The protein kinase C-activated MAP kinase pathway of *Saccharomyces cerevisiae* mediates a novel aspect of the heat shock response

Yoshiaki Kamada, Un Sung Jung, Julia Piotrowski, and David E. Levin

Department of Biochemistry, The Johns Hopkins University School of Public Health, Baltimore, Maryland 21205 USA

The *PKC1* gene of budding yeast encodes a homolog of the α, β, and γ isoforms of mammalian PKC that is proposed to regulate a MAPK-activation pathway. Mutants in this pathway undergo cell lysis resulting from a deficiency in cell wall construction when they attempt to grow at elevated temperatures. We show that the *PKC1*-regulated pathway is important for induced thermotolerance and that the MPK1 protein kinase (the MAPK of this pathway) is strongly activated by mild heat shock. This activation is sustained during growth at high temperature and is dependent on the function of pathway components proposed to function upstream of *MPK1*, including *PKC1*. Expression of genes under the control of known heat shock-inducible promoter elements (HSEs and STREs) was not compromised in *PKC1* pathway mutants, indicating that this pathway mediates a novel aspect of the yeast heat shock response. We propose that the heat-induced signal for pathway activation is generated in response to weakness in the cell wall created during growth under thermal stress, perhaps as a result of increased membrane fluidity. Evidence is presented that the mechanism by which the cell detects this weakness is by measuring stretch of the plasma membrane.

*Key Words: PKC; MAP kinase; heat shock; yeast; cell integrity*

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Members of the family of phospholipid-dependent, serine/threonine-specific protein kinases, known collectively as protein kinase C (PKC), respond to extracellular signals that act through receptor-mediated hydrolysis of phosphatidylinositol-4,5-bisphosphate to diacylglycerol (DAG) and inositol-1,4,5-trisphosphate (IP3) (Hokin 1985). DAG serves as a second messenger to activate PKC (Takai et al. 1979; Kishimoto et al. 1980; Nishizuka 1986, 1988), and IP3 functions to mobilize Ca2+ from intracellular stores (Berridge and Irvine 1984). A dozen distinct subtypes of mammalian PKC have been reported to date (Dekker and Parker 1994). The four initially identified isozymes, α, βI, βII, and γ, are structurally closely related and display similar catalytic properties. The activity of these enzymes can be stimulated by Ca2+, but at physiological Ca2+ concentrations DAG is also required for activation. Five additional isozymes, δ, ε, θ, η(λL), and μ, are related more distantly to the α, β, and γ subtypes and lack a conserved region thought to be important for Ca2+-binding. Consistent with this structural feature, these subtypes display Ca2+-independent protein kinase activity (Nishizuka 1992). Three recently identified subtypes, ζ, λ, and κ also lack the putative Ca2+-binding domain, but are unusual in that they lack one of the pair of DAG-binding zinc fingers.

Mammalian PKC is thought to play a pivotal role in the regulation of a host of cellular functions through its activation by growth factors and other agonists. These functions include cell growth, proliferation, and differentiation (Rosengurt et al. 1984; Kaibuchi et al. 1985; Persons et al. 1988; Michak et al. 1993), release of various hormones (Negro-Vilar and Lapetina 1985; Ohmura and Friesen 1985), and control of ion conductance channels (Farley and Auerbach 1986; Madison et al. 1986; Hardy et al. 1995). PKC induces the transcription of a wide array of genes, including the proto-oncogenes *c-myc, c-fos, c-sis,* and *c-jun* (Kelly et al. 1983; Greenberg and Ziff 1984; Krujer et al. 1984; Colamonici et al. 1986; Angel et al. 1988), human collagenase (Angel et al. 1987), metallothioneine IIα, and the SV40 early genes (Imbra and Karin 1986). Several transcription factors have been implicated in this response, including components of the AP-1 complex, AP-2, AP-3, and NF-κB (Chiu et al. 1987; Imagawa et al. 1987; Lee et al. 1987; Baehrle and Baltimore 1988).

Although substantial progress has been made toward elucidating the pathways leading to PKC activation, the steps between this activation and subsequent nuclear events are not well understood. However, members of a
family of enzymes called mitogen-activated protein kinases (MAPKs) have been implicated in PKC-dependent signaling. These enzymes are thought to function as intermediaries between membrane-associated signaling molecules and the nucleus (Pelech and Sanghera 1992; Thomas 1992). The observation that at least some members of the MAPK family translocate between the cytoplasm and the nucleus (Chen et al. 1992; Sanghera et al. 1992, Lenormand et al. 1993) makes them excellent candidates for messengers from the membrane. In support of this notion is the finding that a number of nuclear transcription factors are targets for MAPKs (Hill and Treisman 1995). MAPKs are activated through protein kinase cascades that are comprised of three highly conserved core components. The MAPKs themselves are activated in response to dual phosphorylation on a threonine and tyrosine residue by MAPK kinases (or MEKs) (Ahn et al. 1992). MEKs are phosphorylated and activated by members of a third family of protein kinases called MEK kinases (MEKKs; Herskowitz 1995).

The initially described p42/p44 isoforms of MAPK (ERKs) are activated in response to a wide array of extracellular agonists, including those that stimulate PKC, tyrosine kinases, and G proteins (Cobb et al. 1991). Additional subtypes of mammalian MAPKs reported recently include the p46 and p54 c-Jun amino-terminal kinases (JNKs) (Hibi et al. 1993; Derijard et al. 1994; Galcheva-Gargova et al. 1994, Kyriakis et al. 1994) and p38 MAPKs (Han et al. 1994; Rouse et al. 1994). Apparently distinct in function from the p42/p44 isoforms, these enzymes are activated in response to cytokines and a variety of stresses including heat shock, high osmolarity, hydrogen peroxide, UV light, sodium arsenite, and inhibition of protein glycosylation or synthesis. The mechanisms by which such varied stress signals activate these novel MAPK isotypes are not yet clear. However, JNK activation requires costimulation of both Ca2+ and PKC-dependent pathways in T cells (Su et al. 1994).

We are studying signal transduction from PKC in the budding yeast Saccharomyces cerevisiae. The PKC1 gene of S. cerevisiae encodes a homolog of the Ca2+-stimulated subtypes of mammalian PKC that is essential for cell growth (Levin et al. 1990). Loss of PKC1 function results in a cell lysis defect that is attributable to a deficiency in cell wall construction (Levin and Bartlett-Heubusch 1992; Paravicini et al. 1992, Levin et al. 1994). Several additional components of the signaling pathway regulated by PKC1 were identified by isolating genes whose mutational activation, or expression at high levels, could suppress the cell lysis defect resulting from pathway inactivation. Among these are four genes that encode protein kinases proposed to constitute a MAPK activation cascade that is under the control of PKC1. These include a MEKK homolog (BCK1; Lee and Levin 1992), a redundant pair of MEK homologs (MKK1/MKK2; Irie et al. 1993), and a MAPK homolog (MPK1; Lee et al. 1993, initially designated SLT2, Torres et al. 1991). Deletion of any of these components results in cell lysis only when cells are cultivated at 37°C. Because loss of PKC1 function results in a more severe defect (lysis at any temperature) than does loss of function of the downstream components, we have proposed that PKC1 regulates a bifurcated pathway, on one branch of which functions this MAPK activation cascade (Lee and Levin 1992). Genetic interactions among these protein kinase-encoding genes suggest that they function in a linear pathway from PKC1 to BCK1 to MKK1/MKK2 to MPK1 (Irie et al. 1993, Lee et al. 1993). This pathway is one of five physiologically distinct MAPK activation cascades known to exist in budding yeast (Herskowitz 1995; Levin and Errede 1995).

To identify the signals that activate the PKC1-regulated cell integrity pathway, we have developed an immune-complex protein kinase assay for the MPK1-encoded MAPK. We show here that this signaling pathway is strongly activated by growth under mild heat shock conditions. We also present evidence that heat-induced activation of this pathway is in response to the development of weakness in the cell wall that is detected by stretch of the plasma membrane.

Results

The PKC1-regulated signaling pathway is important for induced thermotolerance

Wild-type S. cerevisiae cells exhibit the ability to survive conditions of severe heat shock if they are first exposed to a mild heat shock (Sanchez et al. 1992). This behavior, known as induced thermotolerance, is attributable in part to the induced expression of a set of so-called heat shock genes (Parsell and Lindquist 1994). Because deletion mutants in the PKC1-regulated MAPK-activation pathway (bck1A, mkk1/mkk2A, and mpk1A) grow normally at 23°C, but undergo cell lysis when cultivated at 37°C (Torres et al. 1991; Lee and Levin 1992, Irie et al. 1993; Lee et al. 1993), we explored the possibility that this pathway plays a role in induced thermotolerance. Figure 1A shows that a bck1A mutant was no more sensitive than an isogenic BCK1+ strain to killing by shift from logarithmic growth at 23–50°C. However, the bck1A mutant was partially deficient in the acquisition of thermotolerance induced by brief pretreatment at 37°C (30 min; Fig. 1B). Similar results were obtained using an mpk1A mutant [data not shown]. In addition, a constitutive allele of BCK1 (BCK1-20), which alleviates the requirement for PKC1 (Lee and Levin 1992), conferred partial resistance to heat shock (Fig. 1A). These results indicate that the PKC1-regulated pathway plays a role in induced thermotolerance and suggest that this pathway might be activated in response to heat shock.

The MPK1-encoded MAPK is activated by heat shock in a PKC1-dependent manner

To test directly the effect of growth temperature on PKC1-regulated pathway activity, we developed an immune complex protein kinase assay for the MPK1-encoded MAPK. The MPK1 gene was fused to a sequence encoding an epitope from the influenza virus hemagglu-
MPK1 HA was immunoprecipitated from extracts of cells growing at various temperatures. The protein kinase activity associated with MPK1 HA was measured in immune complex using bovine myelin basic protein (MBP) as substrate, an excellent substrate for mammalian MAPks. MPK1 HA from cells growing at 23°C was very weakly active, enzyme from cells at 30°C was moderately active, and enzyme from cells at 37°C was strongly active (Fig. 2B). Because extracts were made after long-term cultivation of cells in log phase (12–15 hr), we conclude that MPK1 HA activity remains persistently high in cells growing at elevated temperature. The range of activity at different temperatures reflects differences in MPK1 HA specific activity because the amount of this protein in the immunoprecipitates did not vary appreciably in cells grown at different temperatures (Fig. 2B, bottom). In addition, the observed differences in protein kinase activity are not a function of growth rate because the optimal growth temperature for this species is 30°C. This protein kinase activity is an intrinsic property of the MPK1 HA protein and is not attributable to a contaminating protein kinase, because a biologically inactive mutant form of this protein was biochemically inactive (see below).

To examine the kinetics of MPK1 HA activation in response to temperature change, growing cells were shifted rapidly from 23°C to 39°C by dilution into prewarmed medium. Figure 2C shows that activity remained low until 20 min after temperature shift and peaked by 30 min. Later time points showed no further increase in activity [not shown]. This activation represents a 170-fold increase in MPK1 HA specific activity. The delay in MPK1 HA activation in response to elevated temperature is considerably longer than that observed for activation of two other yeast MAPks in response to extracellular signals. The FUS3 and HOG1 protein kinases become fully active within 1 or 2 min after.

**Figure 1.** BCK1 is important for induced thermotolerance. Cells growing in YEPD at 23°C were either shifted directly to 50°C for the indicated times [A] or pretreated at 37°C for 30 min before the challenge temperature [B]. After heat shock, cells were diluted and plated for viability. For B, percent survival is relative to value after pretreatment. Strains were wild-type [1788], BCK1-20 bearing YEp352[MPK1::HA] (DL1183), and bck1A (DL251). Overproduction of MPK1 from YEp352[MPK1::HA] enhanced thermal resistance conferred by BCK1-20. Each value is the mean and standard deviation of at least three experiments.

**Figure 2.** Activation of MPK1 by growth at elevated temperatures. [A] Immunoblot detection of MPK1 HA. Immunoprecipitations using the anti-HA antibody [12CA5] were done with cell extracts [50 µg protein] prepared from a wild-type strain expressing either untagged MPK1 (DL744), or MPK1 HA (DL922) in the presence [lanes 1,2] or absence [lanes 3,4] of antibody. Immune complexes were subjected to SDS-PAGE on a 7.5% acrylamide gel. Immunodetection was performed as described in Materials and methods. The molecular masses of the standards are indicated [in kD]. The asterisk indicates commmunoprecipitated 12CA5 IgG. [B] MPK1 activity is dependent on growth temperature. MPK1 HA was immunoprecipitated from extracts of cultures [DL1101] expressing MPK1 HA growing at 23°C [lanes 1,2], 30°C [lanes 3,4], or 37°C [lanes 5,6]. In vitro protein kinase assays were conducted using MBP as a substrate, and the reactions were subjected to SDS-PAGE on a 12.5% acrylamide gel, as described in Materials and methods [top]. Autophosphorylated MPK1 HA and phosphorylated 12CA5 IgG [asterisk] are also indicated. All other bands derive from the MBP preparation. A parallel set of immune complexes were subjected to immunodetection of MPK1 HA [bottom]. [C] MPK1 is activated by heat shock. MPK1 HA was immunoprecipitated from extracts of DL1101 cultures shifted from 23°C to 39°C for the indicated times. Protein kinase assays [top] and immunodetection of MPK1 HA [bottom] were carried out on parallel sets of immune complexes.
exposure of cells to their respective activating signals (Gartner et al. 1992; Brewster et al. 1993). This suggests that elevated growth temperature does not induce a direct signal for MPK1\(^\text{HA}\) activation but that this activation is the result of a physiological change arising from growth under the new environmental condition. To determine whether MPK1\(^\text{HA}\) activation depends on the synthesis of a new protein, cells growing at 23°C were pretreated with the protein synthesis inhibitor cycloheximide before a 30-min shift to 39°C. Initial MPK1\(^\text{HA}\) activation was not impaired by the absence of protein synthesis (Fig. 3A, cf. lanes 2 and 5), indicating that the delay in thermally induced activation is not attributable to a requirement for the synthesis of new proteins. Cycloheximide treatment alone had a slight activating effect on MPK1\(^\text{HA}\) (threefold; Fig. 3A, cf. lanes 1 and 4).

To confirm that the protein kinase activity detected was an intrinsic property of MPK1\(^\text{HA}\), rather than a contaminating activity in the immune precipitates, a mutant form of this protein that lacks the two phosphorylatable residues predicted to be required for activation was tested for protein kinase activity. Mutation of either (or both) Thr-190 to Ala, or Tyr-192 to Phe, severely diminishes the biological activity of MPK1 (Lee et al. 1993). An MPK1\(^\text{T190A, Y192F::HA}\) double mutant allele was shifted from growth at 23°C to 39°C for 30 min. Immunoprecipitated MPK1–TAYF\(^\text{HA}\) displayed no detectable protein kinase activity (Fig. 3A, lane 7).

To test the veracity of the model proposed for the architecture of the PKC1-regulated pathway (Lee et al. 1993), we examined the requirement for BCK1 function in MPK1\(^\text{HA}\) activation. MPK1\(^\text{HA}\) isolated from a temperature-shifted strain bearing a bck1Δ mutation possessed no detectable activity (Fig. 3A, lane 9), confirming the dependence on BCK1 for MPK1\(^\text{HA}\) activity. As an additional test of the proposed model, we examined the requirement for PKC1 function in the activation of MPK1\(^\text{HA}\). The MPK1\(^\text{HA}\) protein was expressed in a strain bearing a temperature-sensitive allele of PKC1 (pkc1-2; Levin and Bartlett-Heubusch 1992). The mutant strain was subjected to a 30-min shift from 25°C to 37°C, a treatment that had the dual effect of creating the appropriate conditions for MPK1 activation while inactivating the thermolabile PKC1. In addition, this treatment does not cause appreciable killing of the mutant over the time course of the experiment (Levin and Bartlett-Heubusch 1992). Figure 3B shows that this temperature shift failed to activate MPK1\(^\text{HA}\) in the pkc1 mutant, confirming the requirement for PKC1 in signal transmission to this MAPK.

**Figure 3.** Activation of MPK1 does not require protein synthesis, but it does require signals from PKC1 and BCK1. (A) Stimulation of MPK1 protein kinase activity in response to shift from growth at 23°C–39°C for 30 min was detected as described in Fig. 2 (top). Strains were wild-type [DL922, expresses MPK1\(^\text{HA}\), mpk1-TAYF\(^\text{::HA}\) [DL1182, expresses MPK1-T190A, Y192F::HA], and bck1A::URA3 [DL923, expresses MPK1\(^\text{HA}\)]. (Lanes 4, 5) Cycloheximide [100 μg/ml] was added to the culture 30 min before heat shock. A parallel set of immune complexes were subjected to immunodetection of MPK1\(^\text{HA}\) [bottom]. (B) Stimulation of MPK1 protein kinase activity in response to shift from growth at 25°C to 37°C for 30 min in a temperature-sensitive pkc1 mutant. Strains were wild type [DL1101, expresses MPK1\(^\text{HA}\)] and pkc1-2 [DL1102, expresses MPK1\(^\text{HA}\)]. Protein kinase assays [top] and immunodetection of MPK1\(^\text{HA}\) [bottom] were carried out on parallel sets of immune complexes.

**Heat shock gene expression is not regulated by the PKC1 pathway**

Mild heat shock induces the expression of an array of genes under the control of the heat shock transcription factor (HSF; Sorger and Pelham 1988; Weiderrecht et al. 1988). Many of these genes encode molecular chaperones that aid in the refolding or proteolytic degradation of unfolded proteins under thermal stress conditions (Mortimoto et al. 1994). HSF binds to a promoter element in these genes known as the heat shock element (HSE; Munro and Pelham 1985). To determine whether heat shock-induced gene expression is dependent on the PKC1-regulated pathway, we measured the expression of a reporter gene (Escherichia coli lacZ) driven by either a synthetic- or natural HSE-associated promoter in an mpk1Δ mutant and a pkc1Δ mutant. Table 1 shows that heat shock-induced expression from these promoters was normal in both mutants, although the pkc1Δ mutant displayed an elevated constitutive level of expression. This indicates that the PKC1-regulated pathway is not responsible for transcriptional activation of genes under HSE control. But because the expression of some heat shock genes is also regulated post-transcriptionally (e.g., HSP104; S. Lindquist, pers. commun.), we examined the heat shock-induced accumulation of the HSP104 protein in a bck1Δ mutant. Here again, no difference was observed between the mutant and wild-type strain (not shown).

A second promoter element that is activated in response to heat shock is the stress-responsive element [STRE; Marchler et al. 1993]. To determine whether STRE-driven gene expression is dependent on the PKC1-mediated pathway, we measured expression of the CTT1...
gene in an \textit{mpk1}\Delta mutant and a \textit{pkc1}\Delta mutant. The \textit{CCT1} gene encodes the major yeast catalase [catalase T] and is induced transcriptionally by stresses such as heat shock, hydrogen peroxide, and high osmolarity (Marchler et al. 1993; Schuller et al. 1994). Both mutants displayed normal induction of catalase activity in response to heat shock (Table 1), indicating that the \textit{PKC1} pathway is not responsible for heat shock-induced expression of genes under STRE control. Therefore, thermal activation of this pathway must serve some other purpose.

Stresses other than heat shock that cause proteins to unfold also induce the expression of heat shock genes. Ectopic expression of denatured proteins in cells is sufficient to stimulate heat shock gene expression (Parsell and Lindquist 1994), suggesting that unfolded proteins constitute the primary signal for activation of HSF. We examined the activation of MPK1\textsuperscript{HA} in response to several protein denaturing stresses. Treatment of cells growing at 23°C with 400 \textmu M hydrogen peroxide, 6% ethanol, or 750 \textmu M sodium arsenite, all of which induce expression of HSE-driven genes (Sanchez et al. 1992), failed to activate MPK1\textsuperscript{HA} (less than twice basal activity). Therefore, it is not likely that unfolded proteins resulting from heat shock are the signal that stimulates the \textit{PKC1} pathway.

\textbf{The \textit{PKC1}-regulated signaling pathway is activated in response to plasma membrane stretch}

If the \textit{PKC1} pathway is not activated by the same signal that induces heat shock gene expression, then to what aspect of thermal stress might this pathway respond? In addition to unfolding proteins, heat shock also increases the fluidity of membranes. Because mutants in the \textit{PKC1}-regulated pathway undergo cell lysis resulting from a deficiency in cell wall construction, we speculated that the activating signal might be generated as a consequence of weakness in the cell wall created during growth at high temperature, possibly attributable to increased membrane fluidity. The observed delay in MPK1\textsuperscript{HA} activation in response to elevated temperature can be explained by the development of weakness in the cell wall during growth under thermal stress. One mechanism by which the cell might detect weakness in the wall is by measuring stretch of the membrane. The yeast plasma membrane possesses mechanosensitive ion channels that are activated by stretch (Gustin et al. 1988).

As a first test of the model that the \textit{PKC1}-regulated pathway is activated by plasma membrane stretch, we examined the effect of altering the extracellular osmolarity on MPK1\textsuperscript{HA} activity. One prediction of this model is that growth in the presence of high extracellular osmolarity, which prevents outward stretch of the plasma membrane, should prevent activation of MPK1\textsuperscript{HA}. Cells growing in the presence or absence of 0.5 M potassium acetate were subjected to a 30-min temperature shift from 24°C to 39°C. Figure 4 shows that activation of MPK1\textsuperscript{HA} by elevated temperature was prevented to a large degree in cells growing in the high osmolarity medium (cf. lanes 2 and 5). This inhibition of MPK1\textsuperscript{HA} activation was not specific to potassium acetate but was also caused by 10% sorbitol [not shown], indicating that this is a general effect of increased osmolarity. Moreover, prevention of heat-induced MPK1\textsuperscript{HA} activation by high osmolarity is not a consequence of activating the high osmolarity glycerol [HOG] response pathway, because potassium acetate prevented activation of MPK1\textsuperscript{HA} in a mutant in this pathway (\textit{pbs2}\Delta; Fig. 4, cf. lanes 7 and 9). Although mechanisms other than the HOG pathway may exist to signal high extracellular osmolarity, these results suggest that elevated osmolarity prevents MPK1\textsuperscript{HA} activation as a direct consequence of the increased external pressure on the plasma membrane. This conclusion is supported by the observation that increasing the osmolarity at the time of temperature shift also prevented MPK1\textsuperscript{HA} activation effectively [not shown].

\begin{table}
\centering
\caption{Heat shock gene expression is normal in \textit{mpk1} and \textit{pkc1}\Delta mutants}
\begin{tabular}{|c|c|c|c|c|}
\hline
Genotype & Temperature & HSE-driven \beta-galactosidase & STRE-driven catalase T activity [relative units]
& & activity [relative units]
& & SSA1/HSP26 & synthetic & \\
\hline
Wild type & 23 & 1.0 & 3.2 & 1.0 \\
& 39 & 19.0 & 20.6 & 23.9 \\
\textit{mpk1}\Delta & 23 & 1.2 & 4.2 & 1.1 \\
& 39 & 17.8 & 21.8 & 22.0 \\
Wild type & 23 & 1.2 & N.T. & 2.1 \\
+ salt & 39 & 14.4 & N.T. & 25.8 \\
\textit{pkc1}\Delta & 39 & 4.6 & N.T. & 5.7 \\
+ salt & 39 & 30.8 & N.T. & 36.7 \\
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Strains were wild-type (DL1161 and DL1162), \textit{mpk1}\Delta (DL1157 and DL1158), and \textit{pkc1}\Delta (DL1186). Plasmid pB11 (Susek and Lindquist 1990) expresses \textit{E. coli lacZ} under the control of the yeast \textit{CYC1} promoter driven by upstream activating sequences from \textit{HSP26} and \textit{SSA1}. Plasmid CS11 is a similar construct with four synthetic HSEs fused to the \textit{CYC1} promoter. The elevated basal level of catalase T activity in the presence of salt (0.5 M potassium acetate) results from transcriptional activation of \textit{CTT1} by high osmolarity (Schuller et al. 1994). Each value for \beta-galactosidase activity is the mean of at least two experiments. [N.T.] Not tested.
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\end{table}
Another prediction of this model is that MPK1\textsuperscript{HA} should be activated by environmental changes that induce plasma membrane stretch. We used two mechanisms for inducing stretch. First, we tested the ability of a rapid reduction in extracellular osmolarity to activate MPK1\textsuperscript{HA}. Cells growing at 23\textdegree C in high osmolarity medium (YEPD + 1 M sorbitol) were diluted with YEPD [to 0.2 m sorbitol]. MPK1\textsuperscript{HA} was activated transiently by this treatment [Fig. 5A]. Although activation was weak as compared with heat shock activation, the kinetics were strikingly rapid, with activity peaking 1 min after osmotic shock. This is the expected result if MPK1\textsuperscript{HA} is activated in response to plasma membrane stretch induced by the instantaneous reduction in external osmolarity. MPK1\textsuperscript{HA} activity returned to the preactivation level within 5–10 min after this shock. This too is expected, because as the cell opens membrane ion channels to collapse the osmotic gradient, the outward pressure on the membrane is relieved.

The second mechanism used to induce stretch of the plasma membrane was treatment of cells with chlorpromazine (CPZ), a cationic amphipath, which preferentially inserts into the cytoplasmic leaflet of the plasma membrane lipid bilayer. This agent is very effective at inducing inward membrane stretch in vivo (Sheetz and Singer 1974), and activating mechanosensitive ion channels in vitro (Martinac et al. 1990). Although these ion channels can be activated by either inward or outward membrane stretch, we chose an agent that induces inward stretch to avoid constraint by the cell wall. Treatment of cells growing at 25\textdegree C with 250 \mu M CPZ strongly activated MPK1\textsuperscript{HA} (Fig. 5B, cf. lanes 1 and 4). The level of MPK1\textsuperscript{HA} activity in the presence of CPZ was nearly as high as that induced by heat shock (cf. lanes 2 and 4). CPZ has also been reported to inhibit calmodulin in a Ca\textsuperscript{2+}-dependent manner (Babu et al. 1988). To exclude the possibility that CPZ exerts its activating effect on MPK1\textsuperscript{HA} through an indirect mechanism involving calmodulin, we tested a strain with a mutant calmodulin that is unable to bind Ca\textsuperscript{2+} (cmd1-6; Geiser et al. 1991), and therefore, should not interact with CPZ. This mutant behaved identically to the isogenic CMD1\textsuperscript{+} strain with respect to MPK1\textsuperscript{HA} activation by CPZ (Fig. 5B; lanes 5,7). Although it is possible that CPZ has additional effects on cells that could result in the indirect activation of MPK1\textsuperscript{HA}, it seems most likely that the observed activation results from CPZ-induced membrane stretch.

**Intracellular localization of MPK1**

Because some MAPKs have been reported to translocate between the cytoplasm and the nucleus (Chen et al. 1992; Sanghera et al. 1992; Lenormand et al. 1993), we examined the intracellular localization of MPK1\textsuperscript{HA} under various thermal conditions. Indirect immunofluorescence of MPK1\textsuperscript{HA} revealed that this MAPK resides in
Both the cytoplasm and the nucleus in growing cells. MPK1\(^{HA}\) appeared to be concentrated in the nucleus in cells growing at 24\(^\circ\)C (Fig. 6A). In contrast, in cells growing at 37\(^\circ\)C, staining was distributed more uniformly between the two compartments (Fig. 6C). The pattern of localization in cells shifted from 24\(^\circ\)C to 39\(^\circ\)C for 30 min (Fig. 6B) resembled the uniform staining observed after long-term cultivation at high temperature, except that MPK1\(^{HA}\) appeared to be excluded from the enlarged vacuoles that develop in response to heat shock. Because expression of MPK1\(^{HA}\) is not altered by heat shock or prolonged growth at elevated temperature (data not shown), the observed staining patterns reflect a redistribution of nuclear MPK1 to the cytoplasm at high temperature.

Discussion

The PKC1-regulated signaling pathway is important for induced thermotolerance

Heat shock disrupts a variety of cellular structures and processes by causing protein unfolding and increased membrane fluidity (for review, see Parsell and Lindquist 1994). The yeast heat shock response is a complex process that allows cells to grow at elevated temperatures and survive brief exposure to extreme temperatures. The best studied aspect of the heat shock response is the phenomenon of induced thermotolerance. This refers to the resistance to killing by high temperatures (e.g., 50\(^\circ\)C) that is acquired by brief pretreatment at a moderately elevated temperature (e.g., 37\(^\circ\)C). The thermally induced synthesis of a set of proteins, known collectively as heat shock proteins (HSPs), plays an essential role in induced thermotolerance (Parsell and Lindquist 1994). HSPs function in two ways to prevent the accumulation and aggregation of denatured proteins. Some act as molecular chaperones that promote protein refolding, whereas others are involved in the degradation of denatured proteins (for review, see Morimoto et al. 1994).

In this study we demonstrate that the PKC (encoded by \(PKC1\))-regulated cell integrity signaling pathway mediates a novel aspect of the yeast heat shock response. Mutants in the protein kinases under the control of \(PKC1\) (BCK1, MKK1/MKK2, and MPK1) undergo cell lysis because of a deficiency in cell wall construction when they are induced to grow at high temperatures (e.g., 37\(^\circ\)C) (for review, see Levin and Errede 1995). We have demonstrated here that (1) such mutants are deficient in the ability to survive a severe heat shock after brief pretreatment with a mild heat shock; (2) a constitutive mutant in the \(PKC1\)-regulated pathway (BCK1-20) is partially resistant to severe heat shock, indicating that one aspect of induced thermotolerance is activated constitutively in this mutant; (3) the activity of the MAPK (encoded by MPK1) under the control of \(PKC1\) is stimulated 170-fold in growing cells in response to a mild heat shock (i.e., 23\(^\circ\)C to 39\(^\circ\)C). This increased activity is sustained indefinitely during growth at elevated temperature; and (4) mutants in the \(PKC1\) pathway are not compromised for the ability to induce expression of genes under the transcriptional control of known heat shock-activated promoter elements (i.e., HSEs or STREs), indicating that this pathway plays a novel role in induced thermotolerance. To our knowledge, this is the first report linking PKC to a heat shock response.

The PKC1-regulated pathway is activated by plasma membrane stretch

Mutants in the \(PKC1\) pathway are deficient in remodeling their cell walls during growth. We propose that the heat-induced signal for pathway activation is generated as a consequence of weakness in the cell wall created during growth under conditions of thermal stress, perhaps resulting from increased fluidity of the plasma membrane (Fig. 7). An increase in membrane fluidity could disrupt the interaction between the plasma membrane and the cell wall, thereby adversely affecting wall biosynthesis. This model is consistent with the observation that heat shock-induced activation of MPK1 followed a delay of ~20 min, suggesting that the signal develops as a consequence of growth under the new environmental condition.

We propose further that the \(PKC1\)-regulated signaling pathway detects and responds to weakness in the cell wall by measuring plasma membrane stretch (Fig. 7). Three lines of evidence support this model. First, growth in high osmolarity medium greatly inhibited the heat

**Figure 6.** Intracellular localization of MPK1\(^{HA}\) by indirect immunofluorescence. Diploid cells, expressing MPK1\(^{HA}\) (DL1232) or untagged MPK1 (1788), were grown in log phase for 12–15 hr at 24\(^\circ\)C (A) or 37\(^\circ\)C (C), or shifted from growth at 24\(^\circ\)C to 39\(^\circ\)C for 30 min (B). Cells were stained with anti-HA antibodies (12CA5) and DAPI (to visualize DNA).
shock-induced activation of MPK1. This effect appears to be a direct consequence of the increased external pressure exerted on the plasma membrane, because it did not require the function of the HOG pathway. Second, a reduction in extracellular osmolarity induced a very rapid, but transient, activation of MPK1, consistent with the instantaneous membrane stretch such treatment would induce. Tyrosine phosphorylation of MPK1 (as an indirect measure of MPK1 activation) has also been detected in response to reduced osmolarity (M.C. Gustin, pers. comm.). Third, treatment of cells with CPZ, an amphipathic drug that induces membrane stretch by preferentially inserting into the cytoplasmic leaflet of the lipid bilayer strongly activated MPK1. Taken in the aggregate, these results indicate that membrane stretch is sufficient to activate the PKC1 signaling pathway.

Yeast cells possess mechanosensitive ion channels that are activated by membrane stretch and are capable of importing Ca^{2+} (Gustin et al. 1988). Yeast cells experience a rapid influx of Ca^{2+} in response to mild heat shock (H. Iida, pers. comm.) that may result from the opening of such ion channels. Because PKC1 is related to the Ca^{2+}-activated family of PKC isoforms (Levin et al. 1990), it may be activated directly in response to Ca^{2+} influx. Alternatively, PKC1 may be activated indirectly through the stimulation of a Ca^{2+}-dependent phospholipase C [PLC]. Although such a phospholipase has been described in yeast [the product of the PLC1 gene] (Flick and Thorner 1993; Payne and Fitzgerald-Hayes 1993; Yoko-o et al. 1993), this enzyme does not appear to function within the PKC1-regulated signaling pathway. It is also possible that activation of a PLC is coupled directly to ion channel activation.

It is not yet clear what types of pathway output are engaged in response to activation of the PKC1-regulated pathway. Persistent pathway activation in response to elevated temperature might induce the expression of genes whose products can reduce membrane fluidity, perhaps by increasing the level of saturation of the fatty acids in the membrane. Expression of genes involved in cell wall biosynthesis might also be regulated by this pathway. However, if the PKC1 pathway regulates gene expression in response to heat shock, it must do so through a mechanism other than activation of the known heat-inducible promoter elements. Alternatively, it is possible that this pathway regulates functions other than gene expression. For example, the actin cytoskeleton, which controls polarized cell growth (Drubin 1991), may be reorganized in response to pathway activation. The observed translocation of MPK1 from the nucleus to the cytoplasm at elevated temperature suggests that this MAPK serves a function outside of the nucleus.

Transient pathway activation caused by a reduction in extracellular osmolarity may induce a qualitatively different output than that resulting from sustained activation in response to heat shock. For example, ion channel conductance may be regulated by pathway activity during the equilibration of internal and external osmolarity. This may be relevant to the observation that mammalian PKC regulates the function of some ion channels (Farley and Auerbach 1986; Madison et al. 1986; Hardy et al. 1995). Precedents exist for different responses to transient versus sustained activation of mammalian MAPKs, which may be explained by translocation of MAPKs only under conditions of sustained activation (Marshall 1995).

The PKC1 pathway may be an appropriate model for mammalian signaling pathways that regulate stress-activated MAPKs. The JNK and p38 MAPKs are regarded as functional homologs of the yeast MAPK encoded by the HOG1 gene because all of these enzymes are activated in response to high extracellular osmolarity (Galcheva-Gargova et al. 1994; Han et al. 1994). However, unlike HOG1, the mammalian enzymes are activated by a variety of other stresses, including heat shock (Hibi et al. 1993; Kyriakis et al. 1994; Rouse et al. 1994). The requirement for PKC activation in the stimulation of JNK in T cells [Su et al. 1994] suggests that a pathway similar to that regulated by PKC1 might function in mammalian cells to signal membrane stretch. Mammalian cells possess mechanosensitive ion channels (Martinac et al. 1990), which presumably allow them to detect plasma membrane stretch. These cells may monitor the state of...
their extracellular matrices by a mechanism similar to that by which we propose yeast cells to monitor the state of their cell walls.

Other signals that lead to activation of PKC1

The PKC1-regulated pathway is also required for cell integrity during mating factor-induced morphogenesis (Errede et al. 1995). Mutants in BCK1 or MPK1 lyse when treated with mating factor at temperatures that are permissive for vegetative growth. The kinetics of cell lysis indicate that mutant cells die at the onset of morphogenesis (Errede et al. 1995). These observations may be explained by the finding that during mating factor-induced morphogenesis, a large region of the cell wall undergoes remodeling in a relatively short period, potentially creating weakened areas that would result in stimulation of the PKC1 pathway.

Cells depleted of PKC1 lyse at a uniform point in the cell cycle, with small-to-medium sized buds (Levin et al. 1990). We have proposed that this is because a deficiency in cell wall remodeling is manifested most severely in the early stages of bud formation (Levin and Bartlett-Heubusch 1992), during which growth is polarized to the bud tip (Farkas et al. 1974; Lew and Reed 1993). This is supported by the observation that PKC1-depleted cells lyse specifically at their bud tips (Levin et al. 1994). Therefore, it might be expected that PKC1-regulated pathway activity would fluctuate with the cell cycle. In addition, a mutation in MPK1 was isolated in a search for mutations that enhance the growth defect of a temperature-sensitive allele of the CDC28 cell cycle gene (Mazzeno et al. 1993), further suggesting that the cell cycle engine plays a role in cell wall construction. However, measurements of MPK1 activity in synchronized cells have failed to reveal fluctuations through the cell cycle (Y. Kamada and D.E. Levin, unpubl.). This observation seems to preclude any model in which PKC1 activity is regulated by the cell cycle engine.

Materials and methods

Strains, growth conditions, transformations, and nucleic acid manipulation

All yeast strains used in this study (Table 2) were derivatives of EG123 [MATa leu2-3,112 ura3-52 trp1-1 his4 can11 (Siliciano and Tatchell 1984), except CRY1 and JCY149, which were the gift of Trisha Davis (University of Washington, Seattle). Yeast cultures were grown in YEPD medium (1% Bacto-yeast extract, 2% Bacto-peptone, 2% glucose). Synthetic minimal medium (SD; Rose et al. 1990) supplemented with the appropriate nutrients was used to select for plasmid maintenance and gene replacements. The pbs2A::LEU2 plasmid (pMA11) used to delete the PBS2 gene was the gift of M.C. Gustin (Rice University, Houston, TX). A 3.7-kb SacI-SpeI fragment was used for yeast transformation. Yeast transformation was by the lithium ace-

Table 2.  S. cerevisiae strains

| Strain   | Genotype                                      | Source                                      |
|----------|-----------------------------------------------|---------------------------------------------|
| EG123    | MATa leu2-3,112 ura3-52 trp1-1 his4 can11    | I. Herskowitz (University of California)    |
| 1783     | MATa EG123                                    | I. Herskowitz (University of California)    |
| 1788     | MATa/MATa isogenic diploid of EG123           | I. Herskowitz (University of California)    |
| DL521    | MATa/MATa 1788 bck1Δ::URA3/bck1Δ::URA3       | Lee and Levin (1992)                        |
| DL456    | MATa/MATa 1788 mpk1Δ::TRP1/mpk1Δ::TRP1       | Lee et al. (1993)                           |
| DL744    | MATa/MATa DL456 [YEp351[MPK1]]               | this study                                  |
| DL807    | MATa/MATa DL456 [YEp352[MPK1::HA]]           | this study                                  |
| DL952    | MATa/MATa DL922 bck1Δ::URA3/bck1Δ::URA3      | this study                                  |
| DL1049   | MATa/MATa DL807 pbs2Δ::LEU2/pbs2Δ::LEU2      | this study                                  |
| DL1101   | MATa 1783 [YEp352[MPK1::HA]]                 | this study                                  |
| DL1102   | MATa DL1101 pkc1Δ::LEU2 [YCp50[URA3[PKC1-2]] | this study                                  |
| DL1157   | MATa/MATa DL456 [pB11]                       | this study                                  |
| DL1158   | MATa/MATa DL456 [CS11]                       | this study                                  |
| DL1161   | MATa/MATa DL1788 [pB11]                      | this study                                  |
| DL1162   | MATa/MATa DL1788 [CS11]                      | this study                                  |
| DL1182   | MATa/MATa DL1788 [YEp352[mpk1-T190A, Y192F::HA]] | this study                                  |
| DL1183   | MATa/MATa DL1788 BCK1-20/BCK1-20 [YEp352[MPK1::HA]] | this study                                  |
| DL1186   | MATa DL1783 pkc1Δ::LEU2 [pB11]               | this study                                  |
| DL1210   | MATa ade2-1 his3-11,15 leu2-3,112 trp1-1 ura3-1 can1-100 [YEp351[MPK1::HA]] | this study                                  |
| DL1211   | MATa DL1210 cmd1-6                           | this study                                  |
| DL1232   | MATa/MATa DL456 [YCp50LEU2[MPK1::HA]]        | this study                                  |

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tate method (Ito et al. 1983). General genetic manipulation of
breedization, and DNA sequence analysis were also carried out as
described previously {Rose et al. 1990}. Induced thermostolerance of yeast strains was tested as
described by Sanchez et al. (1992). Oligonucleotide labeling, hy-
bridization, and DNA sequence analysis were also carried out as
described previously {Levin et al. 1987}.

Epitope tagging of MPK1

MPK1 was tagged at its carboxyl terminus with an epitope from the
influenza virus HA protein {Wilson et al. 1984}. In the first
of two steps, a 2.7-kb EcoRI-SphI fragment bearing various al-
leles of the entire MPK1 gene in M13mp18 was subjected to
mutagenesis with the primer 5'-GATAGAAAATATTTTG-
sequence analysis. E. coli cells were cultured in Luria broth or YT
medium (0.75% Bacto-tryptone, 0.5% Bacto-yeast extract, 0.5%
NaCl) and transformed or infected with M13 by standard meth-
ods {Maniatis et al. 1982}.

Indirect immunofluorescence

Indirect immunofluorescence was done by modification of the
procedure described in Berkower et al. {1994}. Cultures of log
phase cells (~10e6 cells) growing in YEPD were fixed by direct
addition of formaldehyde to the medium to a final concentra-
tion of 4%. Cells were fixed for 30 min at room temperature
before harvest by centrifugation. Fixed cells were washed twice
with 5 ml of KP buffer [0.1 M potassium phosphate, pH 6.5] and
resuspended in 2 ml of KP buffer [KP with 250 mM sorbitol]. To
remove the cell wall, 0.5 ml of cell suspension was incubated
with 20 μg/ml of Zymolyase 100T (ICN) and 0.2% β-me-
captopothenol at 30°C for 30 min. Spheroplasts were washed with 5
ml of KPS and harvested by gentle centrifugation (3000g for 2
min) and resuspended in 0.5 ml of KPS [KPS with 0.05%
agarase] (gift of B. Futcher, Cold Spring Harbor Laboratory), which carries the
9-amino-acid HA epitope in tandem triplication, was subcloned
into the newly created NotI site of MPK1.

Immunoprecipitations

MPK1HA was immunoprecipitated from cell extracts using the
12CA5 monoclonal antibody {BabCo}, which recognizes the HA
epitope. Cell extracts (50 μg of protein) were added to immuno-
precipitation buffer [lysis buffer without protease inhibitors,
with 1% NP-40 and sodium pyrophosphate at pH 7.5] to a final
volume of 0.5 ml, to which was added 2 μl of mAb 12CA5 raw
ascites diluted 1:10 in Tris-buffered saline [TBS; 50 mM Tris-
HCl [pH 7.4], 8% NaCl, 0.2% KCl]. This mixture was incubated
for 1 hr at 4°C with gentle agitation. To the extract/antibody
mixture was added 20 μl of a 50% suspension of protein
A-Sepharose beads [Sigma] equilibrated in immunoprecipita-
tion buffer, followed by a further incubation for 1 hr at 4°C with
gentle agitation. Immune complexes were washed four times
with immunoprecipitation buffer [without sodium pyrophos-
phate or KF], twice with buffer M [25 mM MOPS [pH 7.5], 1 mm
EGTA, 0.1 mM Na3VO4, 15 mM p-nitrophenylphosphate] and
once with kinase assay buffer [buffer M with 15 mM MgCl2].

In vitro phosphorylation assay

Phosphorylation assays were conducted using immunoprecipi-
tated MPK1HA. Immunoprecipitates were suspended in 18 μl of
kinase assay buffer with 10 μg of MBP [from bovine brain, Sigma]. This mixture was preincubated for 3 min at 30°C before
the reaction was initiated by adding 2 μl of 1 mM [γ32P]ATP [16
μCi/μmol], and incubation for 20 min at 30°C, the re-
action was terminated by addition of 30 μl of 2× Laemml’s
sample buffer (Laemmli 1970), and samples were boiled and
subjected to SDS-PAGE on 12.5% acrylamide gels. After electrophoresis, gels were immersed in 12.5% trichloroacetic acid
for 1 hr, followed by three washes for 30 min in 10% methanol,
10% acetic acid, to reduce background. Dried gels were sub-
jected to autoradiography or phosphorimaging {LAS 1000, Fujix}.

Immunoblots

Immunoprecipitated complexes were subjected to SDS-PAGE
on 7.5% acrylamide gels followed by electroblooting onto nitro-
cellulose using a Hoefer semidry blotting system. Blots were
blocked by incubation for 30 min at room temperature in TBS
with 4% nonfat dry milk. Blots were then washed twice for 10
min with TBST (TBS with 0.05% Tween 20) and incubated with the mAb 12CA5 diluted 1:10,000 in TBST for 1.5 hr. After two washes with TBST, blots were incubated for 1 hr with peroxide-linked secondary antibody (Amersham) diluted 1:10,000 in TBST. After two final washes with TBST, blots were developed using the Enhanced chemiluminescence detection kit (Amersham).

β-Galactosidase and catalase assays

HSE-bearing reporter plasmids were pB11 (carries HSP26/SSA1/ CYC1) chimeric promoter, Sussek and Lindquist 1990 and CS11 (carries four synthetic HSEs fused to CYC1 promoter), which express E. coli lacZ in response to heat shock [gift of S. Lindquist, University of Chicago, IL]. Cells bearing either of these plasmids were grown at 33°C to ~A600 = 1.0 in the presence or absence of 0.5 mM potassium acetate, and heat-shocked at 39°C by diluting 1:1 in medium prewarmed to 55°C. After 30 min at 39°, the temperature of the culture was reduced to 23°C for 1.5 hr to allow expression of β-galactosidase (and catalase) without further stress. The temperature was reduced to protect PKC1 pathway mutants from thermal killing. Cell extracts were made and β-galactosidase assays were done with 15 ~tg of these plasmids were grown at 23~ to ~A60 o = 1.0 in the pres-

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