

The Hsp70 peptide-binding domain determines the interaction of the ATPase domain with Tim44 in mitochondria

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Ssc1, a molecular chaperone of the Hsp70 family, drives preprotein import into the mitochondrial matrix by a specific interaction with the translocase component Tim44. Two other mitochondrial Hsp70s, Ssc3 (Ecm10) and Ssq1, show high sequence homology to Ssc1 but fail to replace Ssc1 *in vivo*, possibly due to their inability to interact with Tim44. We analyzed the structural basis of the Tim44 interaction by the construction of chimeric Hsp70 proteins. The ATPase domains of all three mitochondrial Hsp70s were shown to bind to Tim44, supporting the active motor model for the Hsp70 mechanism during preprotein translocation. The peptide-binding domain of Ssc1 sustained binding of Tim44, while the peptide-binding domains of Ssc3 and Ssq1 exerted a negative effect on the interaction of the ATPase domains with Tim44. A mutation in the peptide-binding domain of Ssc1 resulted in a similar negative effect not only on the ATPase domain of Ssc1, but also of Ssq1 and Ssc3. Hence, the determination of a crucial Hsp70 function via the peptide-binding domain suggests a new regulatory principle for Hsp70 domain cooperation.

Keywords: Hsp70/protein interaction/mitochondrial/*Saccharomyces cerevisiae*/Tim44

Introduction

Molecular chaperones of the 70 kDa class (Hsp70) perform critical functions in many cellular processes (Craig *et al.*, 1994; Hartl, 1996). Hsp70 chaperones promote protein folding and assembly by binding to unfolded or partially folded polypeptide chains. In *Saccharomyces cerevisiae*, 14 Hsp70 homologs have been identified. Despite their functional diversity, proteins of the Hsp70 family have retained a remarkably high sequence conservation and exhibit a well-conserved structural composition (Bukau and Horwich, 1998). The N-terminal part forms a 44 kDa domain exhibiting ATPase activity. Hsp70 proteins can switch between two main conformational states depending on the nucleotide bound (McCarthy *et al.*, 1995). The nucleotide state of the ATPase domain controls the affinity of the C-terminal part for substrate polypeptides. The C-terminal part can be divided into the 18 kDa peptide-binding domain (PBD) and a small 10 kDa variable domain at the extreme

C-terminus. In some types of eukaryotic Hsp70 proteins the variable domain has been shown to act as a binding site for co-factors that modulate their activity (Demand *et al.*, 1998; Horton *et al.*, 2001). When ATP is bound the affinity of the PBD for the substrate polypeptide is low, with a high on and off rate. Hydrolysis of the nucleotide leads to conformational changes in both domains and stabilizes the interaction with the substrate. Two types of co-factors stimulate the cellular activity of Hsp70 family members (Cyr *et al.*, 1994; Rassow *et al.*, 1995). Proteins of the DnaJ family mainly regulate substrate interaction while proteins of the GrpE family catalyze nucleotide exchange.

Specific interactions with specialized partner proteins but also the subcellular localization determine the functional specificity of the different Hsp70s. In mitochondria from *S.cerevisiae* three different Hsp70 molecules have been identified, Ssc1, Ecm10 and Ssq1, all located as soluble proteins in the matrix (Schilke *et al.*, 1996; Baumann *et al.*, 2000). The essential protein Ssc1, the most abundant mitochondrial Hsp70, plays a major role in the membrane translocation of cytosolic preproteins through the mitochondrial inner membrane into the matrix and in subsequent folding reactions. Together with the proteins Tim44 and Mge1, Ssc1 forms an essential import motor complex that is responsible for the coupling of ATP hydrolysis to polypeptide translocation (Matouschek *et al.*, 2000; Strub *et al.*, 2000). By the ATP-regulated interaction with Tim44, Ssc1 is recruited to the import site at the inner membrane pore, thus favoring binding of Ssc1 to the incoming preprotein. A functional interaction between Ssc1 and Tim44 is indispensable for the full efficiency of the preprotein translocation reaction. The introduction of mutations in both Ssc1 and Tim44 that destabilize or abolish their interaction result in significant preprotein import defects (Voos *et al.*, 1996; Merlin *et al.*, 1999; Milisav *et al.*, 2001). In particular, the import of preproteins containing stable folded domains is affected by a defective Ssc1–Tim44 interaction. Here, Ssc1 is not able to exert an inward-directed translocation force on the polypeptide in transit that would lead to the active unfolding of preprotein domains exposed on the cytosolic face of the mitochondrial membranes (Voisine *et al.*, 1999; Lim *et al.*, 2001).

In contrast to Ssc1, Ssq1 and Ecm10 seem to be less important for the mitochondrial metabolism. Mutations in Ssq1 have only minor effects on cellular growth, and no preprotein import defect has been observed (Schilke *et al.*, 1996); however, the maturation of the yeast frataxin homolog is delayed and mitochondria exhibit a significantly increased iron content (Knight *et al.*, 1998). Both genetic and biochemical data seem to indicate that Ssq1 plays a role in the mitochondrial iron metabolism and Fe/S cluster assembly (Craig *et al.*, 1999; Lutz *et al.*, 2001), although its direct cellular role is not fully understood.

Although there is extensive sequence overlap Ssq1 is not able to take over Ssc1 function (Schilke *et al.*, 1996). A possible reason is the inability of Ssq1 to bind to the preprotein translocase component Tim44 (Schmidt *et al.*, 2001). The protein Ecm10, encoded by the open reading frame YEL030w, was initially classified in a group together with the mitochondrial Hsp70 Ssc1 due to the extremely high sequence similarity and a potential mitochondrial targeting sequence (Rassow *et al.*, 1997). Although *ECM10* was also described as being involved in cell wall formation (Lussier *et al.*, 1997), recent evidence demonstrated clearly a mitochondrial localization rendering a function in cell wall biogenesis highly unlikely (Baumann *et al.*, 2000). Based on these results we recommend using the name Ssc3 instead of Ecm10. Despite their functional differences, both Ssq1 and Ssc3 (Ecm10) show typical chaperone properties and also interact specifically with the Hsp70 nucleotide exchange factor Mge1 (Baumann *et al.*, 2000; Schmidt *et al.*, 2001). However, the functional relevance of Ssc3 (Ecm10) under normal growth conditions is unclear, since it is expressed at extremely low levels and no phenotype was observed in a deletion mutant.

These observations raised the question of whether the closely related *SSC3* is capable of complementing a *ssc1Δ* mutation when its expression levels were raised considerably. Since we found that Ssc3 could not replace Ssc1, we asked which structural properties are responsible for the distinct functional differences between the mitochondrial Hsp70s. We investigated the relative influence of the Hsp70 domains on the interaction with the essential partner protein Tim44 by the construction of chimeric proteins consisting of different combinations of ATPase, peptide-binding and variable C-terminal domains derived from the three mitochondrial Hsp70s. We found that the ATPase domains of all three Hsp70s are principally able to bind to Tim44, while the PBDs of Ssq1 and Ssc3 can exert a negative influence on the ATPase domain, which results in the inability of the respective full-length proteins to interact with Tim44.

Results

The mitochondrial Hsp70, Ssc3 (Ecm10), cannot replace the essential function of the related chaperone Ssc1

Of the three species of Hsp70 proteins in the mitochondrial matrix, Ssc1 and Ssc3 (Ecm10) exhibit an exceptionally high degree of sequence conservation. Ssc3 and Ssc1 share 82% identical amino acids, slightly more in the ATPase domain than in the PBD. The other mitochondrial Hsp70, Ssq1, is less well conserved and has an overall identity of 52%. It has been shown that Ssq1 is not able to take over the Ssc1 function *in vivo* even if expressed at similar levels as Ssc1, but the much higher sequence conservation between Ssc3 and Ssc1 indicated an overlapping function. To determine the functional relationship between Ssc3 and Ssc1 *in vivo* we tested whether Ssc3 might be able to complement the lethal effect of a *ssc1Δ* mutation. To increase the expression level of Ssc3, we inserted the *SSC3* (*ECM10*) gene including the promoter and downstream region into a high copy number vector and performed an *in vivo* complementation assay with an *ssc1Δ* mutant

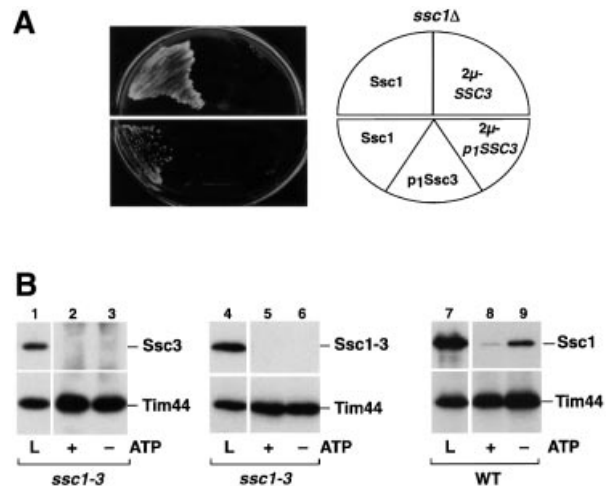


Fig. 1. Ssc3 (Ecm10) is not able to functionally replace Ssc1. (A) Overproduction of Ssc3 cannot rescue the lethal phenotype of a mutant lacking Ssc1. Plasmids expressing *SSC1* (pASC07) or the *SSC3* gene in an overexpression vector (pASE10) were tested for complementation of the *ssc1Δ* mutation (upper panel). In addition, the mature Ssc3 protein with the pre-sequence and the promoter of Ssc1 (p₁Ssc3) was expressed in a low copy (pASE08) and high copy (pNZ46) vector. (B) Ssc3 does not interact with the essential Ssc1 partner protein Tim44. Interaction of Ssc3 with Tim44 was assayed by co-immunoprecipitation with antibodies against Tim44 from mitochondrial lysates overproducing Ssc3 in *ssc1-3* mutant cells in the presence (+) or absence (-) of ATP as described in Materials and methods. Cells were pre-incubated for 15 min at 37°C to induce the mutant phenotype prior to lysis. Five percent of the lysed mitochondria were applied as a control (L). Precipitated proteins were separated by SDS-PAGE and the presence of Ssc3 (lanes 1–3) or Ssc1 (lanes 4–9) was detected by western blot.

strain. No colonies were detected with cells containing the high copy plasmid expressing Ssc3, whereas the control cells expressing Ssc1 grew normally (Figure 1A, upper panel). However, a biochemical characterization of mitochondria isolated from these cells revealed that the amount of Ssc3 was still lower than the amount of Ssc1 in wild-type mitochondria. When we tested the mitochondrial import efficiency *in vitro* it became apparent that a low intrinsic import rate of the Ssc3 precursor protein was the reason for the low mitochondrial abundance (data not shown). To further raise the mitochondrial levels of Ssc3 we exchanged the endogenous pre-sequence of Ssc3 for the highly efficient pre-sequence of Ssc1 without altering the cleavage site for the matrix processing peptidase (MPP) and additionally replaced the endogenous promoter by the promoter region of *SSC1*. The resulting protein construct, p₁Ssc3, was efficiently imported and processed in isolated mitochondria. However, despite mitochondrial protein amounts similar to Ssc1 (data not shown), Ssc3 was not able to complement the lethal phenotype of the *ssc1* null mutation (Figure 1A, lower panel). Even when expressed from a high copy 2μ vector, p₁Ssc3 did not take over the function of Ssc1 (Figure 1A, lower panel). In summary, Ssc3 cannot complement the essential function of Ssc1 despite their high sequence similarity.

Both Ssc3 and Ssq1 show no interaction with the essential partner protein Tim44

We wished to determine what property of Ssc3 might be responsible for the observed inability to take over the function of the related Ssc1. The import of cytosolic

preproteins into the mitochondrial matrix is considered to be the essential function of Ssc1, requiring an efficient interaction with the inner membrane translocase component Tim44. Hence, we tested whether Ssc3 is able to interact specifically with Tim44 by performing co-immunoprecipitation experiments *in organello*. Detergent lysates of mitochondria isolated from the temperature-sensitive mutant strain *ssc1-3* overexpressing *p1SSC3* were prepared under native buffer conditions. The temperature-sensitive mutant protein Ssc1-3 is completely inactivated under non-permissive conditions, and unable to bind to Tim44 (Voos *et al.*, 1996; Figure 1B, lanes 4–6); therefore, a possible competition between Ssc1 and Ssc3 for binding sites at Tim44 could be excluded. However, no signal for Ssc3 could be detected when Tim44 was precipitated from the mitochondrial lysates with affinity-purified antibodies in either the presence or absence of nucleotides (Figure 1B, lanes 1–3). As control, Ssc1 co-precipitated efficiently with Tim44 in an ATP-sensitive manner, indicating a specific and functional interaction (Figure 1B, lanes 7–9). In summary, Ssc3 failed to interact with the Hsp70 partner protein Tim44, and this is a possible reason for the inability of Ssc3 to complement the deletion of Ssc1.

The conserved structural organization of Hsp70 family members allowed us to determine the properties of Hsp70s that are responsible for the interaction with Tim44 using a domain-swap approach. In previous experiments we were able to demonstrate that the ATPase domain of Ssc1 is crucial for interaction with Tim44 (Krimmer *et al.*, 2000). We asked whether it is possible to convert the non-binding proteins Ssc3 and Ssq1 into proteins capable of an interaction with Tim44 by exchanging their ATPase domain for that of Ssc1. For this purpose we constructed chimeric proteins that contained the ATPase domain of Ssc1 (A_1) fused to the PBD and variable domains of Ssc3 (P_3V_3) or Ssq1 (P_QV_Q) (Figure 2A). The resulting chimeric proteins $A_1P_3V_3$ and $A_1P_QV_Q$ were tested for their ability to form a complex with Tim44. In order to be able to detect minor amounts of co-precipitated proteins we performed the analysis with radiolabeled precursor proteins that were imported into isolated mitochondria. After stopping the import reaction and removal of excess preprotein, mitochondria were lysed and the immunoprecipitation with Tim44 antibodies in the presence or absence of ATP was performed. Efficient binding to Tim44 was detected with the imported Ssc1. Up to 3.5% of the imported and processed preprotein was found bound to Tim44 (Figure 2B, lane 3), reflecting the relative amounts of Tim44 and Hsp70 in the matrix. Since endogenous Ssc1 is in high excess over Tim44, only a small part of the total amount of Ssc1 is found in a complex with Tim44 in mitochondria. Similar to the co-immunoprecipitation experiments *in organello*, the Tim44–Ssc1 complex was dissociated in the presence of ATP (Figure 2B, lane 2), demonstrating that the imported proteins are functional. Unexpectedly, neither $A_1P_3V_3$ (Figure 2B, lanes 11 and 12) nor $A_1P_QV_Q$ (Figure 2B, lanes 14 and 15) was found in complex with Tim44, even though they contain the functional ATPase domain of Ssc1. Here the chimeric constructs behaved indistinguishably from wild-type Ssc3 and Ssq1 (Figure 2B, lanes 4–9).

To ascertain that the surprising inability to interact with Tim44 was not a consequence of the *in vitro* import assay

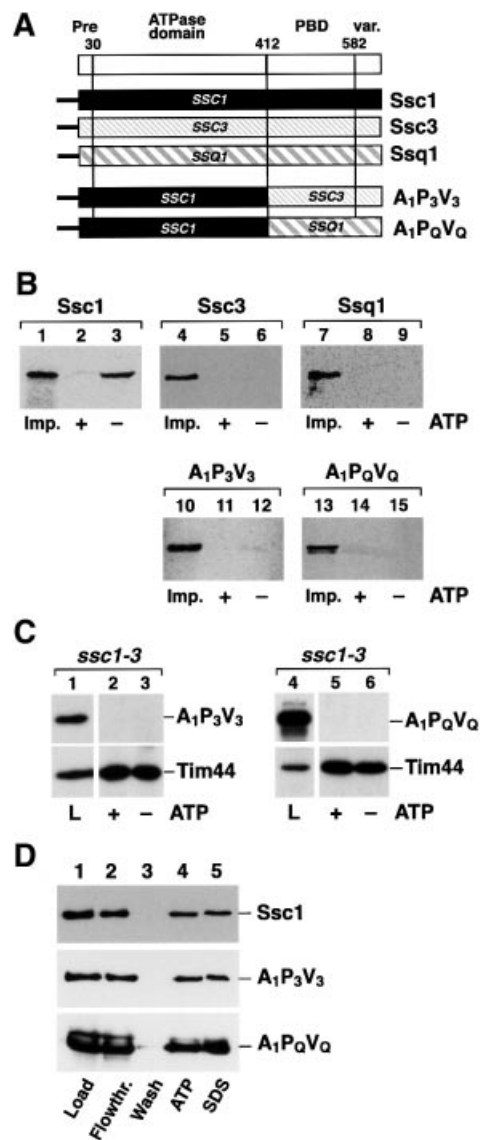


Fig. 2. Chimeric Hsp70 constructs containing the ATPase domain of Ssc1 fused with the PBD and C-terminus of Ssc3 or Ssq1 do not interact with Tim44. (A) Scheme of the used Hsp70 constructs. Indicated are the N-terminal pre-sequence (Pre), the ATPase domain, the peptide-binding domain (PBD) and the C-terminal variable domain (var.). The numbers indicate the first amino acid of each Ssc1 subdomain. (B) The wild-type proteins or the chimeric constructs $A_1P_3V_3$ and $A_1P_QV_Q$ were imported into isolated mitochondria as described in Materials and Methods. Mitochondria were lysed in the presence (+) or absence (-) of ATP. Five percent of the lysis fraction was taken as import control (Imp.). Proteins interacting with Tim44 were isolated by co-immunoprecipitation and detected by SDS-PAGE and autoradiography. (C) The constructs $A_1P_3V_3$ and $A_1P_QV_Q$ do not bind to Tim44 *in vivo*. $A_1P_3V_3$ (pASC31) and $A_1P_QV_Q$ (pASC26) were expressed in *ssc1-3* mutant cells and co-immunoprecipitation analysis was performed as described in (A). Proteins were detected by western blot using antibodies raised against the C-terminal parts of Ssc3 or Ssq1. Five percent of the mitochondrial lysates were included as control (L). (D) ATPase domains of $A_1P_3V_3$ and $A_1P_QV_Q$ are functional. ATP-agarose binding assays were performed under non-permissive conditions with mitochondrial lysates expressing the constructs $A_1P_3V_3$ and $A_1P_QV_Q$ in *ssc1-3*. Samples were split and bound proteins were eluted by incubation in lysis buffer that contained 1 mM ATP or in SDS sample buffer, respectively. Ten percent lysate (Load) and flowthrough (Flowthr.), and 25% of the final wash (Wash) fractions were included. Proteins were detected by western blot using antibodies recognizing Ssc1 (first panel), the C-terminus of Ssc3 (second panel) or Ssq1 (third panel).

procedure, we repeated the co-immunoprecipitation experiments with mitochondrial extracts from cells that expressed the chimeric proteins *in vivo*. Immunoprecipitations were performed in the *ssc1-3* strain background in order to avoid an interference of endogenous Ssc1. As was observed with the radiolabeled imported versions the chimeric proteins A₁P₃V₃ and A₁P_QV_Q were not found in a complex with Tim44 under the *in organello* conditions (Figure 2C, lanes 1–6). To confirm that the ATPase domain of Ssc1 in the heterologous protein context was functional, we tested the ability of A₁P₃V₃ and A₁P_QV_Q to bind nucleotides. We performed ATP-agarose binding experiments with mitochondrial extracts of either wild-type Ssc1 or of A₁P₃V₃ and A₁P_QV_Q expressed in *ssc1-3*. Owing to its mutation in the ATPase domain, Ssc1-3 is not able to interact with nucleotides under non-permissive conditions (von Ahsen *et al.*, 1995). The mitochondrial extracts were incubated with ATP-agarose and bound proteins were eluted either by an incubation in buffer containing ATP or in SDS sample buffer. Similar to the wild-type Ssc1 (Figure 2D, upper panel), both constructs, A₁P₃V₃ (Figure 2D, middle panel) and A₁P_QV_Q (Figure 2D, lower panel), showed efficient and specific binding to ATP (Figure 2D, lane 5) and could be released from the agarose matrix upon addition of excess ATP (Figure 2D, lane 4). In addition, all constructs also showed an efficient and specific interaction with the nucleotide exchange factor Mge1 (data not shown). The binding to ATP and the physical interaction with Mge1 are consistent with a functional ATPase domain in the chimeric protein background of the constructs A₁P₃V₃ and A₁P_QV_Q.

The Ssc1 PBD enables Ssc3 and Ssq1 to interact with Tim44

Since the presence of the ATPase domain of Ssc1 does not restore the ability to interact with Tim44, the possibility remained that the Tim44 interaction is governed by the PBD and variable domain. To determine the influence of these domains, we exchanged the ATPase domain of the Tim44-binding protein Ssc1 by the ATPase domains of the non-binding proteins Ssc3 or Ssq1. The resulting proteins A₃P₁V₁ and A_QP₁V₁ both contained the full PBD (P₁) and variable domain (V₁) of Ssc1, but the ATPase domains of Ssc3 (A₃) or Ssq1 (A_Q) (Figure 3A). To ensure efficient import *in vitro*, all constructs also contained the pre-sequence of Ssc1. Both chimeric constructs were efficiently imported into isolated mitochondria dependent on the inner membrane potential and showed correct maturation in the matrix (data not shown). We then analyzed the interaction of A₃P₁V₁ and A_QP₁V₁ with Tim44 by co-immunoprecipitation experiments after import. Surprisingly, both, A₃P₁V₁ and A_QP₁V₁ were detected in a complex with Tim44. The efficiency of complex formation of both A₃P₁V₁ (Figure 3B, lane 6) and A_QP₁V₁ (Figure 3B, lane 9) was similar to that observed with full-length Ssc1 (Figure 3B, lane 3). As in the case of Ssc1, the complex of A_QP₁V₁ and Tim44 dissociated in the presence of ATP (Figure 3B, lane 8), demonstrating that the ATPase domain of Ssq1 is functional and the protein interaction is specific. Compared with wild-type Ssc1 (Figure 3B, lane 2) the complex of A₃P₁V₁ with Tim44 showed a lower sensitivity to ATP (Figure 3B, lane 5), but a significant amount of the bound protein was still released by ATP.

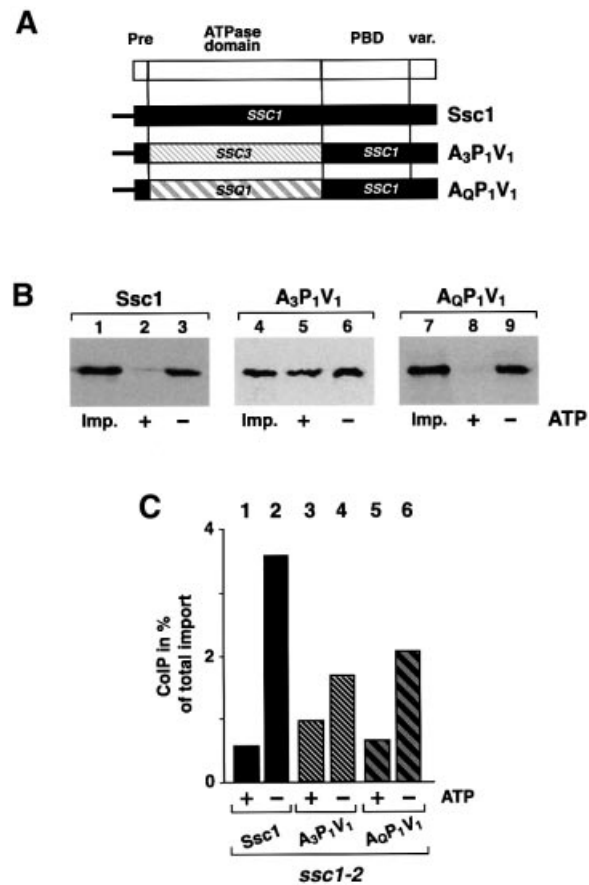


Fig. 3. Chimeric constructs containing the ATPase domains of the different Hsp70 homologs fused to the PBD and C-terminus of Ssc1 can be efficiently co-precipitated with Tim44. (A) Scheme of the chimeric constructs A₃P₁V₁ and A_QP₁V₁. (B) Tim44 interaction of A₃P₁V₁, A_QP₁V₁ and Ssc1. Binding assays were performed after preprotein import by co-immunoprecipitation in the presence (+) or absence (-) of ATP as described. Five percent of total mitochondria was added as import control (Imp.). (C) Binding of A₃P₁V₁ and A_QP₁V₁ to Tim44 is direct and independent of functional Ssc1. Ssc1, A₃P₁V₁ and A_QP₁V₁ were imported into isolated *ssc1-2* mutant mitochondria (PK81) under permissive conditions as described in Materials and methods. Mitochondria were heat-treated for 15 min at 37°C to induce the phenotype of Ssc1-2. Proteins bound to Tim44 were isolated by co-immunoprecipitation in the presence (+) or absence (-) of ATP. The efficiency of the co-immunoprecipitation is given as a percentage of the total imported protein.

The possibility remained that the occurrence of the chimeric proteins in the precipitation with Tim44 was due to a formation of heterodimers with endogenous Ssc1. In order to exclude this we performed the binding assays in mitochondria from the mutant strain *ssc1-2*. Under the chosen conditions Ssc1-2 is not able to form a stable complex with Tim44, while it retains the ability to import denatured preproteins (Gambill *et al.*, 1993; von Ahsen *et al.*, 1995). After import of A₃P₁V₁ or A_QP₁V₁ into *ssc1-2* mitochondria both chimeric proteins (Figure 3C, lanes 4 and 6) were found in complex with Tim44. Essentially no difference from the situation in wild-type mitochondria was observed. Again, the interaction of A₃P₁V₁ was less sensitive to ATP (Figure 3C, lane 3), whereas A_QP₁V₁ showed the typical sensitivity of the Hsp70-Tim44 interaction to nucleotides (Figure 3C, lane 5). As a control, wild-type Ssc1 was imported into *ssc1-2* mitochondria, showing a specific and efficient interaction

with Tim44, as expected (Figure 3C, lanes 1 and 2). Taken together, it is highly unlikely that the observed interaction of the imported proteins with Tim44 did occur via a dimerization with endogenous Ssc1. The co-precipitation experiments therefore indicate clearly that chimeric proteins containing the ATPase domains of Ssc3 and Ssq1 and the C-terminal domains of Ssc1 can interact specifically with Tim44.

The interaction of chimeric Hsp70 constructs with Tim44 does not restore preprotein import

Since the chimeric proteins $A_3P_1V_1$ and $A_QP_1V_1$ were able to bind to the preprotein translocase component Tim44, we tested whether the constructs were able to complement the growth defect of *ssc1Δ* mutants *in vivo*. The constructs were expressed under control of the *SSC1* promoter and growth in the absence of functional Ssc1 was tested as described above. Surprisingly, neither construct was able to restore growth of the *ssc1Δ* cells (Figure 4A). To analyze the phenotype in more detail, we expressed the chimeric constructs in the *ssc1-3* temperature-sensitive mutant. Mitochondria were isolated under permissive conditions and preprotein import was tested after a short pre-incubation at 37°C to inactivate the endogenous Ssc1-3. Interestingly, both chimeric constructs exhibited a pronounced import defect of the reporter preprotein Su9-DHFR (Figure 4B). Mitochondria containing the chimeric constructs behaved virtually indistinguishably from those derived from the parent strain *ssc1-3*. The significant preprotein translocation defect shown by both chimeric proteins despite a specific interaction with Tim44 is the most likely reason for their inability to replace Ssc1.

To elucidate which particular functional property of the chimeric proteins is responsible for the defect, we assayed their basic biochemical behavior, substrate binding and nucleotide interaction. As substrate we chose carboxy-methylated α -lactalbumin (CMLA), a typical unfolded model substrate for Hsp70 proteins. We incubated $A_QP_1V_1$ that was expressed in *ssc1-3* with immobilized CMLA and eluted the bound proteins either in the presence of ATP or with SDS sample buffer. Interestingly, the construct $A_QP_1V_1$ showed significant binding to the substrate CMLA, indicating a functional PBD (Figure 4C, lane 4, upper panel). However, in contrast to wild-type Ssc1 (Figure 4C, lower panel), no release from the immobilized substrate in the presence of ATP was observed (Figure 4C, lane 3, upper panel), although the Tim44-binding assays showed full reactivity of the ATPase domain in $A_QP_1V_1$ to ATP. Ssc1-3, with its inability to bind ATP, shows a comparable substrate binding behavior to the construct $A_QP_1V_1$ (Figure 4C, middle panel), indicating that the information of the nucleotide state of the ATPase domain can not be propagated to the PBD and alter the substrate affinity. Since Ssc1-3 does not show any binding activity to ATP (Figure 4D, lanes 4 and 5, lower panel), we were able to analyze the nucleotide interaction of the construct $A_3P_1V_1$ by binding assays with ATP-agarose. $A_3P_1V_1$ was able to bind ATP-agarose but, in contrast to Ssc1, was not released in the presence of excess ATP (Figure 4D, lanes 4 and 5, upper panel). The chimeric construct behaved very similar to the full-length Ssc3 (Figure 4D, lanes 4 and 5, middle panel), indicating a possible lower nucleotide exchange rate of the A_3 domain. Hence, both chimeric

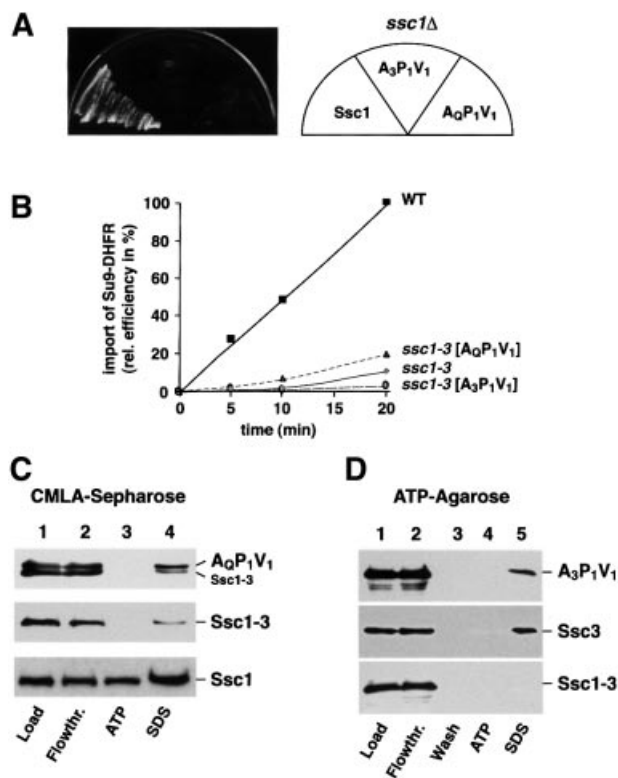


Fig. 4. The chimeric constructs $A_3P_1V_1$ and $A_QP_1V_1$ are not able to replace Ssc1. (A) $A_3P_1V_1$ and $A_QP_1V_1$ cannot rescue the lethal phenotype of a mutant lacking Ssc1. Complementation assays with strain WVY37 expressing Ssc1 (pASC07), $A_3P_1V_1$ (pASC25) and $A_QP_1V_1$ (pASC11) were essentially performed as described in Figure 1A. (B) $A_3P_1V_1$ and $A_QP_1V_1$ cannot complement the import defect of a *ssc1* mutant. Mitochondria isolated from the strains ASYC32 (WT) or *ssc1-3* expressing $A_3P_1V_1$ or $A_QP_1V_1$ were incubated at 37°C for 15 min to induce the mutant phenotype and the model protein Su9(70)-DHFR was imported for the indicated times. The import reactions were quantified and the amount imported into wild-type after 20 min was set to 100%. (C) Substrate proteins bound to $A_QP_1V_1$ are not released by addition of ATP. Mitochondria from strain *ssc1-3* expressing the construct $A_QP_1V_1$ (upper panel), a control plasmid (middle panel) and wild-type mitochondria (lower panel) were lysed under non-permissive conditions and binding assays with CMLA-Sepharose were performed as described. Samples were split and proteins were eluted with lysis buffer containing 3 mM ATP or SDS sample buffer. Load (10%), and flowthrough (Flowthr.; 10%) were included. Bound proteins were detected by western blot using antibodies recognizing the C-terminus of Ssc1. (D) Ssc3 and $A_3P_1C_1$ are not able to release bound ATP. *ssc1-3* mitochondria expressing the constructs p_1 Ssc3 (pASE08; upper panel), $A_3P_1V_1$ (middle panel) or a control plasmid (lower panel) were lysed under non-permissive conditions and binding assays with ATP-agarose were performed as described. Samples were split and proteins were eluted with lysis buffer containing 1 mM ATP or SDS sample buffer. Load (10%), flowthrough (Flowthr.; 10%) and final wash (25%) were included. Bound proteins were detected by western blot using antibodies recognizing the C-terminus of Ssc3 (upper panel) or Ssc1 (middle and lower panels).

constructs $A_3P_1V_1$ and $A_QP_1V_1$ have severe biochemical defects concerning nucleotide reactivity or interdomain communication, resulting in a significant preprotein import defect and eventually in a pronounced growth inhibition *in vivo*.

Tim44 is not able to bind to the C-terminal part of any mitochondrial Hsp70

The observed interaction of $A_3P_1V_1$ and $A_QP_1V_1$ with Tim44 raised the question of whether the PBD derived

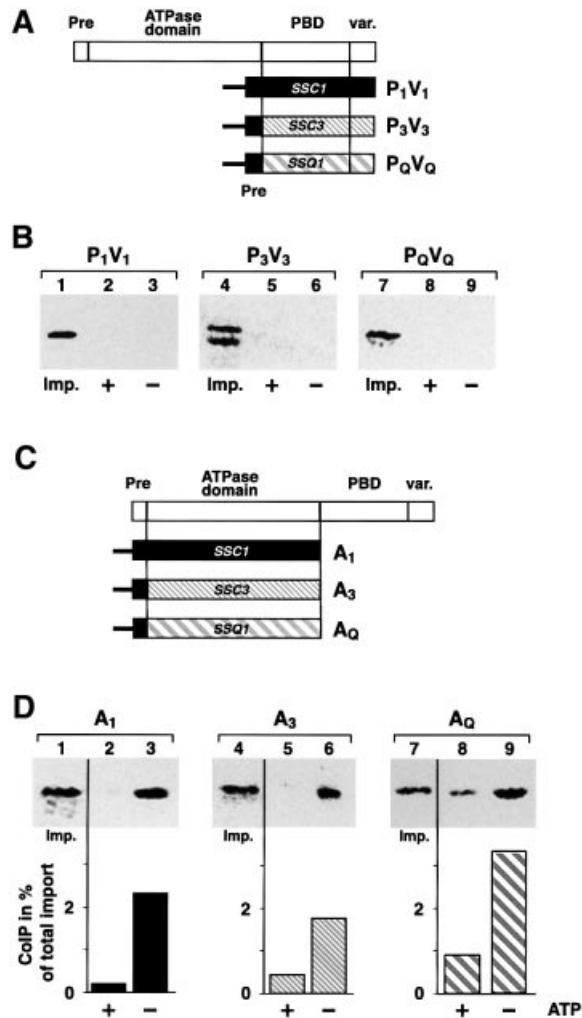


Fig. 5. The PBDs of all three Hsp70s do not interact with Tim44, while the ATPase domains bind specifically to Tim44. (A) Scheme of the constructs P₁V₁, P₃V₃ and P_QV_Q. The pre-sequence of Ssc1 was fused to the PBDs of either Ssc1, Ssc3 or Ssq1. (B) *In vitro* import and Tim44 co-immunoprecipitation of P₁V₁, P₃V₃, P_QV_Q and Ssc1. After import into mitochondria, binding to Tim44 was analyzed in the presence (+) or absence (-) of ATP. Five percent of mitochondria were added as control (Imp.). (C) Scheme of the ATPase domain constructs A₁, A₃ and A_Q. (D) Co-immunoprecipitation analysis of the single ATPase domains A₁, A₃ and A_Q with Tim44. Immunoprecipitations with Tim44 antibodies in the presence (+) or absence (-) of ATP after import into mitochondria were performed as described. As a control, 2% of the mitochondrial extracts after import were applied (Imp.). Precipitation efficiency is given as a percentage of the total imported protein.

from Ssc1 rather than the ATPase domain would be able to interact with Tim44. The pre-sequence of Ssc1 was fused to the PBD and the adjacent variable domain of Ssc1, Ssc3 or Ssq1 (Figure 5A). The constructs P₁V₁, P₃V₃ and P_QV_Q were imported into isolated mitochondria. Although the processing of P₃V₃ was not as efficient as P₁V₁ and P_QV_Q (Figure 5B, lane 4), import into mitochondria was specific, as shown by its full protection against protease digestion. When the interaction with Tim44 was analyzed by co-immunoprecipitation, no binding was observed for P₁V₁, P₃V₃ and P_QV_Q (Figure 5B), either in the presence or absence of ATP. Similar results were obtained with the PBD and variable domain of Ssc1 that were expressed *in vivo* (data not shown). Since the C-terminal domains did

not bind to Tim44, we tested the ATPase domains for their binding behavior. We constructed truncated versions of Ssc1, Ssc3 and Ssq1 that contained the pre-sequence of Ssc1 fused to the respective ATPase domains, resulting in the constructs A₁, A₃ and A_Q (Figure 5C). After import into isolated mitochondria, we observed that the ATPase domain of Ssc1 (A₁) bound to Tim44 in the absence of ATP, as expected (Figure 5D, lane 3). Interestingly, we found that the ATPase domains of the other two mitochondrial Hsp70s, A₃ and A_Q, were also able to interact efficiently with Tim44 (Figure 5D, lanes 6 and 9). These results confirmed that the observed binding of the chimeric constructs A₃P₁V₁ and A_QP₁V₁ (Figure 3B) indeed occurred via the ATPase domain. For all three ATPase domains, between 2 and 3.5% of the imported protein was found in complex with Tim44 (Figure 5D, lanes 3, 6 and 9), a similar value as was found for full-length Ssc1 (Figure 3B). The sensitivity of the Tim44 complex of all three proteins to the presence of ATP indicated that the interaction is specific and functional (Figure 5D, lanes 2, 5 and 8). In summary, the ATPase domains of all three mitochondrial Hsp70s were sufficient to bind to Tim44 with characteristics comparable to the full-length Ssc1. The interaction was also observed in the context of the full-length protein as long it contains the PBD and variable domains of Ssc1. In contrast, no interaction was found for the wild-type or chimeric proteins that contained the C-terminal domains of either Ssq1 or Ssc3. We conclude that the C-terminal domains of Ssc3 and Ssq1 exert a negative influence on the binding behavior of their corresponding ATPase domains.

The variable domain of Ssc1, but not of Ssc3 and Ssq1, has a stabilizing influence on the Tim44 interaction

We then tried to dissect the relative influence of the PBD and the variable domain on the Tim44 interaction. We exchanged the variable domain of Ssc1 by the corresponding segments of Ssc3 or Ssq1, resulting in the proteins A₁P₁V₃ and A₁P₁V_Q (Figure 6A). The constructs were imported into isolated mitochondria and processed to the mature protein (data not shown). Both A₁P₁V₃ and A₁P₁V_Q were able to bind to Tim44 in an ATP-sensitive fashion (Figure 6B, lanes 1–3, middle and lower panel). In the construct A₁P₁V₃ the C-terminus of Ssc3 had no influence on Tim44 binding efficiency compared with the full-length Ssc1 (Figure 6C, column 2 and 4). The construct A₁P₁V_Q also showed significant and ATP-sensitive binding to Tim44 (Figure 6B, lanes 2 and 3, lower panel), but its interaction was less efficient than wild-type Ssc1 (Figure 6C, columns 2 and 6). The truncated protein A₁P₁, which did not contain the variable domain, was able to bind to Tim44 (Figure 6B, lane 6, upper panel), but with decreased efficiency. The interaction of the shortened construct A₁P₁Q was completely abolished, while the construct A₁P₁ was reduced to almost background levels (Figure 6B, lane 6, middle and lower panels). Interestingly, the binding efficiency of A₁P₁ was lower than wild type, but almost identical to the value obtained with A₁P₁V_Q. This indicates that the variable domain of Ssc1 could contribute to the overall efficiency of the Tim44 interaction. The variable domain of Ssc3 was able to exercise this stabilizing effect, but the correspond-

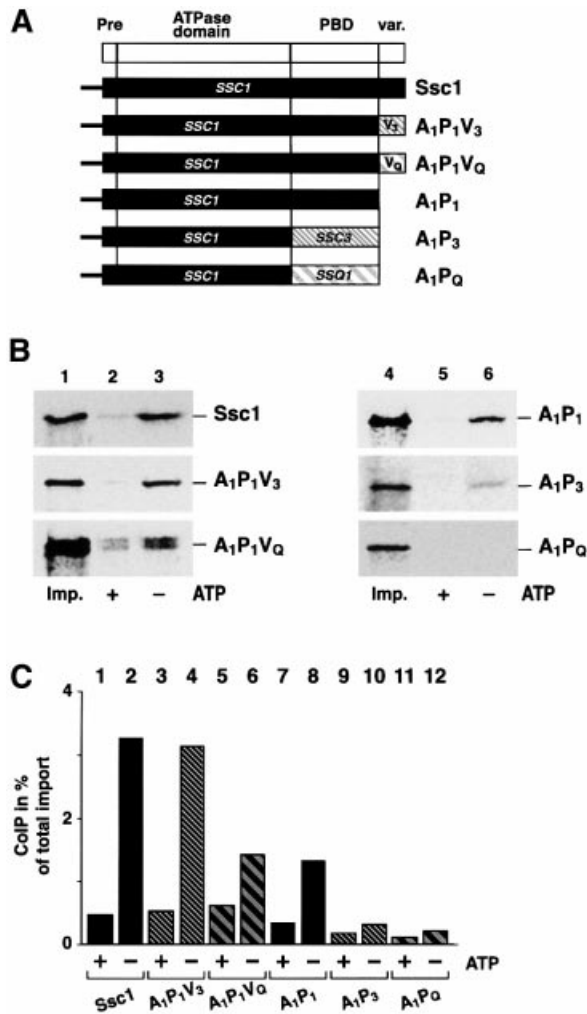


Fig. 6. The variable C-terminal domains of Ssc1 and Ssc3 have a stabilizing effect on the interaction with Tim44. (A) Scheme of the Hsp70 constructs. The variable C-terminal domain of Ssc1 was replaced by the domains of Ssc3 or Ssq1 (A₁P₁V₃ and A₁P₁V_Q). The constructs A₁P₃ and A₁P_Q were obtained by replacing the PBD and C-terminus of Ssc1 with the PBD of Ssc3 or Ssq1. (B) Interaction with Tim44. Import into isolated mitochondria and co-immunoprecipitation with Tim44 antibodies in the presence (+) or absence (-) of ATP was performed as described. As a control, 5% of the mitochondrial lysates after the import reactions were applied (Imp.). (C) Relative efficiencies of the interaction of Tim44 with the chimeric constructs. The co-immunoprecipitation experiments shown in (B) were quantified and binding efficiencies are given as a percentage of the total imported preprotein.

ing domain of Ssq1 was not. This effect of the variable domain was qualitatively different from the influence of the PBDs on the Tim44 interaction. In the absence of a stabilizing effect exerted by variable domains the residual binding of the truncated proteins was again abrogated by the negative influence of the PBD derived from Ssc3 or Ssq1.

A mutation in the PBD of Ssc1 abolishes Tim44 interaction

The negative influence of the PBD of Ssq1 and Ssc3 was reminiscent of the behavior of the mutant Ssc1-2 containing a point mutation in the PBD (substitution of Pro442 by serine) that alters its reactivity towards Tim44. We asked whether the same amino acid exchange would be able to

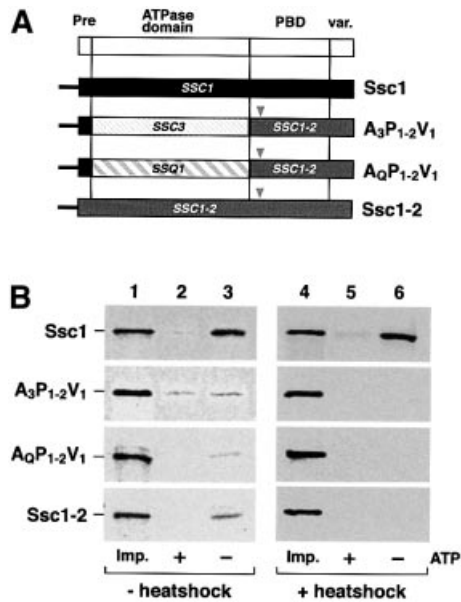


Fig. 7. A point mutation in the PBD of Ssc1 exerts a negative influence on the Tim44 interaction of all three Hsp70s. (A) Scheme of the Hsp70 constructs. The point mutation of the *ssc1-2* mutant (substitution of Pro442 by serine, arrowhead) was introduced into the constructs A₃P₁₋₂V₁ and A_QP₁₋₂V₁. (B) Co-immunoprecipitation of constructs carrying the *ssc1-2* point mutation in the PBD. After import, mitochondria were divided into two fractions and either directly lysed and subjected to co-immunoprecipitation analysis (lanes 1–3) or resuspended in import buffer and incubated for 15 min at 37°C to induce the phenotype of the mutation (lanes 4–6). Samples were divided into two fractions and mitochondria were isolated and lysed in the presence (+) or absence (-) of ATP as described. As a control, 5% of the mitochondrial lysates were applied (Imp.).

exert a similar negative influence on the binding of the ATPase domains derived from Ssc3 and Ssq1. We fused the PBD of Ssc1-2 with the ATPase domains of Ssc3 and Ssq1, resulting in the constructs A₃P₁₋₂V₁ and A_QP₁₋₂V₁ (Figure 7A). After import of the radiolabeled Ssc1, A₃P₁₋₂V₁, A_QP₁₋₂V₁ and Ssc1-2, mitochondria were either lysed directly (Figure 7B, lanes 1–3) or incubated at 37°C for 15 min before lysis (Figure 7B, lanes 4–6), and the complex formation with Tim44 was analyzed. Similar to the original mutant protein Ssc1-2, the binding of A_QP₁₋₂V₁ and A₃P₁₋₂V₁ to Tim44 was already decreased at permissive temperature (Figure 7B, lanes 2 and 3, second and third panels) compared with Ssc1 (Figure 7B, lane 3, first panel). Compared with wild-type Ssc1, all constructs carrying the *ssc1-2* mutation showed <20% binding efficiency. Under non-permissive conditions the complex formation between Tim44 and A₃P₁₋₂V₁ or A_QP₁₋₂V₁ was completely abolished (Figure 7B, lanes 5 and 6, second and third panels). In contrast, binding of wild-type Ssc1 to Tim44 was not affected (Figure 7B, lane 6, first panel). The binding behavior of the chimeric constructs containing the PBD derived from Ssc1-2 was identical to the binding defect of the original Ssc1-2 mutant protein (Figure 7B, lanes 1–6, fourth panel). Thus, the destabilizing effect of the *ssc1-2* mutation on the interaction with Tim44 can also be observed in the case of the ATPase domains of Ssc3 and Ssq1. The point mutation Pro442→Ser essentially converted the stabilizing effect of the PBD into a destabilizing effect, as was found for the PBDs of Ssc3 and Ssq1.

Discussion

In this study we addressed the question of the functional correlation between the three mitochondrial Hsp70s. We were able to show by *in vivo* complementation assays that Ssc3 cannot functionally replace the major mitochondrial Hsp70, Ssc1, despite an extremely high sequence conservation. It was previously observed that overproduction of Ssc3 could partially rescue the temperature-sensitive mutant phenotype of the *ssc1-3* mutation (Baumann *et al.*, 2000). Hence, some functional overlap between both proteins must exist, but this is not sufficient to rescue the lethal effect of a *ssc1* null mutation. As was revealed by database analysis, the *ECM10* gene, encoding Ssc3, most probably arose from an ancient gene duplication of the *SSC1* locus (Wolfe and Shields, 1997). A block of several genes from chromosome X, containing *SSC1*, is found repeated on chromosome V, resulting in the *ECM10* gene, a situation similar to the genes for two cytosolic Hsp70s, *SSB1* and *SSB2*. The other mitochondrial Hsp70, Ssq1, is also unable to complement an *ssc1* deletion strain (Schilke *et al.*, 1996), despite showing typical chaperone properties (Schmidt *et al.*, 2001). All three mitochondrial chaperones have previously been shown to interact with the Hsp70 nucleotide exchange factor Mge1 (Baumann *et al.*, 2000; Schmidt *et al.*, 2001). Thus, the deficiency of both Ssc3 and Ssq1 to bind to the inner membrane translocase component Tim44 is the most likely reason for the inability of Ssc3 and Ssq1 to take over the essential cellular function of the related Ssc1. Efficient binding to Tim44 has been shown to be indispensable for the successful membrane translocation of nuclear-encoded precursor proteins into the mitochondrial matrix, especially under *in vivo* conditions (Merlin *et al.*, 1999; Voisine *et al.*, 1999; Milisav *et al.*, 2001). This indicates that the differential interaction of the mitochondrial Hsp70s with Tim44 is a very specific property, with functional relevance.

The conserved domain structure of Hsp70 proteins provided the opportunity to investigate the structural determinants of the Tim44 interaction by examining domain exchange constructs for their ability to bind to Tim44. The results of the interaction assays are summarized in Table I. Interestingly, the constructs that contained the ATPase domain derived from Ssc1 ($A_1P_3V_3$ and $A_1P_QV_Q$) were not able to interact with Tim44, while chimeric proteins with the C-terminal domains of Ssc1 ($A_3P_1V_1$ and $A_QP_1V_1$) showed an efficient interaction. Since the ATPase domains seemed to be functional in the foreign protein context these results seemed contradictory to our previous work in which the ATPase domain of Ssc1 was determined to be the principal Tim44 binding site (Krimmer *et al.*, 2000). Very recently, a report was published that also indicated a possible interaction of Tim44 with the C-terminal domain of Ssc1 (Moro *et al.*, 2002). However, our experiments demonstrate that a specific interaction of the C-terminal domains of the mitochondrial Hsp70s (PBD and variable domain) with Tim44 is very unlikely, since the single domains (P_1V_1 , P_3V_3 and P_QV_Q) showed no binding to Tim44 at all. Most interestingly, when we analyzed the ATPase domains separately for their interaction, the ATPase domain of Ssc1 (A_1), as well as A_3 and A_Q , showed specific and

Table I. Tim44 interaction of mitochondrial Hsp70 constructs

| Hsp70 domains | / | P_1V_1 | P_3V_3 | P_QV_Q | P_1V_3 | P_1V_Q | P_1 | P_3 | P_Q |
|---------------|----|----------|----------|----------|----------|----------|-------|-------|-------|
| / | | - | - | - | | | | | |
| A_1 | ++ | ++ | - | - | ++ | + | + | - | - |
| A_3 | ++ | ++ | - | | | | | | |
| A_Q | ++ | ++ | | - | | | | | |

efficient binding to Tim44. This interaction was not been noted by Moro *et al.* (2002), probably due to the use of non-physiological buffer conditions in the binding assays that did not include potassium ions. Potassium ions have been shown to be indispensable for the full enzymatic activity of the Hsp70 ATPase domain (O'Brien and McKay, 1995; von Ahsen *et al.*, 1995).

A co-precipitation of the chimeric proteins with Tim44 that has been caused by a formation of heterodimers (Azem *et al.*, 1997) between endogenous Ssc1 and the imported constructs can be excluded by two arguments. Binding assays using *ssc1-2* mutant mitochondria that do not show binding of endogenous Ssc1-2 to Tim44 did not affect the behavior of the chimeric proteins. In addition, only trace amounts of A_1 could be precipitated by an antibody predominantly recognizing the C-terminal domains of Ssc1, while almost all of the imported full-length Ssc1 was precipitated (data not shown). Taken together, the results demonstrate that the ATPase domains of all three mitochondrial Hsp70s are principal binding sites for Tim44. The PBDs of Ssc3 and Ssq1 exert a negative influence on their ATPase domains that results in the inability to interact with Tim44 and to perform the translocation function of Ssc1. Interestingly, the interaction of the ATPase domains of Ssc3 or Ssq1 with Tim44 was destabilized by the *ssc1-2* mutation in the PBD, a defect that was indistinguishable from the effect of the mutation in the full-length Ssc1-2. This indicates that a slight alteration of the PBD can abolish interaction with Tim44 even when the ATPase domain is unaltered. In contrast to the PBD the variable domain at the C-terminus of Ssc1 exerted a stabilizing effect on the Tim44 interaction, while the variable domains of Ssq1 and Ssc3 did not contribute to the negative effect of the PBD.

The determination of the Ssc1 interaction site with Tim44 has direct consequences for the definition of the molecular mechanism of Ssc1 during mitochondrial preprotein translocation (Jensen and Johnson, 1999; Matouschek *et al.*, 2000). A spatial separation between the substrate-binding site and the interaction site with the membrane anchor Tim44 would be a prerequisite to allow the generation of a leverage on the preprotein in transit by Ssc1. A functional interaction with Tim44 is required but not sufficient for the full translocation function of Ssc1, as described by the active motor model (Voisine *et al.*, 1999; Strub *et al.*, 2000). This fact is reflected especially by the behavior of the chimeric construct $A_QP_1V_1$, which showed an ATP-dependent interaction with Tim44 and also an efficient binding of substrate proteins. However, in this construct the conformational change in the ATPase domain induced by nucleotides can not be propagated to

Table II. Yeast strains and plasmids

| Strain or plasmid | Genotype | Source |
|-------------------|--|------------------------------|
| WVY37 | <i>trp1-1; ura3-1; leu2-3,112; his3-11, 15; ade2-1; can1-100; met2-Δ1; lys2-Δ2; ssc1ΔClal::LEU2</i> [pNZ05] | This study |
| ASYC32 | <i>trp1-1; ura3-1; leu2-3,112; his3-11, 15; ade2-1; can1-100; met2-Δ1; lys2-Δ2; ssc1ΔClal::LEU2</i> [pASC07] | This study |
| PK81 | MATα; <i>ade2-101; lys2; ura3-52; trp1; leu2-3,112; ssc1-2::LEU2</i> | Gambill <i>et al.</i> (1993) |
| PK82 | MATα; <i>his4-713; lys2; ura3-52; trp1; leu2-3</i> | Gambill <i>et al.</i> (1993) |
| PK83 | MATα; <i>ade2-101; lys2; ura3-52; trp1; leu2-3,112; ssc1-3::LEU2</i> | Gambill <i>et al.</i> (1993) |
| pASC07 | <i>Amp^R CEN6 TRP1 ARSH4 SSC1</i> | This study |
| pASC08 | <i>Kana^R A_Q</i> | This study |
| pASC10 | <i>Kana^R A_QP₁V₁</i> | This study |
| pASC11 | <i>Amp^R CEN6 TRP1 ARSH4 A_QP₁V₁</i> | This study |
| pASC24 | <i>Amp^R CEN6 TRP1 ARSH4 A₁P₁V₃</i> | This study |
| pASC25 | <i>Amp^R CEN6 TRP1 ARSH4 A₃P₁V₁</i> | This study |
| pASC26 | <i>Amp^R CEN6 TRP1 ARSH4 A₁P_QV_Q</i> | This study |
| pASC27 | <i>Amp^R CEN6 TRP1 ARSH4 A₁P₁V_Q</i> | This study |
| pASC31 | <i>Amp^R CEN6 TRP1 ARSH4 A₁P₃V₃</i> | This study |
| pASC33 | <i>Amp^R CEN6 TRP1 ARSH4 A₁</i> | This study |
| pASC35 | <i>Amp^R CEN6 TRP1 ARSH4 A_Q</i> | This study |
| pASC36 | <i>Amp^R CEN6 TRP1 ARSH4 A₃</i> | This study |
| pASC37 | <i>Amp^R CEN6 TRP1 ARSH4 A₃P₁₋₂V₁</i> | This study |
| pASC38 | <i>Amp^R CEN6 TRP1 ARSH4 A_QP₁₋₂V₁</i> | This study |
| pASC40 | <i>Amp^R CEN6 TRP1 ARSH4 P₁V₁</i> | This study |
| pASC41 | <i>Amp^R CEN6 TRP1 ARSH4 P₃V₃</i> | This study |
| pASC42 | <i>Amp^R CEN6 TRP1 ARSH4 P_QV_Q</i> | This study |
| pASE02 | <i>Amp^R CEN6 TRP1 ARSH4 SSC3</i> | This study |
| pASE06 | <i>Kana^R p₁SSC3</i> | This study |
| pASE08 | <i>Amp^R CEN6 TRP1 ARSH4 p₁SSC3</i> | This study |
| pASE10 | <i>Amp^R 2μ TRP1 ARSH4 SSC3</i> | This study |
| pASQ04 | <i>Amp^R SSQ1</i> | This study |
| pNZ05 | <i>Amp^R CEN6 URA3 ARSH4 SSC1</i> | This study |
| pNZ12 | <i>Amp^R SSC3</i> | This study |
| pNZ46 | <i>Amp^R 2μ TRP1 ARSH4 p₁SSC3</i> | This study |
| pRS314 | <i>Amp^R CEN6 TRP1 ARSH4</i> | Sikorski and Hieter (1989) |
| pRS316 | <i>Amp^R CEN6 URA3 ARSH4</i> | Sikorski and Hieter (1989) |
| pRS424 | <i>Amp^R 2μ TRP1 ARSH4</i> | Sikorski and Hieter (1989) |

the PBD, resulting in a strong translocation defect and a lethal phenotype *in vivo*. Since in contrast to Ssc1 substrate binding of A_QP₁V₁ was not influenced by the presence of ATP, the observed interaction with Tim44 correlates with the behavior of the ATPase domain but not with that of the PBD. Although it has been proposed that Tim44 functions in analogy to members of the conserved DnaJ protein family (Merlin *et al.*, 1999), the interaction properties of the chimeric constructs indicate that the binding between Hsp70 and Tim44 is structurally different from the binding of DnaJ homologs. For the bacterial Hsp70 DnaK it was proposed that DnaJ has two binding sites, one in the ATPase domain and one near the substrate-binding pocket (Gässler *et al.*, 1998). In contrast to Ssc1, neither the single ATPase domain nor the PBD of DnaK can be found in complex with DnaJ (Suh *et al.*, 1999). Not only the complex formation but also the stimulation of ATPase activity and substrate binding of DnaK requires at least the presence of the PBD adjacent to the ATPase domain. A definition of the molecular mechanism of both substrate-ATPase coupling in the case of DnaJ proteins or the negative effect of the PBD on the Tim44 interaction requires an understanding of the structural cooperation between Hsp70 ATPase and the C-terminal domains. The negative influence on the ATPase domains of the mitochondrial Hsp70s could be either caused by a direct steric hindrance of Tim44 interaction by the PBD or indirectly by the alteration of the properties of the ATPase domains. The characterization of cytosolic members of the

Hsp70 family established that an important determinant of functional specificity is largely based on specific interactions between the ATPase and the substrate-binding domain (James *et al.*, 1997). The substrate specificity set by the particular molecular environment of the peptide-binding pocket is only of minor influence. The activity of Hsp70 proteins like substrate-binding affinity is mostly determined by the nucleotide-state of the ATPase domains. In the case of mitochondrial Hsp70s a functionally important property, i.e. the interaction with Tim44, is determined by a reverse control of the PBD on the ATPase domain. This destabilizing effect of the PBD on the Tim44 interaction observed in Ssc3 and Ssq1 has significant consequences for the cellular function of structurally very closely related proteins.

Materials and methods

Construction of plasmids and yeast strains

Table II contains brief descriptions of all yeast strains and plasmids used in this study. Details of their construction are listed as Supplementary data (available at *The EMBO Journal* Online).

Complementation assays

Plasmids carrying the gene of interest were transformed into strain WVY37 (*ssc1Δ::LEU2* [pNZ5]) containing the wild-type gene of SSC1 on a centromeric plasmid with the *URA3* marker gene following standard procedures. Transformants were plated on solid YPD medium containing 5-fluoro-orotic acid (5-FOA), which only allowed growth of cells without the wild-type *SSC1* gene.

Protein import into mitochondria and co-immunoprecipitation

Mitochondria were isolated from yeast cells according to published procedures (Ryan *et al.*, 2001). Preproteins were synthesized and imported *in vitro* into yeast mitochondria essentially as described previously (Ryan *et al.*, 2001). Mitochondria were resuspended in lysis buffer [0.3% (v/v) Triton X-100, 200 mM KCl, 30 mM Tris–HCl, pH 7.4, 5% (v/v) glycerol, 0.5 mM phenylmethylsulfonyl fluoride, protease inhibitor mix (Roche) and either 5 mM MgCl₂/2 mM ATP or 5 mM EDTA]. After a clarifying spin (20 000 g, 5 min) the supernatant was transferred to Tim44 antibodies that were bound to protein A–Sepharose (Amersham-Pharmacia) in lysis buffer. The sepharose was incubated for 1 h at 4°C and washed three times with lysis buffer. Bound proteins were eluted by addition of electrophoresis sample buffer, applied to SDS–PAGE, and detected and quantified using a storage PhosphorImaging system (Molecular Dynamics).

Co-immunoprecipitation of Tim44–Hsp70 complexes

The co-immunoprecipitation experiments using affinity-purified Tim44 antibodies were performed essentially as described previously (Schmidt *et al.*, 2001).

Binding assays with ATP-agarose and CMLA–Sepharose

The binding assays to ATP-agarose and CMLA–Sepharose were performed as described previously (Schmidt *et al.*, 2001).

Supplementary data

Supplementary data for this paper are available at *The EMBO Journal* Online.

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