound fraction of each sample represent Dsk1, GST-Srp1, GST-Srp2, GST-Prp2, and GST, respectively, as indicated on the right side of the figure.

Dsk1 complexes isolated by the GST pulldown procedure were regenerating system for 30 min at 23°C. Following the kinase reaction, samples were centrifuged to pellet the beads, and proteins released from the complex in the presence of ATP (lanes 4 and 8), but it is not dissociated from the complex in the absence of ATP (lanes 1, 2, 5, and 6) an ATP regenerating system for 30 min at 23°C. Following the kinase reaction, protein-bound beads were pelleted by centrifugation. The supernatant (S) and bead (P) portions of each sample were resolved on an SDS–10% polyacrylamide gel and subsequently processed for immunoblotting with anti-T7-Tag MAb. The bound Dsk1 phosphorylated Srp1 and Srp2 in the complex (lanes 1 and 3). (A) The bound Dsk1 phosphorylates Srp1 and Srp2 in the complex in the presence of ATP. The pulldown complexes GST-Srp1/Dsk1 (lanes 1 and 2) and GST-Srp2/Dsk1 (lanes 3 and 4), as described in Fig. 3, were incubated with (lanes 2 and 4) or without (lanes 1 and 3) purified Dsk1 protein in the presence of [γ-32P]ATP. GST protein was also used in place of the GST fusion proteins as a negative control (lanes 5 and 6). Samples were resolved on an SDS–10% polyacrylamide gel and visualized by autoradiography. The bound Dsk1 phosphorylated Srp1 and Srp2 in the complex (lanes 1 and 3). (B) After the kinase reaction, Dsk1 is released from the Srp1/Dsk1 and Srp2/Dsk1 complexes. GST-Srp1/Dsk1 and GST-Srp2/Dsk1 protein complexes were incubated individually with (lanes 3, 4, 7, and 8) or without (lanes 1, 2, 5, and 6) an ATP regenerating system for 30 min at 23°C. Following the kinase reaction, protein-bound beads were pelleted by centrifugation. The supernatant (S) and bead (P) portions of each sample were resolved on an SDS–10% polyacrylamide gel and subsequently processed for immunoblotting with anti-T7-Tag MAb. Dsk1 was released from the complex to the supernatant in the presence of ATP (lanes 4 and 8), but it is not dissociated from the complex in the absence of ATP (lanes 2 and 6). Note the phosphorylated Srp proteins (indicated with a circled P) have slower mobility than that of their nonphosphorylated forms.
bound (P) and the released (S) fractions were analyzed by immunoblotting (Fig. 4B). Dsk1 was released from the complex to the supernatant after incubation with ATP (Fig. 4B, lanes 4 and 8), while no Dsk1 was released in the absence of ATP (Fig. 4B, lanes 2 and 6). Note that GST-Srp1 (Fig. 4B, lane 3) and GST-Srp2 (lane 7) migrated more slowly on the gel upon phosphorylation. Therefore, Dsk1 was dissociated from the complex after phosphorylating the GST-Srp1 or GST-Srp2 protein. Some Dsk1 protein was retained in the pellet fraction of the samples with ATP (Fig. 4B, lanes 3 and 7). This may be due to trapping of some released Dsk1 molecules in the pellet fraction since, once separated from the supernatant, the beads were not washed following the kinase reaction. Based on these results the Dsk1 reaction is dissected into three distinct steps: substrate binding, substrate phosphorylation, and release of the kinase from the product. Quantitative measurement for the percentage and rate of Dsk1 release from the complex has not been carried out.

Genetic interaction between prp2<sup>+</sup> and dsk1<sup>+</sup>. To understand the biological functions of Dsk1 protein kinase, it is necessary to investigate interactions of Dsk1 with the RS domain-containing proteins in vivo. For example, if Prp2 protein is an in vivo target of Dsk1, overexpression of prp2<sup>+</sup> may confer a phenotype, which is only apparent in the strain with the dsk1<sup>+</sup> gene. To test this, we placed prp2<sup>+</sup> gene under the control of a thiamine-repressible nmt1<sup>+</sup> (no message in thiamine) promoter of *S. pombe* (20, 21), so that Prp2 protein could be produced at a high level by growing cells in medium without thiamine. Consistent with a recent report (37), induction of prp2<sup>+</sup> expression from the nmt1<sup>+</sup>-driven plasmid, pREP1<sub>prp2</sub>, in wild-type cells leads to smaller colonies than those transformed with the vector pREP1 alone (data not shown). Exponentially growing cells in liquid culture were transferred to thiamine-depleted medium and grown for 21 h. The cells were then stained with DAPI and examined by phase-contrast (Fig. 5, top panels) and fluorescence (bottom panels) microscopy. Elongated cells were observed when the expression of the plasmid-borne prp2<sup>+</sup> gene was induced in the wild-type strain 1913; the average cell length increased about 60% (from 8.8 to 14.2 μm) compared to that of the cells harboring the vector pREP1 (Table 1). In addition, more than 40% of the cells had a cell length exceeding the regular range for 1913/pREP1 cells (Fig. 6). Although multiple nuclei were observed in some cells, many elongated cells seemed to have a single nucleus (Fig. 5, second column, bottom panel). In contrast, overexpression of prp2<sup>+</sup> gene in a dsk1-null strain (∆dsk1), B8, did not display any elongation phenotype (Fig. 5, third column) under the same condition; the average size of the cells (9.2 μm) was consistent with a recent report (37).
FUNCTIONAL CONSERVATION OF SR KINASES IN S. POMBE

1913/pREP1

% of Total Cells

Range of Cell Length (μm)

1913/pREP1

1913/pREP1ppr2−

B8/pREP1

B8/pREP1ppr2−

2A5/pREP1

2A5/pREP1ppr2−

FIG. 6. Size distribution of cell population in strains with ppr2+ gene overexpression. The cell length of the four samples in Fig. 5 was measured. The cell populations with a size range as indicated for 1913/pREP1, 1913/pREP1ppr2−, B8/pREP1ppr2−, and 2A5/pREP1ppr2− are displayed as histograms. A population of cells longer than 16 μm was observed in 1913/pREP1ppr2− and 2A5/pREP1ppr2−. The distribution pattern of 2A5/pREP1ppr2− is similar to that of 1913/pREP1ppr2−, while the pattern of B8/pREP1ppr2− resembles that of the negative control, 1913/pREP1.

remained similar to that of the wild-type strain, i.e., 1913/pREP1 (8.8 μm) (Table 1). Moreover, the “elongated” population as seen in 1913/pREP1ppr2− disappeared in strain B8/pREP1ppr2− (Fig. 6). Therefore, the elongation characteristic of ppr2+ overexpression requires the presence of dsk1+ gene.

To address the specificity of the genetic interaction between dsk1+ and ppr2+, we examined the ppr2+ overexpression phenotype in another kinase-deletion strain. The S. pombe kath+ gene (GenBank accession number Q10156) encodes a protein closely related in sequence to mammalian Clk/Sty. Interestingly, overexpression of the kath+ gene in S. pombe leads to branched cells with multiple septa and nuclei, which was different from the phenotype conferred by dsk1− overexpression (unpublished data). Since the name kath+ is not conventional nomenclature for a S. pombe gene, we changed it to kic1+ for “kinase in Clk” family. The kic1+ gene was disrupted, and a haploid strain with a null allele was found to be viable (unpublished data). The pREP1ppr2+ plasmid was transformed into a kic1-null mutant strain (Δkic1), 2A5. Similar to wild type, overexpression of ppr2+ gene in the kic1-null mutant strain, 2A5 (Δkic1), resulted in elongated cells (Fig. 5, fourth column). The average size of the cells in 2A5/pREP1ppr2+ was 16.8 μm, increased approximately 90% compared to 8.8 μm in 1913/pREP1 (Table 1). An “elongated” population representing more than 40% of the cells was again observed (Fig. 6). Thus, cell elongation caused by Prp2 overproduction is specifically dependent on the presence of the dsk1+ gene but does not require the kic1+ gene. These in vivo results substantially support the notion that Prp2 protein is a target of Dsk1 action in fission yeast and reinforce the in vitro data demonstrating the binding of the two proteins and phosphorylation of Prp2 protein by Dsk1.

The ppr2+ overexpression phenotype in strains 1913 and 2A5 displayed two distinct populations, one with normal length distribution and the other elongated. This is perhaps due to the leakiness of the ppr2+ overexpression phenotype. This is consistent with the observation that overexpressing ppr2+ did not kill the cell but instead produced smaller colonies. The dual population phenomenon indicates that the ppr2+ overexpression may block cell cycle progression only part of the time. Alternatively or additionally, it suggests that the ppr2+ overexpression may affect multiple steps of the cell cycle. Since plasmid in S. pombe is not stable, cells that lost the pREP1-ppr2+ plasmid might also contribute to the population with apparently normal cell sizes. Finally, because the ppr2+ overexpression was induced in an asynchronous cell population, a portion of the cells may already pass the elongation phase and resume the normal cell cycle.

Human SRPK1 protein is a functional homologue of fission yeast Dsk1 protein in vivo. Sequence analysis and kinase assays indicate that S. pombe Dsk1 is homologous to human SRPK1 (12). We here tested the functional similarity between Dsk1 and SRPK1 in vivo. When the dsk1+ gene is overexpressed, it results in highly elongated cells with a delay in the progression from G2 to M phase (46). Thus, we first examined whether overexpression of the human SRPK1 gene in fission yeast would produce a phenotype similar to that of dsk1+ overexpression. Plasmid pREP1 SRPK1 was introduced into a wild-type S. pombe strain, and the expression of the SRPK1 gene was induced in the absence of thiamine. The majority of the SRPK1-overproducing cells became elongated (Fig. 7A, right panel) compared to cells containing only the pREP1 vector (left panel). Therefore, like dsk1+ overexpression, overexpression of the human SRPK1 gene in S. pombe leads to elongated cells that are indicative of a delay at the G2/M-phase transition.

One stringent evaluation for functional homology is complementation of the loss of one gene by another gene. We anticipated that if human SRPK1 is a true functional homologue of Dsk1, it should compensate for the loss of Dsk1 in the cell. The genetic complementation test had not been accomplished because dsk1− is not essential for the viability of the cell. Interestingly, dsk1− mutant yeast cells became very sick when a related kinase gene in the cell, kic1−, was also disrupted. With the double deletions, cells grew extremely slowly and formed microcolonies (see Fig. 7B). Taking advantage of this recent finding, we transformed the dsk1−/kie1 double-null mutant, 2D4 (Δdsk1−Δkie1), with either pREP1 or pREP1 SRPK1. Since the nmt1+ promoter is leaky, a considerable amount of expression occurs even in the presence of thiamine (28). Transformants were first selected and subsequently restreaked for growth analysis on thiamine-containing plates (repressed condition). Cells carrying pREP1 SRPK1 formed healthy colonies (Fig. 7B, right panel), whereas cells containing the pREP1 vector alone...
RS domain-containing proteins Prp2, Srp1, and Srp2 in vitro. Consistent with its substrate specificity, Dsk1 forms kinase-competent complexes with those RS domain-containing proteins. The kinase-substrate interaction is supported by the in vivo evidence for the dependency of prp2\textsuperscript{+} overexpression phenotype on dsk1\textsuperscript{-} gene. Despite the evolutionary gap separating fission yeast and human, SRPK1 not only shares similar biochemical properties with Dsk1 but also compensates for the loss of Dsk1 in fission yeast cells. The functional conservation of these kinases at the molecular and cellular level illustrates the importance of the SR protein-specific kinases in eukaryotic systems.

The evidence accumulated in recent years indicates that SR networks exist in the fission yeast \textit{S. pombe} which consist of RS domain-containing proteins and their kinases. Our studies suggest that the phosphorylation patterns and interactions of the SR networks are conserved from fission yeast to mammals. We have shown for the first time that all four \textit{S. pombe} RS domain-containing proteins, including Prp2, Srp1, Srp2, and Rsd1 (T.-L. Tseng and A. R. Krainer, personal communication), are phosphorylated by Dsk1 in vitro, and these phosphorylated proteins are recognized by 3C5 MAb (this report; Z. Tang, R.-J. Lin, T.-L. Tseng and A. R. Krainer, unpublished data), indicating that the kinase reaction generates a phosphoepitope identical to that found in mammalian SR proteins. In agreement with the in vitro observation, Rsd1 isolated from wild-type fission yeast is also recognized by MAb 3C5 (T.-L. Tseng and A. R. Krainer, personal communication), providing in vivo evidence for the conservation of phosphorylation specificity.

Here we dissected the Dsk1-mediated kinase reaction in vitro into three discrete steps: substrate binding, substrate phosphorylation, and release of the Dsk1 from the complex after phosphorylating its substrate. Thus, Dsk1 forms transient complexes with RS domain-containing proteins in the presence of ATP. Similar kinase-substrate complexes were recently observed between human SRPKs and SR proteins. Both SRPK1 and SRPK2 bind and subsequently phosphorylate GST-SF2/ASF. The expression of a kinase-deficient mutant SRPK2 leads to trapping SF2/ASF in the cytoplasm, possibly by forming a stable complex between the two proteins (16).

The phosphorylation by Dsk1 may affect the interactions between and/or the activity of these proteins in splicing. In agreement, pre-mRNA splicing is partially impaired in dsk1 deletion strain of \textit{S. pombe} (unpublished data), and the interaction of Srp1 and Srp2 proteins is inhibited by Dsk1-mediated phosphorylation in vitro (2; Tang et al., unpublished data). It has been shown that the phosphorylation status of SF2-ASF exerts distinct effects on its association with various protein targets in vitro (55). Additionally, changes in SR protein phosphorylation play a role in the activation of pre-mRNA splicing during early development in the nematode (43). It was also established in \textit{Drosophila} that SR protein phosphorylation is essential for developmentally regulated alternative splicing (7).

Dsk1 influences the activity of Prp2 in vivo. Overexpression of \textit{prp2}\textsuperscript{+} in different strains of \textit{S. pombe} demonstrated that the ability of Prp2 to cause cell elongation is Dsk1-dependent (Fig. 5 and 6). Moreover, the observation that \textit{kic1}\textsuperscript{-} gene does not have obvious effect on the phenotype of \textit{prp2}\textsuperscript{+} overexpression substantiates the specific interaction between Dsk1 and Prp2. The effect on Prp2 probably is through phosphorylation of Prp2 by Dsk1, especially that Dsk1 displays high activity in phosphorylating Prp2 in vitro (47). It will be very interesting to investigate whether Dsk1 is indeed required for the phosphorylation of Prp2, Srp1, and Srp2 in vivo. The phosphorylation levels of these target proteins can be determined by in vivo \textsuperscript{32}P labeling of wild-type and dsk1-null mutant cells followed by

**DISCUSSION**

Together with our previous studies (47), we show in this report that Dsk1 protein specifically phosphorylates \textit{S. pombe}
immunoprecipitation of individual proteins with antibodies specifically against each protein. Alternatively or additionally, it can be done by using 3C5 MAb, which is specific to the SR-phosphoepitope, to probe these target proteins isolated from wild-type and dsk1-null mutant cells. We plan to address this important issue in the future.

It was reported that SRPK1 and Clk/Sty also phosphorylate human U2AF65 protein in vitro (4), although the consequence of the phosphorylation on the function of U2AF65 is not known. Perhaps Dsk1-mediated phosphorylation changes the ability of Prp2/Mis11 protein to interact with other splicing factors, such as the fission yeast homologue of human U2AF35, spU2AF23 (51). Therefore, as in mammalian systems, phosphorylation and/or dephosphorylation of RS domain-containing proteins may regulate the properties of these proteins and the organization of the protein relay in fission yeast.

We performed the first cross-species test for viability complementation of SR protein-specific kinases. Since the discovery of human SRPK1 (12), members of SRPK and Clk/Sty families were identified from various eukaryotic organisms, including mammals, Drosophila, and yeasts, based on sequence analysis and kinase specificity (4, 5, 18, 44, 47, 50, 57). Recently, the SRPK homologue in Saccharomyces cerevisiae, Sky1, was shown to phosphorylate Npl3, a budding yeast RNA binding protein containing SR/RS diad repeats (44) and several mammalian SR proteins in vivo (56). The phosphorylation by Sky1 affects the cellular localization and protein interactions of these mammalian SR proteins in yeast cells (56).

Interestingly, mammalian SRPK1 and Clk/Sty specifically substitute the activity of Sky1 in mediating RS domain interactions in vivo (56). However, the viability complementation strategy had not been applied to measure the functional similarity of these protein kinases prior to this study, perhaps partly due to their redundancy in cells, so that single mutation in one protein kinase lacks the apparent phenotype. Our genetic result exhibited that human SRPK1 compensates for the loss of the S. pombe Dsk1 in vivo and thus is a functional homologue of Dsk1. Collectively, these data provide both in vitro and in vivo evidence for the conservation of the SR networks through evolution.

A common feature shared between Dsk1 and Prp2 is their dual functional potential. Dsk1 protein plays a role in mitotic control (46) and is an SR protein-specific kinase involved in pre-mRNA splicing (12, 47). Prp2 is essential for pre-mRNA splicing (34, 35) and also affects chromosome segregation (45). A similar type of dual functional feature is also found in other proteins such as Ran, a small guanosine triphosphatase. It was recently shown that Ran functions to trigger the formation of the mitotic spindle, in addition to its well-characterized role in nuclear trafficking (32, 52). Another example is fission yeast Cdc5 protein, which is required for G2/M transition and is a component of a 40S snRNP-containing complex essential for pre-mRNA splicing (22). The putative dual functions of Dsk1 and Prp2 in both splicing and the cell cycle may be fulfilled through the action of Dsk1 on Prp2/Mis11, since we demonstrated that Dsk1 and Prp2/Mis11 proteins genetically interact with each other. It is conceivable that differential phosphorylation of Prp2/Mis11 may regulate its ability to either participate in chromosome segregation or to be engaged in splicing. Phosphorylation by Dsk1 may also modulate the activity of Prp2/Mis11 through altering its cellular or subnuclear localization. In agreement with its connection to the cell cycle, the phosphorylation state, cellular localization, and kinase activity of Dsk1 all change in a cell-cycle-dependent fashion (46). Supporting the model in the differential effects of phosphorylation, distinct phosphorylation sites on budding yeast transcription factor Pho4 play separable roles in altering its subcellular localization and interaction with another transcription factor, providing multiple levels of regulation to control the activity of Pho4 (17). Thus, studying the S. pombe RS domain-containing proteins and their kinases may help determine the regulatory pathways that link pre-mRNA splicing with the cell division cycle.

Our studies provide novel information about the fission yeast SR networks. The functional conservation of SRPKs from fission yeast to human S. pombe a valuable system for studying the biological roles of the kinase family. The powerful genetics of S. pombe will facilitate the elucidation of functions of the SR networks in eucaryotic gene expression.

ACKNOWLEDGMENTS

We thank Xiang-Dong Fu (University of California at San Diego) for providing the human SRPK1 gene, Mitsuhito Yanagida (Kyoto University, Kyoto, Japan) for dsk1-null mutant strain, and Paul Salavaterra for advice in microscopic analysis. We also thank Xiang-Dong Fu, David Horowitz, Adam Bailis, and Glenn Manthey for critical reading of the manuscript and for constructive suggestions. We thank the reviewers for their valuable suggestions for improving the manuscript. This work was supported by City of Hope Beckman Endowment Grant.

REFERENCES

28. Murone, M., and V. Simanis. 1996. The fission yeast dma1 gene is a component of the spindle assembly checkpoint, required to prevent septum formation and premature exit from mitosis if spindle function is compromised. EMBO J. 15:6605–6616.