Folding in vivo of a newly translated yeast cytosolic enzyme is mediated by the SSA class of cytosolic yeast Hsp70 proteins

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ABSTRACT The nature of chaperone action in the eukaryotic cytosol that assists newly translated cytosolic proteins to reach the native state has remained poorly defined. Actin, tubulin, and Gα-transducin are assisted by the cytosolic chaperonin, CCT, but many other proteins, for example, ornithine transcarbamoylase (OTC), a cytosolic homotrimeric enzyme of yeast, do not require CCT action. Here, we observe that yeast cytosolic OTC is assisted to its native state by the SSA class of yeast cytosolic Hsp70 proteins. In vitro, refolding of OTC diluted from denaturant was assisted by crude yeast cytosol and ATP and found to be directed by SSA1/2. In vivo, when OTC was induced in a temperature-sensitive SSA-deficient strain, it exhibited reduced specific activity, and nonnative subunits were detected in the soluble fraction. These findings indicate that, in vivo, the Hsp70 system assists in folding at least some newly translated cytosolic enzymes, most likely functioning in a posttranslational manner.

An important question in the study of cellular-protein folding concerns the nature of chaperone action in assisting newly translated eukaryotic cytosolic proteins to the native state. Studies of the bacterial system have established an essential role at all temperatures of the double-ring chaperonin, GroEL, in cooperation with its cochaperonin, GroES, in assisting a large number of proteins to reach native form through the actions of binding and folding in a large central cavity (1-4). However, it has been unclear whether there is similar assistance of folding of newly made proteins in the eukaryotic cytosol (5). Although the cytosolic chaperonin CCT (also known as TCP1 complex or TRiC) is present in the cytosol and has been shown to be essential in Sacccharomyces cerevisiae, it seems to assist only a limited set of substrates, including actin, tubulin, and Gα-transducin (6, 7). Likewise, the Hsp90 class of chaperone is essential but seems to act on only a limited range of proteins, including steroid receptor molecules and signal-transducing kinases (8-10). Thus it remains unclear whether other newly translated eukaryotic cytosolic proteins require chaperone assistance to reach the native state.

A variety of experiments have supported the idea that chaperones are involved in the folding of newly translated eukaryotic cytosolic proteins. Early studies in HeLa cells, for example, identified interaction of a broad collective of newly translated polypeptides with Hsc70 by coimmunoprecipitation (11). Subsequent studies with both intact yeast (12) and translated polypeptides with Hsc70 by coimmunoprecipitation (13) showed interaction of nascent chains with the SSB class of yeast Hsp70 proteins. A functional requirement for Hsp70 action was indicated in studies translating the peroxisomal protein, firefly luciferase, in reticulocyte lysate—in an Hsp70-imunodepleted lysate, the newly translated protein failed to reach native form (14). Other studies suggested an involvement of the essential Hsp90 chaperone in folding cytosolic proteins distinct from recognized substrates such as steroid receptors and signal-transducing kinases. For example, purified Hsp90 could trap nonnative conformations of β-galactosidase after dilution from denaturant; these complexes slowly produced the native state when Hsp70, Hdj1, and ATP were added (15). In other studies, renaturation of firefly luciferase was enhanced by Hsp90 (16-18). However, in vivo studies that expressed these same proteins in a conditional lethal Hsp90 yeast mutant at nonpermissive temperatures did not detect any effect on the acquisition of normal specific activity (10).

To assess further the role of cytosolic chaperones in assisting the folding of newly translated eukaryotic cytosolic proteins, we selected a yeast cytosolic enzyme, ornithine transcarbamoylase (OTC), as reporter, and examined its folding both in vitro and in vivo in yeast lysate and intact cells.

METHODS

Proteins. Yeast OTC with a COOH-terminal myc tag was generated by the addition of 10 codons, encoding EQKLI-SEEDL, to the COOH terminus of the yeast OTC (ARG3) gene. OTC and OTC-myc proteins were produced in Escherichia coli by expression of pET14a derivatives carrying the respective coding regions in BL21 cells. For purification, a soluble fraction from sonicated isopropyl β-D-thiogalactoside-induced cells (centrifuged at 15,000 × g for 15 min) was subjected to two steps of HiTrapQ chromatography (Pharmacia), first at pH 6.0 in 50 mM potassium phosphate, then at pH 8.5 in 50 mM Tris-HCl, followed by gel filtration on S300 (Pharmacia). For renaturation studies, purified OTC or OTC-myc (homotramer) was denatured at 23°C for 2 h in 6 M guanidine-HCl and 10 mM DTT. For assay of ATP-dependent refolding, the denatured enzyme (100 μM in monomer) was diluted 100-fold into cytosol or chromatographic fractions of yeast that had been first diazoyed for 4 h at 4°C against 50 mM Hepes, pH 7.4/100 mM KCl/5 mM MgCl2/1 mM DTT, then split into equal portions, each 100 μl in volume, one of which was supplemented with ATP to 5 mM. The pairs of mixtures were incubated at 30°C for 1 h, then 2–5 μl were assayed for OTC enzymatic activity according to Kalousek et al. (ref. 19; protocol available on request).

SSA1/2 proteins were purified from the pep4-deficient strain, JHRY20–2C (20), according to the method of Levy et al. (21), employing anion-exchange chromatography followed by ATP-agarose chromatography.

Steps of enrichment for OTC renaturing activity were carried out with the pep4-deficient strain, JHRY20–2C, which was grown to late logarithmic phase in yeast extract/peptone

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Abbreviations: OTC, ornithine transcarbamoylase; ts, temperature-sensitive.

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medium containing 2% dextrose. Spheroplasts were prepared by Zymolyase treatment (22), and lysates were prepared by Dounce homogenization in 20 mM Hepes, pH 7.4/50 mM KCl/1 mM EDTA/1 mM DTT/10% glycerol/2 mM phenylmethylsulfonyl fluoride. The lysate was cleared by 15,000 × g centrifugation for 20 min at 4°C. Notably, no mitochondrial Hsp60 was detected in the cleared lysate on immunoblotting with antiyeast Hsp60 antibodies. Subsequent chromatographic steps were carried out as indicated in Fig. 2. A gradient of KCl concentration was used to elute the SourceQ and HiTrap heparin columns. For ATP-agarose chromatography, elution was carried out with 10 mM ATP.

Peptide sequencing was carried out on tryptic peptides prepared from SDS/PAGE bands according to the method of Stone and Williams (23). The peptides identified were RLI-GRNFNDPVEQDGMK, matching the sequence of SSA2, and NFTPEIQSSMVLGKMK, matching the sequences of both SSA1 and SSA2.

Strains. For preparation of yeast cytosolic extract, the strain JHRY20–2C was employed (pep4::URA3 his4–519 ura3–2 leu2–3,112 trp1; ref. 20), SSA1WT (WY12) and ssalts (WY13) strains (produced in E.A.C.’s lab; ref. 24) carried respectively a single intact SSA4 gene, SSA1, or a temperature-sensitive (ts) version in the chromosomal locus, ssa1–45, in the background MATa his3–11,15 leu2–3,112 ura3–52 trp1–Δ1 lys2::LEU2 ssa3::TRP1 ssa4::LYS2 pep4::HIS3. The Hsp90 wild-type and mutant strains, p82a and 1–101a, kindly supplied by Susan Lindquist, are W303 derivatives (MATa leu2–3,112 trp1–Δ1 ura3–1 ade2–1 his3–11,15 his2::LEU2 hsc82::LEU2 bearing CEN-TRP plasmids encoding either wild-type HSP90 or a ts version with a G170D codon change (25). The ts CCT strains, with conditional substitutions in α- or β-subunits, have been described (26) and were all studied in the background of YPH500 (MATa ura3–2 lys2–801 ade2–101 trp1–Δ63 his3–Δ200 leu2Δ1; ref. 27). The strains ts in CCTα, alleles 25 and 47, were produced by plasmid shuffling (28) (a hydroxylamine-mutagenized CCTα CEN-His3 plasmid with an unmutagenized (URA) plasmid, in the chromosomal CCTα-disrupted YPH500 derivative that contains a LEU2 marker replacing the CCTα coding sequence and carries a Ycp50 plasmid with CCTα. The strain deleted of the SSB1 and SSB2 genes, JN208, and an isogenic wild-type strain, JN54, (from E.A.C.’s lab) were MATa his3–11,15 leu2–3,112 lys2::LEU2 ssa3–52 (12). An HSP104 disruptant was produced by transformation of YPH500 with a linear DNA containing the HSP104 gene replaced between the EcoRI and KpnI sites with URA3.

Cell Studies. For induction of OTC-myc in yeast, wild-type or chaperone-deficient transformants (see Strains), containing the plasmid, pGALOTC-myc, bearing the OTC-myc coding segment adjoining with a GAL1 operon promoter in the plasmid, pCGS109 (29), were grown to logarithmic phase in yeast extract/peptone medium containing 2% ethanol/3% glycerol/2% raffinose and then, for induction, were transferred to yeast extract/peptone medium containing 2% galactose that had been preequilibrated to the temperature of induction. After 2 h, cells were collected, washed with 10 mM NaNO3, and disrupted by vortexing (three times for 45 sec at 4°C) with an equal volume of acid-washed glass beads (∼50 μl/OD600 of cells) in a buffer containing 20 mM Hepes (pH 7.4), 50 mM KCl, 1 mM EDTA, and 1 mM phenylmethylsulfonyl fluoride. The lysate was centrifuged at 15,000 × g for 15 min, and the supernatant was either dialyzed, as above, in preparation for OTC enzymatic assay, or directly fractionated in SDS/10% PAG or in a nondenaturing gel. Gels were analyzed by immunoblotting with anti-myc 9E10 monoclonal antibodies (Santa Cruz Biotechnology) as described (26).

Known amounts of purified OTC-myc protein were applied in parallel with cell extract and dilutions thereof (particularly of wild-type extract) to provide standards for quantitative analysis. A linear range of densitometric signal was observed between 1 and 10 ng of OTC-myc. In the case of the SSA1WT and ssa1ts strains, OTC enzymatic activity was also measured in anti-myc immunoprecipitates of cell extract to measure directly the amount of activity from the induced OTC-myc protein as distinct from any significant activity contributed by the endogenous ARG3 gene product. This result was particularly important for specific-activity determination in the ssa1ts strain at 37°C, where only a low level of activity was detected. OTC enzyme assay was carried out by addition of ornithine and carbamoyl phosphate substrates to the protein A Sepharose beads washed in enzyme assay buffer. After 5 min at 30°C, the beads were removed, and the citrulline produced was measured colorimetrically according to the method described by Kalousek et al. (19). Results are expressed as nanomoles of citrulline produced per minute (milliunits). The amounts of enzymatic activity measured in the immunoprecipitates proved to be the same as those measured directly on the soluble fraction, supporting the idea that endogenous ARG3 does not contribute significant levels of activity in these studies.

RESULTS AND DISCUSSION

Cytosolic Yeast OTC as a Reporter of Chaperone Activity. Yeast cytosolic OTC, a product of the ARG3 gene, like the well-characterized homologous mammalian mitochondrial urea-cycle enzyme, is a homotrimer of 36-kDa subunits (30). The mammalian OTC subunit, after import into mitochondria, requires the action of the matrix-localized chaperonin, Hsp60, to reach native form (31). Analogously, when the mature subunit of the mammalian enzyme is expressed in E. coli, the bacterial chaperonin, GroEL, is required to produce active enzyme (2). The yeast OTC subunit bears 38% identity (48% similarity) to the mammalian subunit and, like the mitochondrial protein, largely misfolds after dilution from 6 M guanidine-HCl (Fig. 1A, bar 2). In contrast to the folding of its mammalian homologue by Hsp60, however, the yeast OTC subunit does not seem to be recognized or assisted in folding by the homologously localized CCT. In particular, when yeast OTC subunits were expressed at nonpermissive temperature in ts lethal CCT-deficient yeast strains, they reached active form with the same kinetics as in wild-type cells (data not shown). Correspondingly, yeast OTC subunits, either translated in reticulocyte lysate, which contains CCT, or diluted from denaturant into a mixture with purified CFT, failed to become associated with the chaperonin (not shown).

ATP-Dependent Renaturation of Yeast OTC by Crude Yeast Cytosolic Extract. To aid in the immunological identification of yeast OTC, a variant was produced that contained a COOH-terminal myc epitope, OTC-myc. After OTC-myc was overexpressed in E. coli and purified, it had the same physical characteristics and activity as wild-type yeast OTC (not shown). To determine whether there is an activity in yeast cytosol that assists OTC folding, purified OTC-myc was denatured in 6 M guanidine-HCl and diluted into a crude postmitochondrial yeast cytosolic extract. When unsupplemented extract was used, only ∼20% activity was recovered, as compared with ∼10% recovered when diluted into buffer (Fig. 1A, bar 3). However, when the extract was supplemented with 5 mM ATP at the time of dilution, nearly full recovery of OTC activity was achieved (Fig. 1A, bar 4). These results (identical for wild-type OTC) are consistent with the presence of an ATP-dependent chaperone(s) in the yeast cytosol that supports the renaturation of nonnative OTC subunits. Interestingly, when the addition of ATP to the extract containing denatured OTC was delayed, recovery was reduced progressively (Fig. 1B). Because OTC becomes incapable of reaching native form if ATP is withheld from the mixture initially, it seems that, in the absence of nucleotide, the chaperone component(s) does not compete successfully against misfold-
OTC or OTC-myc was diluted 100-fold from 6 M guanidine. Extent of renaturation is reduced when ATP is added at later times. To identify the component(s) involved in OTC renaturation in the cytosolic extract, steps of chromatographic separation were carried out as outlined in Fig. 2A. A peak of renaturing activity was observed, with the most significant effects occurring between 280 and 320 mM NaCl. Despite excellent recovery of activity at this step, subsequent fractionation steps produced only 10- to 20-fold more enrichment of renaturing activity in relation to total protein concentration, with a large loss of total activity. Neither reordering the fractionation steps, nor use of alternative separation procedures (e.g., sucrose gradient fractionation), nor supplementation with nucleotide during fractionation steps could overcome these losses. Although the activity loss could reflect separation of a component that cooperates with the ATP-dependent one, recombining fractions after chromatography was also unsuccessful in restoring activity. Despite the modest enrichment of activity, there was, nevertheless, a progressive enrichment of a 70-kDa protein species associated with the active fractions.

Tryptic digestion of the 70-kDa protein species and amino acid sequencing of peptides identified peptides matching the SSA2 protein, a constitutively expressed member of the abundant cytosolic Hsp70 chaperone family, consisting of four related proteins (SSA1–4). Although members of this family are functionally redundant, at least one must be active (at relatively high concentration) for viability of yeast (32). To date, the SSA family has been implicated in posttranslational maintenance of precursor proteins in conformations competent for import into the endoplasmic reticulum and mitochondria (24, 33, 34). This action in the cytosol contrasts with the cotranslational interaction of the other class of cytosolic yeast Hsp70 proteins, the SSB class, with nascent chains exiting from the ribosome (12, 13).

To determine whether the SSA proteins were, in fact, responsible for the OTC-renaturation activity observed in the yeast cytosolic fraction, extracts were prepared from SSA-deficient yeast strains, and their renaturing activity was compared with that of wild-type yeast cytosolic extract (Fig. 3). When an extract from a strain containing only a single intact SSA gene, SSA1, was tested, it exhibited only ~20% of the level of activity of the wild-type extract (Fig. 3, SSA1/WT, black bar). SSA1 is normally present at only 30–40% of the level of SSA2 under nonstress conditions.) Even more strikingly, when an extract from a strain bearing a single SSA with a ts mutational defect (ssa1ts) was examined, no significant renaturation activity was detected (Fig. 3, ssa1ts, black bar). These results suggested that the absolute level of functional SSA protein in the cytosolic extract determined the level of renaturation observed. To test this conclusion further, purified SSA protein (a mixture of SSA1 and SSA2) obtained from a wild-type strain was added to the respective extracts, and the renaturing activity was measured. In each case, an increment of additional renaturation was observed, with the most significant effects
in vitro by the SSA is related to the level of SSA (Hsp70) protein, influenced either in vivo, biogenesis SSA1 protein in the yeast cytosolic extract.

To assess whether SSA proteins play a role in OTC renaturation, we carried out an experiment as in Fig. 1 for extracts prepared from three different yeast strains: wild-type strain (WT), with all four SSA genes intact; SSA1WT strain, with only SSA1 intact and SSA2-4 deleted; or ssa1ts strain, with a ssa1-45 ts version of SSA1 and SSA2-4 deleted. A control was also carried out with buffer only (–).

The extracts were tested for renaturing activity unsupplemented (black bars), supplemented with the mixture of purified SSA1 and SSA2 proteins obtained from wild-type yeast grown at 30°C to levels of either 0.75 μM (open bars), or 1.5 μM (hatched bars). Activity is expressed by subtracting activity measured in the absence of ATP from that in its presence and expressing this difference as a percentage of the total input enzymatic activity. Note that provision of SSA1 and SSA2 proteins to a buffer extract is insufficient to promote OTC-myc renaturation (right-hand bars), consistent with requirement for other components that are present in cytosolic extract.

observed on the SSA1 and ssa1ts extracts (Fig. 3, SSA1 WT, open and hatched bars, and ssa1ts, open and hatched bars). Thus the levels of functional SSA protein present in the cytosolic extract, whether determined by endogenous levels present in the cells from which extract is prepared or adjusted by exogenous supplementation, directly correlates with the extent to which OTC was renatured. These results support the idea that SSA is the ATP-dependent component that mediates the renaturation of OTC in the yeast cytosolic extract.

Production of Native OTC in Intact Yeast Is Assisted by SSA Proteins. To assess whether SSA proteins play a role in OTC biogenesis in vivo, we programmed galactose-inducible expression of OTC-myc in yeast bearing either SSA1 as the single intact SSA (referred to as SSA1/WT in Fig. 3) or only a ts ssa1 allele (ssa1ts). The two strains were grown in glucose medium at 24°C and were then shifted to galactose-containing medium at 37°C. After 90 min, cells were disrupted, and a postmitochondrial supernatant was prepared. One portion of this extract was analyzed directly for the amount of induced OTC-myc protein by solubilization, SDS/PAGE, and immunoblotting with anti-myc antibody, whereas the other portion was assayed for the amount of OTC enzymatic activity (Fig. 4 and see Methods). From extracts of identical amounts of SSA1 and ssa1ts mutant cells, the amounts of enzymatic activity were compared with the amounts of OTC-myc protein detected. These data provided a measurement of the specific activity of the induced OTC, indicating how efficiently the newly made protein had folded/assembled to its native active form. At 24°C, there was a 2-fold reduction in the level of enzyme activity in ssa1ts as compared with SSA1 cells, even though nearly the same amounts of OTC-myc protein were produced (Fig. 4A, left-hand lanes), indicating that the specific activity of the protein in the mutant strain was reduced to ~60%. Thus, at 24°C, only ~60% of the newly translated OTC-myc subunits reached enzymatically active form in ssa1ts.

At 37°C, SSA-deficiency affected both the synthesis of OTC subunits and their activity. The amount of OTC-myc protein in ssa1ts was reduced to ~10% that in SSA1, reflecting a general defect of translation observed in pulse-labeling studies (Fig. 4B), but OTC enzymatic activity was reduced to an even greater degree in ssa1ts, measuring only ~2% of that in SSA1. These relative amounts of OTC activity and protein indicate that the specific activity of OTC in the mutant strain at 37°C...
was only \(\approx 25\%\) that in SSA1 (Fig. 4A, right-hand lanes). Assuming that virtually all of the newly made OTC-myc subunits in SSA1 cells reach the native homotrimeric state, it would thus seem that only \(\approx 25\%\) of newly translated OTC-myc becomes active homotrimer in the SSA-deficient strain at nonpermissive temperature, with the remainder in an inactive, presumably misfolded state.

To assess the fate of nonnative OTC-myc subunits in SSA1ts cells, we first addressed whether they became lodged in large, insoluble aggregates. Extracts were subjected to a high-speed centrifugation step (340,000 \(\times\) g for 10 min), and supernatant and pellet fractions were examined by immunoblotting. No significant amount of OTC-myc was found in the insoluble fraction (not shown). Because the nonnative subunits apparently remained soluble, we sought to observe them in nondenaturing gel electrophoresis (Fig. 5). When SSA1 extract was applied, followed by immunoblotting, virtually all of the OTC-myc was found at a single position that corresponded to that of purified homotrimer. When mutant extracts were examined at various times after temperature shift, however, only a small portion of the OTC-myc migrated to this position, and the majority was detected at other positions in the gel. In particular, 1.5 h after temperature shift (the time used in Fig. 4A), \(\approx 80\%\) of the subunits migrated to other positions. This behavior is in agreement with the specific activity measurements, which showed that \(\approx 80\%\) of induced OTC-myc in mutant cells had failed to reach active form at this time after shift. The nonnative protein seemed to occupy several different conformational states. A significant portion was detected in the gel slot, as compared with little or no signal at this position in SSA1, suggesting that some subunits had, in fact, aggregated. Another portion was found as a distinct species migrating faster than homotrimer. These molecules might be a population of unassembled OTC monomers, or alternatively, might comprise a population of OTC-myc bound to a cellular component. Whatever the exact fate of such species, we conclude that a substantial portion of the newly translated OTC produced in the SSA-deficient cells failed to be incorporated into native, active homotrimer.

Studies carried out here with both yeast cytosolic extracts and intact cells indicate that the newly translated subunits of OTC, a yeast cytosolic enzyme, are assisted in reaching their selective insolubility in the mutant cells. Indeed, when ssat1ts cells were subjected to pulse–chase and extracts were separated into soluble and insoluble fractions, we did not observe any change in the amount or identity of insoluble proteins after two-dimensional gel electrophoresis of the urea-solubilized precipitates (S.K., unpublished results).

SSA Action in Polypeptide-Chain Folding vs. Oligomeric Assembly. The action of SSA proteins in facilitating OTC biogenesis likely lies at the level of assisting OTC monomers to reach an assembly-competent conformation, because OTC proteins function to maintain newly made mitochondrial and endoplasmic-reticulum precursor proteins in translocation-competent states apparently by binding and conformational adjustment of monomeric species (24, 33, 34). Likewise, the action of Hsp70 proteins at the “trans” side of membranes in driving membrane translocation seems to involve interaction with monomeric species (35–39). Perhaps even more directly indicative is the fact that purified SSA1 protein, when in the presence of its cooperating partner protein YDJ1, was able to assist the refolding of monomeric firefly luciferase diluted from denaturant (21). Nevertheless, a role in assisting already folded OTC monomers to assemble into native homotrimer cannot be ruled out, particularly in light of the observation of a discrete species that may be unassembled monomers in the nondenaturing gel analysis of SSA-deficient mutant extract (Fig. 5).

SSA Assistance Is Likely to Be Posttranslational. The action of SSA proteins in assisting biogenesis of cytosolic OTC seems to be exerted posttranslationally. In the \(in vitro\) studies with cytosolic extract, full-length OTC subunits were the target of assistance by SSA. More significantly, however, both in a yeast-cell-free translation system and \(in vivo\), SSA proteins do not seem to form cotranslational interactions with translating polypeptides; such interactions are mediated, rather, by the specific activity of the induced OTC-myc protein is displayed, measured as the amount of OTC enzymatic activity per nanogram of OTC-myc protein.
class of Hsp70 proteins known as SSB proteins (12, 13). In the studies referred to below, we observed that mutational alteration of this class of proteins did not interfere with OTC biogenesis. Also consistent with posttranslational action of SSA proteins are the observations concerning their action in maintaining precursor-protein translocation competence, which also seems to be mediated post-translationally on full-length polypeptide chains (24, 33, 34).

**Cooperation with SSA by Other Components.** Having observed a role for SSA proteins in assisting biogenesis of OTC in the yeast cytosol, it seems appropriate to ask whether the normal cooperating partner of SSA proteins, the yeast DnaJ homologue YDJ1 (24, 40), also participates. In preliminary immunodepletion experiments in vitro that examined ATP-dependent OTC renaturation by a yeast cytosolic extract that had been immunodepleted of either SSA proteins or of YDJ1, it seems that YDJ1 is also required. Although the cooperating chaperones SSA and YDJ1 seem to be involved in folding OTC in the context of a cytosolic extract, notably, the two purified chaperones were unable to reanimate OTC in vitro (data not shown). This result raises questions as to whether other components participate or whether a characteristic assembly of SSA and YDJ1 must be formed (41). Parenthetically, we note that in the in vivo experiments, because translation was affected, another involved component may have become limiting at nonpermissive temperature. We tested whether other major cytosolic chaperone systems are involved in vivo by placing the galactose-inducible yeast OTC-myc gene into yeast strains bearing ts lethal versions of Hsp90 or of subunits of CCT or the latter two of which are cold-sensitive (12) or thermointolerant (42), respectively. For each strain, OTC-myc was induced after shift to the nonpermissive temperature (except for Hsp104 where 37°C was employed), and specific activity of the newly made OTC-myc was measured as it had been in the SSA-deficient strain (Fig. 6). This survey showed that OTC biogenesis was affected only in the SSA-deficient strain.

The mechanism by which an Hsp70 system alone could assist folding to native form is interesting to consider. Whereas folding at chaperonin assemblies occurs in an encapsulated central cavity, this mechanism does not seem to be followed by the Hsp70 class of chaperones. Rather, it seems most likely that SSA/YDJ1 bind to local, most likely, hydrophobic, segments of nonnative cytosolic OTC and that productive folding and assembly occur after ATP-directed release from the chaperones. We conclude that, at least for this cytosolic enzyme, the major chaperone assistance in folding/assembly of its newly translated subunits is mediated posttranslationally by the SSA system. Further studies should resolve whether such action is also required by other proteins of the eukaryotic cytosol.

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