Genetic and physical interaction of Ssp1 CaMKK and Rad24 14-3-3 during low pH and osmotic stress in fission yeast

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1. Summary

The Ssp1 calmodulin kinase kinase (CaMKK) is necessary for stress-induced reorganization of the actin cytoskeleton and initiation of growth at the new cell end following division in Schizosaccharomyces pombe. In addition, it regulates AMP-activated kinase and functions in low glucose tolerance. ssp1 cells undergo mitotic delay at elevated temperatures and G2 arrest in the presence of additional stressors. Following hyperosmotic stress, Ssp1-GFP forms transient foci which accumulate at the cell membrane and form a band around the cell circumference, but not co-localizing with actin patches. Hyperosmolarity-induced localization to the cell membrane occurs concomitantly with a reduction of its interaction with the 14-3-3 protein Rad24, but not Rad25 which remains bound to Ssp1. The loss of rad24 in ssp1 cells reduces the severity of hyperosmotic stress response and relieves mitotic delay. Conversely, overexpression of rad24 exacerbates stress response and concomitant cell elongation. rad24 does not impair stress-induced localization of Ssp1 to the cell membrane, however this response is almost completely absent in cells overexpressing rad24.

2. Introduction

Environmental stress causes massive changes to cell physiology, including major metabolic, cytoskeletal and transcriptional responses. Changes in osmolality, temperature, reactive oxygen species and nutrient status stimulate conserved stress-activated Spc1 MAPK pathways [1–5]. In fission yeast, the Wak1 and Win1 [6,7], Wis1 [8,9] and Spc1 MAPK cascade [9,10] relays environmental signals to the nucleus. Phosphorylated Spc1 MAPK enters the nucleus and activates transcription factors, inducing stress response genes including gpd1 (glycerol-3-phosphate dehydrogenase) and tps1 (trehalose-6-P synthase), increasing intracellular concentrations of glycerol and trehalose [10–13]. MAPK signalling impinges on the cell cycle via Skt1, which phosphorylates the mitotic activator Cdc25, inducing 14-3-3 dimer binding and nuclear export of Cdc25 thus reducing the opportunity to activate its CDK nuclear substrate, Cdc2 [14]. spc1 cells experience G2 delay under normal conditions and G2 arrest after osmotic or oxidative stress [9,15,16].

The CaMKK and CaMKI and CaMKV calcium–calmodulin (Ca2+/CaM)-dependent signalling cascade is highly conserved and involved in a number of important cellular processes including cell cycle and neuronal- and immune-cell function. CaMKKs phosphorylate and fully activate Ca2+/CaM-bound
3. Results

3.1. rad24 deletion suppresses the cell-cycle phenotype of ssp1 cells at high temperatures

We identified the 14-3-3 homologues Rad24 and Rad25 [31] multiple times in a yeast two-hybrid screen using full-length Ssp1 as a bait protein (data not shown), corroborating previous mass spectrometry data [19]. 14-3-3 proteins inhibit CaMKKα in mammalian systems [32] and are directly linked to the control of cell-cycle progression by regulating the Cdc2/Cdc13 activator Cdc25 [33] and inhibitor Wee1 [34–36]. In fission yeast, neither 14-3-3 isoform is essential; however, the double deletion is lethal [31]. To test for the influence of Rad24 on the mitotic delay of ssp1 cells at high temperatures [20,23], ssp1 rad24− (Q4101; table 1) and ssp1 rad25− (Q4104) cells (YEA) were shifted from 30 to 36 °C for 4 h (figure 1a). Loss of rad24 is epistatic with respect to the heat-stress-dependent cell elongation phenotype of ssp1− cells at 36°C. Loss of Rad25 has no effect, presumably owing to the small proportion of the rad25 14-3-3 isoform in the overall pool of 14-3-3 proteins (see figure 7d).

3.2. Overproduction of Rad24 and Ssp1 results in an additive cell-cycle phenotype

Plasmid-borne rad24-GFP or GFP-ssp1 (pR2-22) [23], both under control of the strong thiamine-repressible nmt1 promoter (OP-rad24-GFP and OP-ssp1-GFP), were expressed for 20 h in wild-type, ssp1− and rad24− cells, respectively (Q4105, Q4106, Q4107, Q4108) (figure 1b). In an otherwise wild-type background at 25°C, overproduction of Rad24 causes moderate cellular elongation, whereas overproduction of Ssp1 makes cells shorter [23]. Conversely, overproduction of Rad24 at 25°C in ssp1− cells leads to occasional branching, exacerbated at 35°C with extremely elongated cells often displaying aberrant branched morphology. The size reduction from OP-GFP-ssp1 is more conspicuous in rad24− cells, which become spherical. Overexpression of rad24 thus has an additive phenotype with ssp1−, exacerbating the cell elongation phenotype.

At 30°C, wild-type cells expressing a single chromosomally integrated copy of rad24-His6 under control of the nmt1 promoter (OP-rad24-His6) (Q4109) are significantly longer (21.7 ± 4.1 μm) than if expression is repressed (13.2 ± 1.1 μm). By contrast, cells with single-copy expression of GFP-ssp1 under control of the nmt1 promoter (OP-ssp1-GFPint) (Q4111) are significantly shorter (12.9 ± 1.0 μm) than if expression is repressed (13.9 ± 1.2 μm). An intermediate size (19.2 ± 2.4 μm) is found for cells co-overexpressing ssp1 and rad24 (Q4111), suggesting that these gene products antagonize each other in some way (figure 1c).

To compare growth rates, thiamine-repressed OP-GFP-ssp1int and/or OP-rad24-His6int cells were washed in EMM before plating diluted aliquots (EMM, EMM+1 M KCl and EMM pH 3.5; all ± thiamine) (figure 1d), followed by incubation for 5 days (30 or 37°C). At both temperatures, cells overexpressing rad24 displayed inhibited growth compared with cells overexpressing ssp1 or both rad24 and ssp1. Overexpression of rad24 negatively affects proliferation, resistance to hyperosmotic and low pH stress, and increases mitotic delay. Growth inhibition in OP-rad24-His6int cells was exacerbated by 1 M KCl (30 and 37°C) and OP-GFP-ssp1int did not alleviate this effect. Similar growth delay in rad24 overexpressing cells was evident on EMM pH 3.5 media (30 and 37°C) and was not ameliorated by ssp1 overexpression. Low pH conditions may affect nutritional status in these cells. Failure of CaMKK to activate AMPK [24] in cells with a drop in energy may consequently inhibit processes such as protein synthesis and growth, and affect downstream regulators.

3.3. Loss of ssp1 reduces the restrictive temperature for cdc25-22

The cdc25-22Δ (Q300) allele at the restrictive temperature (36–36.5°C) arrests at the G2/M boundary as very elongated single cells [33,37,38]. The ssp1− background exacerbates the cell elongation phenotype of cdc25-22Δ at semi-permissive temperatures (32°C) (figure 2a,b). As ssp1− cells are sensitive to KCl stress and Cdc25 is exported out of the nucleus within 10 min after KCl stress [39], we investigated the effect of ssp1− on the cdc25-22Δ (Q300) strain.
on the nuclear localization of Cdc25. We expressed single-copy, native promoter-integrated cdc25-GFP (cdc25-GFPint) [40] in rad24− ssp1− (Q4113) and ssp1− (Q3974) backgrounds at 25°C and shifted to 35°C for 4 h. The nuclear localization of Cdc25-GFP is independent of ssp1− and rad24− single and ssp1− rad24− double gene deletions in both conditions (figure 2c), suggesting that the loss of ssp1 does not interfere with the nuclear localization of Cdc25-GFP.

3.4. Deletion of rad24 overrides growth sensitivity after 0.6 M KCl stress in ssp1− cells

At high temperatures and in the presence of 0.6 M−1.5 M KCl or low pH (3.5), ssp1− cells arrest at the G2/M boundary (figure 3a) [23]. rad24− alleviates this arrest at high temperatures on YEA and in the presence of KCl. rad24− is therefore epistatic with respect to ssp1− cell-cycle arrest following 0.6 M KCl stress at 36°C. Proliferation at pH 3.5 at 36°C is not rescued and small cell size suggests a block to growth (figure 3b). The response of Ssp1 to 0.6 M KCl stress and to low pH probably occurs through different mechanisms.

3.5. At native expression levels Ssp1-GFP localizes to the cell membrane after KCl stress

GFP-Ssp1 localization was previously examined following strong overexpression (nmt1) from a multi-copy plasmid

| strain | genotype | source |
|--------|-----------|--------|
| Q1618  | URA4::lexAop-lacZ/8LEXA-ADE2::URA3 ura3-1/ura3-1 leu2-3/leu2-3 his-3/11/11 trp1-1/1 trp1-1/1 ade2-2/ade2-2 can1-1/1 can1-1 | laboratory stock |
| Q250   | wild-type (972 h) | laboratory stock |
| Q3677  | leu1-32 ura4-D18 | laboratory stock |
| Q4101  | ssp1::ura4− leu1-32 ura4-D18 | Toda laboratory |
| Q1537  | ssp1::sup3-5 ade-6-704 ura4-D18 leu1-32 | laboratory stock |
| Q4101  | ssp1::ura4− rad24::ura4− ura4-D18 leu1-32 | this study |
| Q4102  | rad24::ura4− ura4-D18 leu1-32 | Carr laboratory |
| Q4103  | rad25::kanMX6 ura4-D18 leu1-32 | this study and Carr laboratory |
| Q4104  | ssp1::ura4− rad25::kanMX6 ura4-D18 leu1-32 | this study |
| Q4105  | pREP1-rad24GFP in ssp1::sup3-5 ade6-704 leu1-32 ura4-D18 | this study |
| Q4106  | pREP1-rad24GFP in leu1-32 ura4-D18 | this study |
| Q4107  | pRH2-22 (nmt1:GFP-ssp1) in leu1-32 ura4-D18 | this study and laboratory stock |
| Q4108  | pRH2-22 in rad24::ura4− ura4-D18 leu1-32 ade6-704 | lab stock |
| Q4109  | nmt1:rad24-His6int ura4-D18 leu1-32 | this study |
| Q4110  | nmt1:rad24-His6int nmt1:GFP-ssp1int ura4-D18 leu1-32 | this study |
| Q4111  | nmt1:GFP-ssp1int ura4-D18 leu1-32 | this study |
| Q4112  | cdc25-GFPint ura4-D18 leu1-32 | laboratory stock |
| Q3974  | cdc25-GFPint ssp1::sup3-5 ade6-704 ura4-D18 leu1-32 | this study |
| Q4113  | cdc25-GFPint rad24::ura4− leu1-32 ura4-D18 | laboratory stock |
| Q300   | cdc25-22 leu1-32 ura4-D18 ade6-10 | laboratory stock |
| Q1530  | cdc25-22 ssp1::sup3-5 ade6-704 ura4-D18 leu1-32 | laboratory stock |
| Q4114  | ssp1-GFPint ura4-D18 leu1-32 | this study |
| Q4115  | ssp1-GFPint rad25::ura4+ ura4-D18 leu1-32 | this study and Carr laboratory |
| Q4116  | ssp1-GFPint rad24::ura4+ ura4-D18 leu1-32 | this study and Carr laboratory |
| Q4117  | ssp1-GFPint ap3YC-YFPint leu1-32 ura4-D18 | this study and Nolen |
| Q4118  | ssp1-GFPint rad24::2HA-His6 (ura4+ int) leu1-32 ura4-D18 | this study and Russell laboratory |
| Q4119  | ssp1-GFPint rad25::His6 ura4-D18 leu1-32 | this study |
| Q4120  | ssp1-GFPint rad24::2HA-His6 (ura4+ int) rad25::His6 leu1-32 ura4-D18 | this study and Russell laboratory |
| Q4121  | ssp1-GFPint rad24::ura4+ ura4-D18 leu1-32 | this study |
| Q4122  | nmt1::GFP-ssp1int ura4-D18 leu1-32 | this study |
| Q4123  | ssp1-GFP::kanMX6int ura4-D18 leu1-32 | this study |
| Q4124  | ssp1-GFP::kanMX6int ura4-D18 leu1-32 | this study |

Table 1. List of strains.
30 and 36°C as indicated and incubated at 30 and 37°C. There is no significant difference in cell lengths of ssp1Δ rad24Δ cells at 30 and 36°C (p = 0.198). There is a significant difference in cell lengths of ssp1Δ rad25Δ cells at 30 and 36°C (p < 0.05) (all n ≥ 37; all Student’s t-test).

The nucleus appeared compressed and irregular in cells overexpressing (23). Cells overexpressing ssp1-GFPint are significantly shorter in the absence than in the presence of thiamine (p < 0.05) (all n ≥ 117; Student’s t-test). Cells (30°C, EMM + thiamine) were washed with EMM and derepressed for 24 h. Cells overexpressing GFP-ssp1int are significantly shorter in the absence than in the presence of thiamine (p < 0.05) (all n ≥ 37; all Student’s t-test).

Vaccules are faintly discernible, suggesting that Ssp1-GFP levels are lower than those in the cytoplasm (figure 4c,d). After switching from YEA to YEA + 0.6 M KCl, cell shapes became more jagged and fission scars became more notable, with a slight decrease in cell volume. Reversal of this cell ‘shrivelling’ with the re-establishment of cell turgor depends upon glycerol synthesis and requires gpd1Δ [42]. The nucleus appeared compressed and irregular in shape. Vacuolar areas became more conspicuous, with low fluorescence intensity similar to the nucleus. Within 2–3 min of 0.6 M KCl treatment, small areas of increased Ssp1-GFP fluorescence (foci) in the cytoplasm and near the cell walls and septum appeared. These foci brightened further and by 7 min, additional fluorescent foci along the cell membrane appeared. These foci brightened further and by 7 min, additional fluorescent foci along the cell membrane appeared, especially at the cell tips (figure 4c,d). Localization of Ssp1-GFP at the cell membrane started to decline by 20 min, decreased further by 34 min and was similar to cytoplasmic levels by 70 min. Cells accommodated to the increase in extracellular osmolytes, regaining volume and rod-shaped cell morphology at 23–34 min, indicating induction of gpd1Δ and glycerol synthesis [42]. Surface plots of fluorescence intensity of cells at t = 0, t = 17, t = 34 and t = 70 min during 0.6 M KCl stress (figure 4d) highlights the formation of the small foci of fluorescence intensity in the stressed cells. The conspicuous placement of fluorescent foci, often on opposing sides along the cylindrical portion of the cell suggested that Ssp1-GFP localization to this area takes place not only as distinct foci, but instead as a ring. Although overall fluorescence declined, Ssp1-GFP protein levels stayed constant (figure 4e).

Expression of multi-copy GFP-ssp1 from the nmt1 promoter (Q4108) facilitated the capture of a multi-image Z-stack. In EMM + 1.5 M KCl (25°C), a band of foci formed around the
circumference of the cell (figure 5a). This band was not visible in control cells. To determine whether Ssp1 co-localizes with actin patches, we co-expressed single-copy, integrated ssp1-CFP and arp3C-YFP (Q4117) which co-localizes with cortical actin patches. Mid-logarithmic cells (YEA, 25°C) were treated with YEA or YEA + 0.6 M KCl. Projection images of deconvolved Arp3C-YFP (Slidebook; Z-stack) and single Ssp1-CFP images revealed that after 15 min of osmotic stress, the vast majority of actin patches in the cell membrane and/or cell wall area do not co-localize with the accumulated Ssp1-CFP at the cell membrane (figure 5b).

3.6. Loss of rad24 or rad25 does not affect localization of Ssp1-GFP after stress

To investigate how loss of 14-3-3 proteins affects the subcellular localization of Ssp1-GFP, we expressed ssp1-GFPint in rad24− (Q4116) or rad25− (Q4115) cells. In YEA (30°C), Ssp1-GFP in mid-logarithmic rad24− or rad25− cells was cytoplasmic and excluded from the nucleus. Ssp1-GFP accumulated along forming and formed septa, and along the cell membrane in a subset of cells. After osmotic stress (YEA, 0.6 M KCl) Ssp1-GFP promptly localized to areas near the cell membrane, forming foci of fluorescence as in rad24+ rad25+ cells (figure 6a), indicating that Ssp1-GFP localization to the cell membrane after 0.6 M KCl stress does not require rad24− or rad25−.

3.7. Ssp1-GFP co-immunoprecipitates with the 14-3-3 proteins Rad24-2HA-His6 and Rad25-His6

14-3-3 proteins are abundant in all cells, and in budding yeast interact with at least 271 proteins representing approximately 4.4% of the proteome [44] indicating that only a small amount of the total pool of 14-3-3 associates with any one protein at any given time. Only a portion of the total Ssp1-GFP present in the lysate immunoprecipitated in the Rad24-2HA-His6 pulldown. To test the relative binding of Rad24 or Rad25 to Ssp1-GFP, protein lysates were prepared from mid-logarithmically growing (YEA, 30°C) cells expressing single-copy ssp1-GFPint and either rad24−2HA-His6 (Q4118) or rad25−His6 (Q4119) integrated at their native promoters. Ssp1-GFP co-immunoprecipitates with Rad24 and Rad25 (figure 6b,c) and a portion of each of the Ssp1 and 14-3-3 pools co-immunoprecipitate.
In budding yeast, the deletion phenotype of the major 14-3-3 isoform BMH1 is complemented by overexpression of the minor isoform BMH2, suggesting that these proteins have similar binding partners [45]. To determine the relative amounts of Rad24 and Rad25 associating with Ssp1, we expressed ssp1-GFP rad24-2HA-His6 rad25-His6 (Q4120), where Rad24-2HA-His6 is distinguishable from Rad25-His6 by size owing to small differences in molecular weight as well as the presence of the 2HA. Rad24-2HA-His6 was present at approximately five-fold higher levels (IMAGEJ) compared with Rad25-His6 protein in the cell lysate (figure 6d) and a similar ratio of Rad24-2HA-His6 and Rad25-His6 co-precipitated with Ssp1-GFP.

Ssp1-GFP is a highly phosphorylated protein [19]. We can see doublet formation on gels (see immunoprecipitates; figure 6c) and can separate these bands by running the SDS-PAGE gel for an extended amount of time. The phosphorylation state of Ssp1-GFP will be discussed below.

### 3.8. Rad25-His6 and Ssp1-GFP physically interact in the absence of Rad24

For the previous co-immunoprecipitation studies, Rad24-2HA-His6 or Rad25-His6 proteins were co-expressed with Ssp1-GFP. We wanted to determine whether Rad25 associates with Ssp1-GFP only as part of a heterodimer with Rad24 or is able to bind Ssp1 as a Rad25–Rad25 homodimer in vivo. Immunoprecipitation of Ssp1-GFP and Rad25-His6 in rad24+ and rad24– (Q4121) cells showed that Rad25-His6 co-precipitates with Ssp1-GFP in the absence of Rad24 (figure 6e). We also found that Ssp1-GFP is less stable in a rad24– background (figure 6e,f).

### 3.9. Stress reduces the interaction between Ssp1-GFP and Rad24-2HA-His6

Harvesting cells by centrifugation at 4°C, followed by washes in ice-cold lysis buffer exposes cells to stressors, including increased gravitational forces and hypoxia. This is manifested by transient increased phosphorylation of MAPK Spc1 [46,47] and Atf1 [46]. Cells undergo a brief cell-cycle delay similar to but shorter than the delay after 0.6 M KCl [23,46] and there is a transient depolarization of actin [46]. Hypoxia in pelleted cells causes activation of hypoxia response genes via Sre1 [48,49]. Moderate thermal downshift (28–15°C) brings about phosphorylation of Spc1 and induction of the stress response genes ctt1, tps1 and ntp1 [46]. To minimize these stressors, cells were treated with pre-warmed YEA (30°C; + KCl to 0.6 M KCl) for 15 min, then rapidly chilled to 0°C with frozen, crushed YEA (± 0.6 M KCl) preceding centrifugation. Very low temperatures greatly delay Spc1 phosphorylation [46]. Ssp1-GFP localizes to the cell membrane after 15 min of 0.6 M KCl stress and activating phosphorylation of Spc1 MAPK is detected [10]. After KCl treatment approximately 10 times less Rad24-2HA-His6 co-immunoprecipitated with Ssp1-GFP in the KCl-treated immunoprecipitate than in the untreated controls (figure 7a), however we consistently failed to detect a decrease in the total amount of Rad25-His6 protein that co-immunoprecipitated with Ssp1-GFP (figure 7b). Ssp1-GFP

![Figure 3. Loss of rad24 relieves cell-cycle delay and KCl stress sensitivity in ssp1− cells at 36°C.](image-url)
binds Rad24-2HA-His6 and Rad25-His6 during unperturbed growth in vivo, however bound Rad24-2HA-His6 but not Rad25-His6 decreases substantially after 0.6 M KCl stress treatment relative to unperturbed conditions in vivo. The limitations of immunoblotting do not allow detection of very subtle changes in Rad25-His6 binding.

3.10. Overexpression of rad24 diminishes Ssp1-GFP localization to the cell membrane

Loss of either 14-3-3 homologue perturbs MAPK hyperosmolarity stress-dependent signalling in budding yeast [44]. BMH2 physically interacts with the NHA1 antiporter protein

Figure 4. Ssp1-GFP expressed as a chromosomal integrant on its native promoter localizes to the cell membrane following 0.6 M KCl stress. (a,b) ssp1-GFPint cells are phenotypically wild-type. (a) Cells (YEA, 30 °C) were diluted to 10⁶, 10⁵, 10⁴, 10³ ml⁻¹ and 5 μl of suspension were spotted onto media and incubated for 5 days (30 and 36 °C). (b) Wild-type and ssp1-GFPint cells (YEA, 30 °C) were incubated for 4 h at 30 and 36 °C. (c) ssp1-GFPint (YEA, 30 °C) cells were analysed in a microfluidic growth chamber supplied with fresh YEA at room temperature. YEA + 0.6 M KCl added at t₀ induced hyperosmotic stress. Cells were imaged 11 times from t = 0 to 70 min. Some images were omitted for the sake of brevity. (d) Surface fluorescence intensity plots of ssp1-GFPint cells. Single-plane images of cells from figure 5c at t = 0 (no stress) and t = 7, t = 34 and t = 70 min (0.6 M KCl stress) were analysed further (surface plot function; IMAGEJ). (e) Ssp1-GFP protein levels after addition of KCl to 0.6 M. Cells were harvested at the indicated times. WBα, western blot with antibody.
at the membrane in a HOG1-independent manner. Loss of BMH1 increases sensitivity to NaCl, KCl and LiCl without affecting plasma membrane potential [45]. In Schizosaccharomyces pombe, MAPK activation after osmotic stress leads to phosphorylation of Cdc25 by Srk1, binding of 14-3-3 and Cdc25 nuclear export [14]. Loss of rad24 or rad25 does not prevent the stress-dependent localization of Ssp1-GFP to the cell membrane, nor does it increase sensitivity to temperature, low pH or KCl stress (figure 3a, b). Strong expression of multi-copy pREP1:rad24-His6 in ssp1-GFPint cells (Q4124) (EMM, 30°C) impairs Ssp1-GFP accumulation at the membrane after 1 M KCl treatment (figure 7c). In rad24+ cells expressing single-copy integrated GFP-ssp1 under control of the nmt1 promoter (Q4111), the majority accumulate GFP-Ssp1 at the membrane after KCl stress. This localization pattern is suppressed by co-overexpression of single-copy, integrated rad24-His6 under control of the nmt1 promoter, where this response is only detected in approximately 12% of the cell population (figure 7d).

3.11. The phosphorylation state of Ssp1 is not altered after 0.6 M KCl stress

Like all CaMKKs [18], Ssp1 is regulated by phosphorylation. Ssp1 has potential phosphorylation sites at Y58, S59, Y63, T82 and S94, where S59 and S94 were identified in phosphopeptides containing only one phosphorylatable Ser residue [19,50]. Ssp1 is dephosphorylatable in vitro [19]. We investigated whether stress response causes changes in Ssp1 phosphorylation. To preserve the phosphorylation state of Ssp1-GFP in unperturbed and KCl-stressed cells, we collected mid-logarithmically growing cells by gentle filtration [35,51] after 15 min treatment with pre-warmed YEA (+ KCl to 0.6 M, 30°C). We detected a higher mobility band in extracts treated with Lambda phosphatase (no phosphatase inhibitors; figure 7e, lane 3). The faint upper band and strong lower band represent Ssp1-GFP protein in a partially phosphorylated and dephosphorylated state. The lower band is absent in the mock-treated extract (lane 2), confirming that it is not due to proteolysis. There was no bandshift evident in extracts prepared in the presence of phosphatase inhibitors in either perturbed or KCl-treated cells (lanes 4 and 5). Bands corresponded to the position of the phosphorylated bands produced by the control lysates. Ssp1-GFP thus appears to be phosphorylated in both unperturbed (YEA, collected on filter [52] and stressed with 0.6 M KCl) cells in vivo. We obtained similar results when lysates were extracted from Ssp1-GFP cells after mild temperature stress (36°C) (not shown). Our results confirm basal phosphorylation of Ssp1.

4. Discussion

Regulation of CaMKK occurs through Ca2+/CaM binding and inhibitory and activating changes in phosphorylation...
Additional inhibition takes place through 14-3-3 binding [17,32]. Here, we explore the interaction of CaMKK Ssp1 with 14-3-3 and their role in cell-cycle regulation and stress response.

4.1. Genetic interaction with 14-3-3 links CaMKK Ssp1 to cell-cycle control machinery

The CaMKK Ssp1 is a mitotic activator [20,23]. Its link to the cell-cycle machinery is supported by suppression of the ssp1'' mitotic delay by rad24'' and the elongation and arrest of ssp1'' cells when overexpressing rad24. Mitotic advancement can be interpreted as an additive effect of high levels of CaMKK activity and loss of rad24 [31]. Human [53] and Xenopus [54] 14-3-3 binds to the mitotic inhibitor Wee1 to negatively regulate the cell cycle, through increasing Wee1 half-life, protein levels and kinase activity [53]. Fission yeast 14-3-3 may act similarly on Wee1, where increases in 14-3-3 contribute to G2/M delay. Ectopically augmented levels of the mitotic activator Ssp1 presumably titrates out some of the excess Rad24, reducing its effect on other binding partners. Ssp1 overexpression by itself causes mitotic advance indicating a codominant relationship upon overexpression where Ssp1 and Rad24 work independently in an opposing manner. Fission yeast Cdk1 Y15 dephosphorylation by Cdc25 (and Pyp3) phosphatase and phosphorylation by Wee1 (and Mik1) kinase provide positive and negative

**Figure 6.** Ssp1-GFP physically interacts with Rad24-2HA-His6 and Rad25-His6.

*Panel (a)* Ssp1-GFP localization in rad24'' and rad25'' cells. Cells (YEA, 30°C) were incubated with pre-warmed YEA, 0.6 M KCl. Fluorescence images were taken prior to and at 15 min after the addition of KCl. (b–f) Ssp1-GFP interacts with Rad24-2HA-His6 and Rad25-His6 in vivo. Cells co-expressing Ssp1-GFP and Rad24-2HA-His6 protein (b), Ssp1-GFP and Rad25-His6 protein (c), Ssp1-GFP with both Rad24-2HA-His6 and Rad25-His6 or Ssp1-GFP and Rad25-His6 (rad24'' or rad24'') proteins (d) were grown at 30°C in YEA. Aliquots of whole cell lysates used for the immunoprecipitations were loaded (5–15 μg total protein) and Ssp1-GFP, Rad24-2HA-His6 and Rad25-His6 fusion proteins were directly detected. (e) Rad25-His6 interacts with Ssp1-GFP in the absence of rad24. (f) Reduced stability of Ssp1-GFP in the absence of rad24. WB, western blot with antibody.
regulation of cell-cycle progression, respectively [34,55–62]. Wee1 is negatively regulated by phosphorylation through the mitotic activators Cdr1 and Cdr2 kinases [58,63–68]. Ssp1 CaMKK involvement in mitotic control is demonstrated by the reduction of the restrictive temperature of cdc25-22 by the loss of ssp1. The negative additive effect on cell-cycle progression suggests that loss of Ssp1 further inhibits Cdk1, presumably via net increase in Y15 phosphorylation probably through impacting Wee1 kinase activity. Together, these findings support a role for CaMKKs in cell-cycle regulation.

4.2. CaMKK Ssp1 does not co-localize with actin patches at the cell membrane

At native expression levels, cytoplasmic Ssp1-GFP accumulates at the cell membrane after perturbation by either 0.6–1.2 M KCl or sorbitol [23]; however ssp1 is required for growth in the presence of KCl but not sorbitol. Similarly, although MAPK Pmk1 is activated by 1.2 M sorbitol, pmk1- cells are not sensitive to this hyperosmotic stress [69]. A small pool of Sep1-GFP accumulates at the cell membrane, while the majority of the protein remains cytoplasmic. This does not support a previous model suggesting that Ssp1 directly localizes actin patches at the cell membrane to support areas of new growth [23]. We show that following hyperosmotic stress GFP-Ssp1 forms a band, while actin patches do not follow this pattern; Arp3C-YFP and Sep1-CFP do not co-localize. The importance of the compartmentalization of Ssp1 at the membrane is not clear. The S. pombe cell wall is most vulnerable to rupture at the extensile tips and is sturdier in the cylindrical portion of the cell [70]. Ssp1 accumulates in areas corresponding to fission scars, which are less vulnerable to damage than the extensile tips [70]. Accumulation at the cell membrane following hyperosmotic stress is transient; however, Ssp1 is required for long-term cell survival under hyperosmotic conditions.

4.3. The role of 14-3-3 binding to CaMKK

At least some cytoplasmic Ssp1 is bound to Rad24 and Rad25 in unperturbed cells. After applying hyperosmotic stress, Sep1 is released from 14-3-3 and Ssp1 accumulates at the cell membrane. Our data also show that Rad24 and Rad25 are dispensable for the translocation of Sep1-GFP to the cell membrane. This response can be repressed in cells overexpressing Rad24 even when there is an excess of Ssp1-GFP.

14-3-3 proteins commonly act as cytoplasmic anchors, providing negative regulation of proteins through sequestration. In Drosophila melanogaster, 14-3-3 binds phosphorylated β-catenin antagonist Chibby (Cby), promoting cytoplasmic sequestration of the β-catenin–14-3-3–Cby complex [71]. In mammalian systems cytoplasmic Bax, a Bcl-2-related protein required for membrane integrity, is released and translocates to the mitochondria after stress stimuli, where it induces cytochrome c release. 14-3-3-bound Bax remains anchored in the cytoplasm [72]. Upon activation, JNK phosphorylates 14-3-3, and constitutively binds to 14-3-3 proteins which are thought to inhibit activation of plasma membrane-anchored Ras by sequestering its activator Raf-1 in the cytoplasm. This mechanism prevents cascade activation in resting cells [74]. The majority of Ras superfamily G proteins are kept in inactive (GDP-bound) or active (GTP-bound) forms by guanine nucleotide exchange factors and GTPase-activating proteins, respectively. The atypical Rho GTPases Rnd1/2/3 are cell morphology regulators and constitutively bind GTP [75,76]. After phosphorylation by
Rock1 kinase or protein kinase Ca (PKCa) [77,78], Rnd3 binds 14-3-3 and translocates from the plasma membrane to the cytoplasm, its function inhibited via sequestration from its site of action [79]. In fission yeast, 14-3-3 proteins also regulate the localization of many proteins. For example, they associate with the primarily cytoplasmic Byr2, preventing binding to Ras1/GTP at the cell membrane during vegetative growth. Loss of 14-3-3 expedites Byr2 translocation [80]. Our data suggest that in fission yeast, 14-3-3 proteins may play a role in the negative regulation of Ssp1 translocation to the cell membrane.

Mammalian 14-3-3 isoforms require only their N-termini to dimerize. Dimerization greatly increases their thermostability and single recombinant isoforms form homodimers even if they function as heterodimers in vivo [81,82]. Particular isoforms in 14-3-3 heterodimers allow the interaction of proteins by bringing the two binding partners closer together ([82–84]; reviewed in [85–87]). The minor isoform Rad25, which associates with Ssp1, forms homodimers and binds Ssp1, at least in the absence of Rad24. A reduction in Rad24 binding to Ssp1 after osmotic stress could occur owing to a decrease in Rad24–Rad24 homodimer and/or Rad24–Rad25 heterodimer binding. Conversely, following hyperosmotic stress, Rad25–Rad25 homodimers remain bound to Ssp1-GFP. This suggests an interesting and distinct role for the minor 14-3-3 isoform in CaMKK regulation. Future studies will confirm whether the Rad24-bound pool is preferentially located in any particular part of the cell.

4.4. Increased stress sensitivity in cells overexpressing rad24 is alleviated by co-overexpression of CaMKK

Rad24 is a negative regulator of mitosis after DNA damage [88,89]. Overexpression of rad24 increases long-term sensitivity to stressors such as KCl and low pH, but sensitivity to high temperature is relieved to some extent by co-overexpression of ssp1. Association with 14-3-3 proteins inhibits CaMKK activity in mammalian systems [32,90], thus if Rad24 binding inhibits Ssp1 activity then augmenting levels of 14-3-3 would diminish Ssp1-mediated stress response. Ssp1-FLAG binds 14-3-3, but it is unclear whether this association affects CaMKK activity [19]. 14-3-3 proteins are involved in many pathways [91,92] and substantially increasing the Rad24 pool causes a complex response in terms of stress sensitivity. Both deletion and overexpression of 14-3-3 BMH1 increase chronological lifespan in nutrient-stressed budding yeast. Cells overexpressing BMH1 survive longer in the absence of additional stressors, probably because an increase in phosphorylated BMH1 S238 decreases the stress response required for longevity [93].

4.5. A role for 14-3-3 in CaMKK turnover

The absence of Rad24 increases Ssp1 turnover, suggesting a role for Rad24 in maintaining Ssp1 protein stability. rad24− cells do not display hypersensitivity to conditions where ssp1− cells are unable to proliferate, indicating that Ssp1 protein levels are maintained at sufficient levels. 14-3-3 proteins are involved in protein stabilization both directly and indirectly in other systems, for example by blocking access of ubiquitin ligases (reviewed in [85,86]). In mammalian systems, association with 14-3-3 prevents Wee1 degradation by masking a degradation motif required for normal Wee1 turnover [94]. Budding yeast 14-3-3 homologues BMH1, BMH2 and ACM1 form a stable complex with the APC/C activator CDH1/CDC20, keeping it inactivated by acting as an APC pseudosubstrate [95,96].

Further studies will determine whether the interaction of 14-3-3 and Ssp1 has a direct or indirect effect on CaMKK catalytic function and whether 14-3-3 proteins play a role in the negative regulation of Ssp1 translocation to the cell membrane after stress.

5. Material and methods

5.1. Plasmid construction and chromosomal integration

All DNA amplification was performed by PCR with Expand High Fidelity Taq polymerase (Roche). T4 ligase and restriction enzymes were used from Promega. For primers, see table 2.

| ssp1-GFPint and ssp1-CFPint: the ssp1 gene (+1000 bp upstream sequence) was amplified from S. pombe genomic DNA [97] (primers Ssp1intfor and Ssp1intrev) adding PstI and SalI restriction sites. The nmt1 promoter was excised from the pREP1-GFP and pREP1-CFP vectors with PstI and SalI [41,98,99] and the ssp1 +1000 bp fragment was inserted, generating ssp1-GFPint and ssp1-CFPint. The plasmids were integrated into ssp1::ura4− leu1-32 ura4-D18 and stable integrants were identified and tested by out-crossing to a ura4-D18 leu1-32 strain. Strains were tested for their ability to rescue the ssp1− phenotype, and normal subcellular localization was confirmed (see Results section). |
| pREP1-rad24-His6, pREP2-rad24-His6 and pREP1-rad24-GFP, the rad24 ORF was amplified with primers rad24OPrev and rad24OPrev, adding a C-terminal His tag, NdeI and SalI restriction sites or the primers SFrnd24-f and SFrnd24-r, adding NdeI and SalI restriction sites, respectively. The fragments were ligated into pREP1, pREP2 [41,98] and pREP1-GFP plasmids [99] forming pREP1-rad24-His6, pREP1-rad24-GFP and pREP1-rad24-GFP. Plasmids rescued rad24::ura4−. A single copy of pREP2-rad24-His6 was integrated into a leu1-32 ura4-D18 strain. |
| Integration of nmt1::GFP-ssp1: a single copy of the plasmid pLR2-22, containing GFP-ssp1 under control of the nmt1 promoter (pLR2-22) [23] was integrated into leu1-32 ura4-D18. |

5.2. Targeted replacement of LEU2 and ura4 with kanMX6 in ssp1-GFPint and rad25::ura4]

A kanMX6 cassette with 80 bp sequence homology to LEU2 at the 5′ and 3′ ends was generated by PCR amplification using primers KanShortForwTM and KanShortRevTM and the pGEM-T (Promega) vector containing kanMX6. The kanMX6 cassette was transformed into ssp1-GFPint and plated on YEA (+0.1 mg ml−1 G418; Gibco) [100,101]. A kanMX6 cassette was amplified with KmrFW-17 and KmrRV-17 primers having homology to URA4 at the 5′ and 3′ ends and transformed into rad25::ura4− ura4-D18, producing rad25::kanMX6 ura4-D18.

5.3. Protein lysates

Lysates were prepared at 4°C unless otherwise indicated. Mid-logarithmic growth phase cells were harvested by centrifugation (5 min, 1876 g), washed in ice-cold stop buffer [97], collected again by pulse centrifugation (13 051 g) and
frozen on dry ice. Lysis by mechanical disruption with glass beads (0.5 mm, BioSpec) and a bead beater (MiniBeadBeater-8 Cell Disruptor, BioSpec) in lysis buffer [102] or modified SUME buffer (1% SDS, 8 M urea, 10 mM MOPS pH 6.8, 10 mM EDTA, 50 mM NaF, 1 mM NaVO₄; [103]) supplemented with complete protease inhibitor (EDTA-free, Roche) was alternated with rest periods in an ice-water slurry (0 °C). Lysates were cleared by centrifugation (13 051 g) and protein concentration determined by Bio-Rad protein assay. Protein extracts were boiled in Laemmli buffer (200 mM Tris–HCl pH 6.8, 8% SDS, 40% glycerol, 3.34% (v/v) 2-mercaptoethanol, 0.01% bromophenol blue).

5.4. KCl treatment

Mid-logarithmic growth phase cells (1000–3000 ml, YEA, 30 °C) were treated for 15 min with warm (30 °C) YEA (control) or YEA + KCl to 0.6 M. Cells were chilled to 0 °C within 1 min by the addition of frozen, crushed YEA or YEA + 0.6 M KCl and immediate immersion of flask into an ice-water/ethanol slurry.

5.5. Immunoprecipitation

Lysate preparation and immunoprecipitation were performed at 4 °C. Cell cultures (YEA, 30 °C) were harvested by centrifugation (9927.3g) and washed once in 10 ml ice-cold HB buffer [97] (pH 7.4; containing 2 mM DTT and Roche complete protease inhibitor (EDTA-free), washed again and followed by mechanical disruption in ice-cold HB buffer. Lysates were centrifuged (35 440g: 40 min) and incubated with 100 µl bed-volume Sepharose G beads (4 Fast Flow, GE Healthcare) for 1 h to remove non-specific binding proteins. Lysates were incubated overnight with 15 µl rabbit anti-GFP polyclonal serum (Invitrogen) or 15 µl mouse anti-His₅ antibody (Roche) on a rotator. Control lysates did not contain antibodies. Lysates were incubated with 100 µl bed-volume Protein G Sepharose beads for 2 h. Beads were washed extensively with HB buffer and protein complexes eluted with 0.2 M glycine (pH 2.2) for 15 min and neutralized by addition of saturated Tris (pH 10). Supernatants were boiled with Laemmli buffer and analysed by immunoblotting.

5.6. Immunoblotting

Extracts and immunoprecipitates were subjected to SDS-PAGE and transferred to a polyvinylidene difluoride (PVDF) membrane (Perkin Elmer). Immunoprecipitated Ssp1-GFP protein was detected with monoclonal anti-GFP antibody (1: 1000) (Roche). Co-immunoprecipitated Ssp1-GFP protein was detected with polyclonal anti-GFP serum (1: 1000) (Invitrogen). Immunoprecipitated and co-immunoprecipitated Rad24-2HA-His₆ and Rad25-His₆ proteins were detected with polyclonal anti-GFP serum (1: 1000) (Invitrogen). Co-immunoprecipitated Ssp1-GFP protein was detected with monoclonal anti-GFP antibody (1:1000) (Roche). Bands were visualized with goat anti-mouse or goat anti-rabbit HRP-conjugated secondary antibody (1:2000) (Santa Cruz Biotechnology) and luminol-based ECL reagent (Perkin Elmer).

5.7. Reprobing of polyvinylidene difluoride membranes

PVDF membranes were stripped as described [104] and reprobed with polyclonal anti-PSTAIRE antibody (1:500) (Upstate Biotechnology).

5.8. Phosphatase treatments

Mid-logarithmic phase cells were treated with 30 °C YEA or YEA + 0.6 M KCl for 15 min. To preserve phosphorylation state, cells were collected on microfibre filters (934-AH; Whatman) and washed with 5 ml ice-cold stop buffer [97] lysed in lysis buffer [97] (buffers contained 15 mM pNPP and 60 mM β-glycerophosphate). Cells collected by centrifugation were washed (150 mM NaCl, 1 mM EDTA, 1 mM PMSF) and lysed in phosphatase-inhibitor-free lysis buffer (modified from [97]) or washed and lysed with phosphatase-inhibitor-enriched buffers. Phosphatase-inhibitor-free protein (5 µg) was treated with 80 units of Lambda Protein Phosphatase (NEB) for 30 min at 30 °C. Mock treatments did not contain phosphatase.

5.9. Microscopy

Images were captured by a high performance CCD (Cooke SensiCam) camera on a Leitz DMRB fluorescence microscope.
or a high performance CCD Hamamatsu Orca-ER camera on a Zeiss AxioImager.Z1 fluorescence microscope. Slidebook image analysis software (Intelligent Image Innovations) was used to perform cell measurements and to analyse Z-stacks.

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