Two protein tyrosine phosphatases, Ptp2 and Ptp3, modulate the subcellular localization of the Hog1 MAP kinase in yeast

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The MAP kinase Hog1 transiently accumulates in the nucleus upon activation. Although Hog1 nuclear export correlates with its dephosphorylation, we find that dephosphorylation is not necessary for export. Unexpectedly, a strain lacking the nuclear protein tyrosine phosphatase, Ptp2, showed decreased Hog1 nuclear retention, while a strain lacking the cytoplasmic Ptp3 showed prolonged Hog1 nuclear accumulation, consistent with Ptp2 being a nuclear tether for Hog1 and Ptp3 being a cytoplasmic anchor. In support of this result *PTP2* overexpression sequestered Hog1 in the nucleus while *PTP3* overexpression restricted Hog1 to the cytoplasm. Thus, Ptp2 and Ptp3 regulate Hog1 localization by binding Hog1.

[**Key Words:** Hog1; PTP; dephosphorylation; nuclear and cytoplasmic anchoring]

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Signal is transmitted to the nucleus through MAP kinase (MAPK) pathways via nuclear accumulation of MAPK (Chen et al. 1992; Gonzalez et al. 1993; Lenormand et al. 1993, 1998; Brunet et al. 1999). For example, the stress-activated MAPKs Hog1 in *Saccharomyces cerevisiae* and Spc1 in *Schizosaccharomyces pombe* transiently accumulate in the nucleus in response to osmotic stress (Ferrigno et al. 1998; Gaits et al. 1998; Reiser et al. 1999), and the vertebrate extracellular signal-regulated kinase (ERK) accumulates in response to growth factors (Traverse et al. 1992; Khokhlatchev et al. 1998; Lenormand et al. 1998; York et al. 1998). One requirement for nuclear accumulation is phosphorylation of the activation loop. Mutation of the activation loop Thr and/or Tyr residues prevents signal-induced nuclear accumulation of Hog1 and Spc1 (Ferrigno et al. 1998; Gaits et al. 1998; Reiser et al. 1999) as does mutation of the ERK activation loop (Lenormand et al. 1993; Khokhlatchev et al. 1998). Although phosphorylation of the activation loop is required for signal-induced nuclear accumulation, MAPK activity is not required. Hog1 and ERK mutated at the ATP-binding loop accumulate in the nucleus in response to signal (Gonzalez et al. 1993; Ferrigno et al. 1998; Khokhlatchev et al. 1998; Reiser et al. 1999).

Regulation of MAPK export from the nucleus is not as well understood. Nuclear export proteins, including the importin β homolog, Xpo1, and the small GTP-binding protein, Ran-Gsp1, have been implicated in Hog1 export

(Ferrigno et al. 1998); however, it is not clear whether the phosphorylation state of Hog1 regulates export. This event seems possible because dephosphorylation of the activation loop phosphotyrosine occurs simultaneously with export (Ferrigno et al. 1998; Reiser et al. 1999). Our results indicate, however, that Hog1-pY dephosphorylation by the nuclear-localized protein tyrosine phosphatase Ptp2 or the cytoplasmic protein tyrosine phosphatase Ptp3, or Hog1-pThr dephosphorylation by a type-2C Ser/Thr phosphatase, Ptc1, does not regulate its export from the nucleus. Instead, Ptp2 modulates Hog1 localization by acting as a nuclear anchor for Hog1 while Ptp3 acts as a cytoplasmic tether.

Results

Dephosphorylation of Hog1 is not required for nuclear export during adaptation

To examine the effect of Hog1 dephosphorylation on its localization, we produced a strain expressing Hog1-GFP from a low-copy plasmid in a *hog1 Δ* background. To quantify the features of this process, we determined the percentage of cells having greater nuclear Hog1 relative to cytoplasmic Hog1 before and during exposure to osmotic stress. As described previously, Hog1 transiently accumulated in the nucleus in response to stress and began redistribution to the cytoplasm between 10–20 min (Figs. 1 and 2) when the level of Hog1-pY began to decrease (Fig. 3).

To test whether protein tyrosine phosphatase-depen-

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Figure 1. Redistribution of nuclear Hog1 to the cytoplasm does not require dephosphorylation of Hog1-pY. The subcellular localization of Hog1-GFP or Hog1-D310N-GFP in response to osmotic stress was examined in wild type or strains lacking *PTP2*, *PTP3*, or both *PTP2* and *PTP3*. Exponentially growing cells were exposed to 0.4 M NaCl or 0.9 M NaCl for the indicated times. GFP and DAPI-stained nuclei were visualized by fluorescence microscopy.

dent dephosphorylation of Hog1-pY is required for its export from the nucleus, we examined Hog1 localization in strains lacking protein tyrosine phosphatases. Ptp2 and Ptp3 inactivate Hog1 by dephosphorylating Hog1-pY, with Ptp2 being the more potent negative regulator (Jacoby et al. 1997; Wurgler-Murphy et al. 1997). Ptp2 is predominantly nuclear while Ptp3 is cytoplasmic and appears to be excluded from the nucleus (Mattison et al. 1999). Therefore, deletion of *PTP2* should delay Hog1 export if its dephosphorylation is necessary for export. Although the *ptp2Δ* strain poorly dephosphorylated Hog1-pY (Fig. 3), Hog1 redistribution to the cytoplasm was not delayed compared with wild type (Figs. 1 and 2A). For example, in the *ptp2Δ* mutant, the level of Hog1-pY remained high up to 40 min of osmotic stress; however, it was exported from the nucleus after 20 min (Figs. 1 and 2A). To examine further the effect of sustained Tyr phosphorylation, we examined Hog1 localization in the *ptp2Δ ptp3Δ* strain, which had an even greater

defect dephosphorylating Hog1-pY (Fig. 3). Hog1 nuclear accumulation and redistribution to the cytoplasm in the *ptp2Δ ptp3Δ* strain was similar to wild type (Figs. 1 and 2A). These results indicate that Hog1-pY dephosphorylation is not required for export of Hog1 from the nucleus.

Because Ptp2 and Ptp3 negatively regulate at least two other MAPK pathways (Zhan et al. 1997; Mattison et al. 1999), deleting PTPs may indirectly affect Hog1 localization by affecting other pathways. Therefore, we produced a mutant, Hog1-D310N, analogous to the sevenmaker mutant in the *Drosophila* rolled MAPK (Biggs et al. 1994; Bott et al. 1994; Brunner et al. 1994), that resists dephosphorylation in vivo (Fig. 3). Although Hog1-D310N was poorly dephosphorylated during adaptation, its subcellular localization was not appreciably altered from wild type (Figs. 1 and 2B), supporting further the idea that Hog1-pY dephosphorylation is not required for its export from the nucleus.

Hog1 activation and nuclear accumulation require phosphorylation of both Thr174 and Tyr176 in the activation loop (Schuller et al. 1994; Ferrigno et al. 1998; Reiser et al. 1999). To test whether Hog1-pThr174 dephosphorylation could be important for its export, we examined Hog1 localization in a strain lacking *PTC1*, encoding a type-2C Ser/Thr phosphatase. *PTC1* was identified as a negative regulator of the HOG pathway by genetic means (Maeda et al. 1993, 1994), and more recently, we showed that Ptc1 inactivates Hog1 in vivo and in vitro by specifically dephosphorylating pThr174 at the activation loop (J. Warmka and I. Ota, unpubl.). In the *ptc1Δ* strain, Hog1 showed an increased basal level of activity, and was inactivated poorly as judged by Hog1 kinase assays (J. Warmka and I. Ota, unpubl.); however, it accumulated in the nucleus and redistributed to the cytoplasm in a manner indistinguishable from wild type (Fig. 2C). These results indicate that dephosphorylation of neither Thr174 nor Tyr176 is necessary for export.

To test whether inhibiting dephosphorylation of both pThr174 and pTyr176 would prevent export, we examined Hog1 localization in a *ptc1Δ ptp2Δ* strain. This strain constitutively activates Hog1 and, therefore, exhibits a strong growth defect (Maeda et al. 1993; Jacoby et al. 1997). Despite this defect, the kinetics of Hog1 nuclear accumulation and redistribution to the cytoplasm were not significantly altered from wild type (Fig. 2C). These results suggest strongly that dephosphorylation of Hog1 at either p-Thr174 or p-Tyr176, and consequent inactivation of kinase activity is not required for redistribution of Hog1 to the cytoplasm. These results also suggest that factors other than phosphorylation are required for nuclear accumulation of Hog1, because active Hog1, present in the *ptc1Δ* and *ptc1Δ ptp2Δ* strains prior to stress, was not found accumulated in the nucleus (Fig. 2C).

Ptp2 and Ptp3 act as nuclear and cytoplasmic anchors for Hog1, respectively

Although the strain lacking nuclear Ptp2 did not slow

Figure 2. Hog1 dephosphorylation is not required for redistribution to the cytoplasm. Cultures expressing Hog1-GFP were exposed to 0.4 M NaCl, unless otherwise noted, and the percentage of cells with greater nuclear relative to cytoplasmic accumulation determined. Approximately 100 cells were counted for each time point, and the average from at least two such experiments is shown. (A) Hog1 export from the nucleus does not require Ptp2 or Ptp3. (●) Wild type; (○) *ptp2Δ*; (■) *ptp3Δ*; (▽) *ptp2Δ ptp3Δ*. (B) Hog1-D310N export is similar to wild type. (●) Wild type; (□) Hog1-D310N. (C) Hog1 export does not require dephosphorylation by Ptc1 or Ptp2. (●) Wild-type; (□) *ptc1Δ*; (◆) *ptc1Δ ptp2Δ*. (D) Strains lacking *PTP2* accumulate less Hog1 in the nucleus whereas strains lacking *PTP3* accumulate more Hog1 in the nucleus than wild type. Cells were exposed to 0.9 M NaCl. Symbols are as in A. (E) PTP catalytic activity is not required for anchoring Hog1. Wild-type or PTP-null strains expressing PTP catalytic mutants (Jacoby et al. 1997) were treated with 0.9 M NaCl. (●) Wild type; (○) *ptp2-C666A* (CEN); (▽) *ptp2-C666A* (2 μm); (■) *ptp3-C804A* (CEN); (□) *ptp3-C804A* (2 μm). (F) Hog1-K52M redistribution to the cytoplasm requires wild-type Hog1. *HOG1* or *hog1Δ* cells carrying pHOG1-K52M-GFP were exposed to 0.9 M NaCl for the indicated times. (■) pHOG1-K52M-GFP expressed in wild type; (○) pHOG1-K52M-GFP expressed in *hog1Δ*. (G) Hog1 kinase activity is not required to activate PTPs to redistribute Hog1-K52M to the cytoplasm. Exponentially growing wild type (●) or *ptp2Δ* (○) carrying pHOG1-K52M-GFP were exposed to 0.9 M NaCl for the indicated times.

Hog1 export from the nucleus, a slight decrease in the level of Hog1 nuclear accumulation was seen before and during osmotic stress (Fig. 2A). Conversely, the strain lacking cytoplasmic Ptp3 showed a slight increase in nuclear Hog1 (Fig. 2A). These results are consistent with Ptp2 acting as a nuclear tether for Hog1, and Ptp3 being a cytoplasmic anchor for Hog1. For example, the *ptp2Δ* strain should accumulate less Hog1 in the nucleus than wild type if Ptp2 tethers Hog1 in the nucleus. Ptp2 and Ptp3 could function in this manner, as catalytically inactive mutants of Ptp2 and Ptp3 coprecipitate with Hog1 in yeast lysates (Jacoby et al. 1997; Wurgler-Murphy et al. 1997).

To test this idea further, we attempted to amplify the difference in Hog1-GFP localization between wild-type and PTP deletion strains by exposing cells to higher osmotic stress. Exposure to 0.9 M NaCl increased the duration of Hog1 in the nucleus in both wild-type and *ptpΔ* strains (Fig. 2, cf. A and D). Prolonged nuclear accumulation of Hog1 at 0.9 M NaCl is due to at least two factors. First, the level of Hog1-pY remained high for up to 4 hr at 0.9 M NaCl compared with 15 min at 0.4 M NaCl in wild type (Fig. 3), suggesting that upstream components of the HOG pathway remain activated for a longer period

of time at 0.9 M NaCl. Second, more Hog1 was activated at 0.9 M NaCl, as Hog1 kinase activity was ~1.5-fold higher at 0.9 M NaCl compared with 0.4 M NaCl (data not

Figure 3. The level of Hog1-pY increases in *ptpΔ* strains and *HOG1* mutants. Hog1-pY in *hog1Δ* strains wild type for PTPs or lacking PTPs was examined in exponentially growing cells carrying pHOG1-GFP (rows 1-4). The level of Hog1-D310N-pY was also examined in a *hog1Δ* strain. Cells were exposed to 0.4 or 0.9 M NaCl, harvested at the indicated times, and lysates analyzed by SDS-PAGE and immunoblotting with anti-phosphotyrosine antibody.

shown). Because dephosphorylation of Hog1 is not necessary for its export from the nucleus, we believe that the sustained nuclear accumulation of Hog1 at 0.9 M NaCl could be due to re-import of phosphorylated Hog1. Under these conditions, the *ptp2Δ* strain retained significantly less Hog1 in the nucleus than wild type, while the *ptp3Δ* strain showed increased nuclear retention of Hog1 (Fig. 2D).

To test whether Ptp2 sequestration of Hog1 in the nucleus and Ptp3 tethering of Hog1 to the cytoplasm was a result of PTPs binding Hog1, or their catalytic activity, Hog1 localization was examined in *ptp*-null strains expressing catalytically inactive PTPs. Ptp2C666A shifted Hog1 to the nucleus, while Ptp3C804A shifted Hog1 to the cytoplasm (Fig. 2E). Expression of these mutants from multicopy plasmids exacerbated these effects (Fig. 2E). These results support strongly the idea that Ptp2 is a nuclear anchor whereas Ptp3 is a cytoplasmic tether for Hog1.

If PTPs function as anchors for Hog1, then their overexpression from the strong *GAL* promoter might sequester Hog1 in the absence of signal. Indeed, *PTP2* overexpression segregated the majority of Hog1 to the nucleus

Figure 4. Overexpression of PTPs tethers Hog1 in different subcellular compartments. The subcellular localization of wild-type and mutant Hog1 proteins fused to GFP was examined in *hog1Δ* strains overexpressing *PTP2* or *PTP3* or their mutant counterparts, *ptp2-C666S* and *ptp3-C804A*. The Hog1 plasmids were pHOG1-GFP, pHOG-Y176F-GFP, pHOG-K52M-GFP, and pHOG1-D310N-GFP. The PTP-overexpressing plasmids were pKT-PTP2, pKT-PTP3, pGST-PTP2-C666S, and pGST-PTP3-C804A. GFP and DAPI-stained nuclei were visualized by fluorescence microscopy.

Figure 5. PTPs physically associate with Hog1. GST fusions to wild-type and mutant PTPs (Ptp2-C666S and Ptp3-C804A), were coexpressed with wild type and mutant Hog1-GFP fusions in yeast. GST fusion proteins were isolated using glutathione-Sepharose and associated Hog1 examined by SDS-PAGE and immunoblotting with anti-GST and anti-GFP antibodies as indicated.

while *PTP3* overexpression sequestered Hog1 in the cytoplasm (Fig. 4). Colocalization was mediated by complex formation as wild-type PTPs coprecipitated with Hog1 (Fig. 5). Next, we examined the requirements for PTP-induced Hog1 compartmentalization. First, PTP catalytic activity was not required because overexpression of Ptp2-C666S sequestered Hog1 in the nucleus, whereas overexpression of Ptp3-C804A, localized Hog1 to the cytoplasm (Fig. 4). Second, neither Hog1 Tyr phosphorylation nor Hog1 activity was required for PTP tethering, as Hog1-Y176F and Hog1-K52M were concentrated in the nucleus by *PTP2* overexpression (Fig. 4) and sequestered in the cytoplasm by *PTP3* overexpression (Fig. 4). Colocalization was due to complex formation, as mutant PTP proteins coprecipitated with wild-type Hog1 as described previously (Fig. 5; Jacoby et al. 1997; Wurgler-Murphy et al. 1997) and mutant Hog1 coprecipitated with wild-type PTPs (Fig. 5). One mutant protein, Hog1-D310N, was not sequestered by overexpression of PTPs (Fig. 4) and bound poorly to PTPs (Fig. 5). Therefore, PTPs regulate Hog1 localization by formation of tight complexes.

Activation of nuclear Ptp2 is not required for export of kinase-inactive Hog1

One factor important for export of Hog1 from the nucleus is its kinase activity. A kinase-inactive mutant, Hog1-K52M, is retained in the nucleus following exposure to 0.4 M NaCl (Ferrigno et al. 1998; Reiser et al. 1999) and 0.9 M NaCl (Fig. 2F). Because osmotic stress increases the level of *PTP2* and *PTP3* mRNAs in a Hog1-dependent manner (Jacoby et al. 1997; Wurgler-Murphy et al. 1997), it has been suggested that Hog1-K52M is retained in the nucleus because of its inability to activate PTPs. In support of this, Hog1-K52M-pY was de-

phosphorylated poorly after exposure to 0.4 M NaCl (Wurgler-Murphy et al. 1997; Ferrigno et al. 1998) and 0.9 M NaCl (data not shown). Furthermore, coexpression of wild-type *HOG1* with Hog1-K52M-GFP, allows Hog1-K52M-GFP to be dephosphorylated and exported from the nucleus at 0.4 M NaCl (Ferrigno et al. 1998; Reiser et al. 1999) and 0.9 M NaCl (Fig. 2F). These results suggest that Hog1-K52M is retained in the nucleus because of its inability to be dephosphorylated by PTPs. Our results with wild-type Hog1, however, predicted that PTP dephosphorylation of Hog1-K52M is not required for export. Therefore, we examined Hog1-K52M-GFP export in a *HOG1 PTP2* strain, and a *HOG1 ptp2Δ* strain. If Hog1-K52M-GFP is rapidly exported in the *HOG1* strain because Ptp2 is activated, then Hog1-K52M export should be retarded when *PTP2* is deleted. In the *HOG1 ptp2Δ* strain, Hog1-K52M-GFP did not accumulate in the nucleus as well as wild type; however, it redistributed to the cytoplasm in a manner similar to the *HOG1 PTP2* strain (Fig. 2G). Thus, the rapid redistribution of Hog1-K52M to the cytoplasm in the *HOG1* strain is likely due to factors other than Ptp2-mediated dephosphorylation.

Discussion

The subcellular localization of MAPKs is regulated by their phosphorylation. MAPK nuclear import requires phosphorylation of the activation loop Thr and Tyr residues. Although dephosphorylation of Hog1 is correlated with its export from the nucleus, we show that its dephosphorylation at pTyr or pThr in the activation loop is not required for export. Dephosphorylation may not regulate the export of other MAPKs. For example, the protein tyrosine phosphatases that inactivate the *S. pombe* stress-activated MAPK, Spc1, are found in the cytoplasm, suggesting that Spc1-pY dephosphorylation is not necessary for export (Gaits and Russell 1999).

One factor critical for Hog1 export from the nucleus is its kinase activity (Ferrigno et al. 1998; Reiser et al. 1999). Here, we show that dephosphorylation of kinase-inactive Hog1 is not necessary for export, suggesting that kinase activity is required to up-regulate or activate proteins other than phosphatases to allow export. For example, export of the vertebrate stress-activated MAPK, p38, from the nucleus is dependent on phosphorylation of its substrate, MAPKAP kinase-2 (Ben-Levy et al. 1998). An alternative explanation for the extended nuclear localization of Hog1-K52M, however, is that it binds nuclear substrates with greater affinity, and thereby prolongs nuclear accumulation. For example, similar mutations in MEK and MAPK bind their substrates with higher affinity (Zhou et al. 1995; Han et al. 1997).

A second factor that regulates MAPK localization is interaction with proteins in the cytoplasm and the nucleus. For example, MEK is proposed to be a cytoplasmic anchor for MAPK, tethering it in the cytoplasm prior to activation and releasing it upon activation to accumulate in the nucleus (Fukuda et al. 1997). In yeast stress-

response pathways, MEK may not function in this manner. Deletion of MEK does not lead to nuclear accumulation of Hog1 (Reiser et al. 1999; C. Mattison and I. Ota, unpubl.) or the *S. pombe* Spc1 (Gaits et al. 1998). Nevertheless, cytoplasmic anchoring is important for Hog1 localization, as overexpression of the activation loop mutant, Hog1-T174A,Y176A, leads to nuclear accumulation (C. Mattison and I. Ota, unpubl.). Because Hog1 is more abundant than Pbs2 (C. Mattison and I. Ota, unpubl.), other cytoplasmic proteins are likely involved in tethering Hog1.

In this work, we identify Ptp3 as one such cytoplasmic tether for Hog1. Deletion of *PTP3* elevated the level of Hog1 in the nucleus prior to and following osmotic stress. We find that Ptp3 is predominantly cytoplasmic and that its localization does not change in response to osmotic stress, consistent with it being a cytoplasmic tether (Mattison et al. 1999). Because there is less Ptp3 protein than Hog1 (C. Mattison and I. Ota, unpubl.), other proteins are also likely to anchor Hog1 in the cytoplasm. Cytoplasmic phosphatases in other MAPK pathways may also function as tethers. For example, overexpression of the catalytically inactive dual-specificity phosphatase, MKP-3, or the protein tyrosine phosphatase, PTP-SL, has been shown to constrain ERK to the cytoplasm and prevent signaling (Brunet et al. 1999; Zuniga et al.).

Nuclear retention factors are also important in regulating MAPK signaling. In *S. pombe*, nuclear accumulation of Spc1 requires the presence of its nuclear-localized substrate, the transcriptional activator, Atf1 (Gaits et al. 1998; Watanabe and Yanamoto 1996). In this work, we show that a protein phosphatase, Ptp2, can also affect the localization of Hog1. Consistent with Ptp2 being a nuclear tether, it is concentrated in the nucleus, and its localization does not change upon osmotic stress (Mattison et al. 1999). Because deletion of *PTP2* reduced but did not inhibit Hog1 nuclear accumulation, other nuclear proteins may also function as anchors for Hog1. For example, deletion of *MSN2* and *MSN4*, general stress transcription factors, also inhibit maximal nuclear accumulation of Hog1 (Reiser et al. 1999).

Therefore, Ptp2 and Ptp3 have more than one role in regulating the HOG pathway. One is to inactivate Hog1 through dephosphorylation, while the second is to modulate Hog1 subcellular localization through binding interactions. These functions may be separated into different domains in the PTPs, which have a noncatalytic amino-terminal domain and a catalytic carboxy-terminal domain (Ota and Varshavsky 1992; Jacoby et al. 1997). PTP-Hog1 compartmentalization does not require PTP activity, Hog1-Tyr phosphorylation, or Hog1 kinase activity. Therefore, PTP-Hog1 binding interactions could occur at sites separate from the PTP catalytic domain and the Hog1 activation loop. Indeed, the isolated amino-terminal domain of PTPs bind Hog1 (C. Mattison, S. Spencer, and I. Ota, unpubl.), and the Hog1 mutation, Hog1-D310N, which shows reduced binding to PTPs, is found on the face opposite Tyr176, suggesting that re-

gions other than the activation loop can interact with PTPs. Therefore, binding between PTPs and Hog1 could regulate signaling independent of catalytic activity. In support of this, overexpression of the catalytically inactive mutant, Ptp2-C666S, inactivates the HOG pathway (Jacoby et al. 1997; Wurgler-Murphy et al. 1997). Further study of the interactions with PTPs that modulate Hog1 localization are underway.

Materials and methods

Strains, media, and genetic techniques

Hog1-GFP was visualized in the *hog1Δ::TRP1* strain, IMY100 (Jacoby et al. 1997), and in *ptpΔ* strains, CMY10 (*ptp2Δ::HIS3 hog1Δ::TRP1*), CMY11 (*ptp3Δ::TRP1 hog1Δ::TRP1*), and CMY12 (*ptp2Δ::HIS3 ptp3Δ::TRP1 hog1Δ::TRP1*), which were constructed by crossing the previously described *ptpΔ* strains (Jacoby et al. 1997) to IMY100. Hog1-GFP was also visualized in *ptc1Δ hog1Δ* and *ptc1Δ ptp2Δ hog1Δ* (Jacoby et al. 1997). PTPs were overexpressed from the plasmids, pKT-Ptp2, pKT-Ptp2C666A, pKT-Ptp3, and pKT-Ptp3C804A (Jacoby et al. 1997) in a *hog1Δ* strain, IMY200, derived from 334 (Hovland et al. 1989). IMY200 was produced by transforming the 334 strain with the *hog1Δ::hisG-URA3-hisG* allele (O'Rourke and Herskowitz 1998) and growing on media containing 5-fluoroorotic acid (5-FOA; American Biorganics, Inc., Swampscott, MA) to isolate the *hog1Δ::hisG* strain. Yeast transformations were performed according to Dohmen et al. (1991). Media to culture yeast were produced as described by Sherman et al. (1986).

Plasmids

Plasmids expressing wild-type and mutant Hog1-GFP were constructed as follows. A *NotI* site was substituted for the *HOG1* stop codon by use of PCR and ligated to a 750-bp *NotI* fragment containing GFP (J. Hirsch, Columbia University, NY) in the CEN-based plasmid, YCplac111 (Gietz and Sugino 1988). The resulting pHOG1-GFP complemented the osmotic stress sensitivity of the *hog1Δ* strain, IMY100 (Brewster et al. 1993; Jacoby et al. 1997). The HOG1-Y176F allele was produced by PCR with the oligonucleotide, 5'-AAGAATTCAAGACCCCTCAA-ATGACAGGCTTTGTTTCCACTAG-3', where the mutations are underlined. The *HOG1-K52M* allele was produced by PCR using the mutagenic oligonucleotide, 5'-CATCTCAGCCAGT-TGCCATTATGAAAATCATG-3', containing the underlined mutations. A 500-bp *EcoRI* fragment containing the K52M mutation was substituted for the wild-type *EcoRI* fragment in a plasmid bearing the wild-type *HOG1* gene. The *HOG1-D310N* allele was produced by site-directed mutagenesis of a 911-bp *PstI* fragment of the *HOG1* ORF by use of the oligonucleotide, 5'-CGGCTACTGGTTCATTTCGTTGGATCGTGG-3', containing the underlined mutations. This fragment was substituted for the wild-type *PstI* fragment in *HOG1*.

Immunoblot analysis

The level of Hog1-pY was monitored by immunoblotting. Cultures were grown to exponential phase at 30°C in synthetic media and were untreated or exposed to osmotic stress by the addition of NaCl to a final concentration of 0.4 M NaCl or 0.9 M NaCl. Approximately 7.5×10^7 cells were harvested by centrifugation, resuspended in 40 μ l of 90 mM Na-HEPES (pH 7.5), 2% SDS, and heated at 100°C for 3 min (Mattison et al. 1999). To this mixture, 40 μ l of ice-cold lysis buffer, 50 mM sodium

HEPES (pH 7.5), 5 mM EDTA, 150 mM NaCl, 1% Triton-X 100, with protease inhibitors (leupeptin, pepstatin A, antipain, aprotinin, and chymostatin each at 20 μ g/ml, and 35 μ g/ml PMSF), and protein phosphatase inhibitors, 1 mM sodium orthovanadate and 40 mM β -glycerophosphate was added. Immunoblotting was performed as described previously (Jacoby et al. 1997) using PY-99 (Santa Cruz Biotechnology) and anti-GFP antibody generously provided by C. Zuker (UCSD). Immunoreactivity was visualized using alkaline phosphatase conjugated to anti-mouse antibody and 5-bromo-4-chloro-3-indolyl phosphate and nitro-blue tetrazolium.

Fluorescence microscopy

Strains expressing GFP-fusion proteins were grown to ~ 1 unit (A_{600}) in synthetic medium lacking leucine to select for plasmids. To visualize the nucleus, cultures were diluted to 0.50 units (A_{600}), 4',6-diamidino-2-phenylindole (DAPI) was added to a concentration of 2.5 μ g/ml, and the cultures grown under low light conditions for 30 min at 30°C. Cultures were untreated or exposed to 0.4 M NaCl or 0.9 M NaCl for various times and the cells adsorbed to polylysine-coated slides (Polysciences, Inc., Warrington, PA). Fluorescence images were captured using a Zeiss Axioplan fluorescence microscope with a Photometrics Sensys digital CCD camera system. Images were processed using IP-LAB Spectrum software. Photographed cells were scored by eye as having nuclear Hog1 accumulation if the nuclear GFP signal was more intense than the cytoplasmic signal.

Coprecipitation assay

Yeast expressing wild type or mutant GST-PTP fusions, and wild-type or mutant Hog1-GFP fusions were grown to ~ 1 unit (A_{600nm}) in selective medium and harvested by centrifugation. Cells were lysed by glass beading in ice-cold lysis buffer, 50 mM Tris-HCl (pH 7.5), 50 mM NaCl, 5 mM $MgCl_2$, 0.2% Triton X-100, 0.1% 2-mercaptoethanol, with protease inhibitors (leupeptin, pepstatin A, antipain, aprotinin, and chymostatin, each at 20 μ g/ml, and 1 mM PMSF). Samples were analyzed by SDS-PAGE and immunoblotting with anti-GFP, anti-pY, and anti-GST (Pharmacia) antibodies.

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