Heterologous Expression of a Plant Small Heat-Shock Protein Enhances* * Escherichia coli Viability under Heat and Cold Stress

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A small heat-shock protein (sHSP) that shows molecular chaperone activity in vitro was recently purified from mature chestnut (*Castanea sativa*) cotyledons. This protein, named here as CsHSP17.5, belongs to cytosolic class I, as revealed by cDNA sequencing and immunoelectron microscopy. Recombinant CsHSP17.5 was overexpressed in *Escherichia coli* to study its possible function under stress conditions. Upon transfer from 37°C to 50°C, a temperature known to cause cell autolysis, those cells that accumulated CsHSP17.5 showed improved viability compared with control cultures. Sodium dodecyl sulfate-polyacrylamide gel electrophoresis analysis of cell lysates suggested that such a protective effect in vivo is due to the ability of recombinant sHSP to maintain soluble cytosolic proteins in their native conformation, with little substrate specificity. To test the recent hypothesis that sHSPs may be involved in protection against cold stress, we also studied the viability of recombinant cells at 4°C. Unlike the major heat-induced chaperone, GroEL/ES, the chestnut sHSP significantly enhanced cell survivility at this temperature. CsHSP17.5 thus represents an example of a HSP capable of protecting cells against both thermal extremes. Consistent with these findings, high-level induction of homologous transcripts was observed in vegetative tissues of chestnut plantlets exposed to either type of thermal stress but not salt stress.

Both prokaryotes and eukaryotes synthesize a set of proteins that can interact with nonnative polypeptide chains to prevent irreversible aggregation reactions and/or nonproductive folding pathways. Most of these so-called molecular chaperones are induced as part of the ubiquitous heat-shock response and accordingly have been classified into HSP families (Vierling, 1991; Parsell and Lindquist, 1993; Boston et al., 1996). Although HSP induction is correlated with the acquisition of thermotolerance in a variety of organisms, the roles played by individual components in such processes are often not well understood (Boston et al., 1996; Hartl, 1996). Different organisms accumulate different HSPs in response to similar levels of stress. For example, HSP104 has been shown to play a major protective role upon heat shock in yeast but not in *Drosophila melanogaster*, which does not even synthesize an HSP100 protein in response to thermal stress (Sanchez and Lindquist, 1990). Stress conditions other than elevated temperatures can also lead to HSP induction (Vierling, 1991; Parsell and Lindquist, 1993).

The heat-shock response in plants has been extensively investigated for more than a decade (Vierling, 1991; Boston et al., 1996; Waters et al., 1996). In contrast to other eukaryotes, the most prominent heat-induced proteins of plants are the sHSPs, a structurally diverse family of polypeptides with sizes ranging from approximately 15 to 30 kD. sHSPs are encoded in higher plants by at least six multigene families and have been localized to the cytoplasm, ER, mitochondria, and chloroplast (Boston et al., 1996; Waters et al., 1996). Considerably fewer sHSPs have been identified in animals, yeast, and prokaryotes, where the most important HSPs belong to the families HSP60, HSP70, HSP90, and HSP110 (Hartl, 1996).

The in vivo function of plant sHSPs is largely unknown at present. A few studies have demonstrated that at least some members can function as molecular chaperones in vitro (Jinn et al., 1989, 1995; Lee et al., 1995, 1997; Collada et al., 1997), as is the case for mammalian sHSPs and the related α-crystallin eye-lens proteins (Horwitz, 1992; Buchner, 1996). As a result of analyses of the progeny of heat-tolerant and nontolerant variants of *Agrostis palustris*, a thermoprotective role has been proposed for specific HSP25 proteins (Park et al., 1996). More recently, enhanced thermostolerance has been reported in recombinant *Escherichia coli* cells expressing a glutathione S-transferase/rice HSP16.9 fusion protein (Yeh et al., 1997). Other studies with heat-stressed tomato fruits have shown a correlation between the accumulation of sHSPs (as well as other heat-

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Abbreviations: HSP, heat-shock protein; IPTG, isopropyl-1-thio-β-galactoside; sHSP, small (low-molecular-mass) HSP.
induced proteins) and the acquisition of chilling tolerance (Sabehat et al., 1996, 1998; Kadyrzhanova et al., 1998).

Aside from heat stress, other environmental or developmental signals regulate the expression of plant sHSPs. The best-characterized example of developmental regulation is the induction of specific members during seed maturation at normal growth temperatures (Hernandez and Vierling, 1993; Coca et al., 1994; DeRocher and Vierling, 1994; zur Nieden et al., 1995). Recently, a sunflower sHSP promoter that is activated during zygotic embryogenesis but not by heat stress was characterized (Carrasco et al., 1997).

In contrast to other seeds, which typically accumulate low to moderate levels of sHSPs, recalcitrant chestnut (Castanea sativa) seeds contain a highly abundant shSP (Collada et al., 1997). Recalcitrant seeds have unusually high water contents and are in principle more sensitive to certain types of stress. Like other sHSPs, the chestnut protein forms high-molecular-mass complexes under nondissociating conditions and can function as a molecular chaperone in vitro (Collada et al., 1997). We report here the isolation of high-molecular-mass complexes under nondissociating conditions and can function as a molecular chaperone in vitro (Collada et al., 1997). We report here the isolation of high-molecular-mass complexes under nondissociating conditions and can function as a molecular chaperone in vitro (Collada et al., 1997). We report here the isolation of high-molecular-mass complexes under nondissociating conditions and can function as a molecular chaperone in vitro (Collada et al., 1997).

RNA Isolation and Northern Hybridization

Total RNA was obtained from chestnut plantlets as described previously (Chang et al., 1993). Poly(A+) RNA was prepared using oligo(dT)-cellulose spin columns (Pharmacia Biotech). Northern analyses were carried out following standard procedures (Maniatis et al., 1982). After hybridization, membranes were washed twice in 2X SSC (1X SSC is 0.15 M NaCl and 15 mM sodium citrate, pH 7.0) and 0.1% (w/v) SDS at room temperature for 15 min, twice in 1X SSC and 0.1% SDS for 15 min, and twice in 0.2X SSC and 0.1% SDS for 15 min. Autoradiographs were taken on Kodak X-Omat-5 film exposed overnight.

Silent Mutagenesis and Bacterial Expression of CsHSP17.5

The coding sequence for CsHSP17.5 was subcloned into the expression vector pRSET (Invitrogen, Carlsbad, CA). Previously, an internal Ni61 site (nucleotides 114–119 in Fig. 1) was eliminated by silent mutagenesis involving two sequential PCR steps. For the first reaction, we used the forward primer 5’-CTCACAGTAGATATGCGG-3’ (nucleotides 105–125, mutation underlined) and the reverse primer 5’-CTCACTGGATATATGGGACCC-3’ (Collada et al., 1997). The coding sequence for CsHSP17.5 was subcloned into the expression vector pRSET (Invitrogen, Carlsbad, CA). Previously, an internal Ni61 site (nucleotides 114–119 in Fig. 1) was eliminated by silent mutagenesis involving two sequential PCR steps. For the first reaction, we used the forward primer 5’-CTCACAGTAGATATGCGG-3’ (nucleotides 105–125, mutation underlined) and the reverse primer 5’-CTCACTGGATATATGGGACCC-3’.

MATERIALS AND METHODS

Plant Material and Stress Treatments

European chestnut (Castanea sativa Mill.) seeds were harvested at either the mature or the late-mature stage (approximately 4 weeks before shedding) in Zarzalejo, a village northwest of Madrid, Spain. After germination plantlets were kept in a growth chamber (16-h day/8-h night, 22°C/18°C, 70% RH). Heat-stress experiments were performed with 14- to 20-week-old plantlets at either 32°C or 40°C and 80% RH for 8 h. Cold treatments were carried out at 4°C for up to 4 weeks. For salt-stress experiments, plantlets were watered profusely with 200 mm NaCl for up to 48 h.

cDNA Cloning

A cDNA library in λ-Uni-ZAP XR was made from immature chestnut cotyledon poly(A+) RNA using the ZAP-cDNA Gigapack III Gold Cloning kit (Stratagene). Before ligation, the cDNA was enriched in small fragments (between approximately 2000 and 400 bp) by gel filtration on Sepharose CL-2B. The library was screened at reduced stringency with the full-length cDNA for sunflower HSP17.6 (Almoguera and Jordano, 1992). Positive cDNA clones were isolated and subjected to in vivo excision into Escherichia coli SOLR cells to render recombinant pBlue-script 5K(+) phages. For each insert both strands were completely sequenced using an automated DNA sequencer (model 373, Perkin-Elmer).

Figure 1. Nucleotide and predicted amino acid sequence of Cs hspa17.5 cDNA. The cDNA is 733 bp long and encodes a polypeptide of 154 residues (predicted M, 17,482). The 3’-untranslated region consists of 217 nucleotides and contains a putative polyadenylation signal (indicated in boldface) 158 nucleotides upstream of the poly(A) tail. The deduced amino acid sequence includes, with agreement at every residue, two internal peptides (underlined) obtained by endoproteasease Asp-N cleavage of the purified seed protein (Collada et al., 1997). CsHSP17.5 also includes two motifs (residues 61–88 and 111–140) conserved in all plant sHSPs (Waters et al., 1996).
primer 5'-CATGCCATACGGATCCACACTCC-3' (nucleotides 611–589, BamHI site underlined). The second reaction yielded the desired mutation by using the product of the first PCR as a primer and a second primer flanking the 5' end of the gene, 5'-GCAGATCATATGGCGCTCAGT-3' (NdeI site underlined; start codon in italics). A polymerase with 3'→5' proofreading activity (Pfu, Stratagene) and experimental conditions described previously (Garcia-Casado et al., 1998) was used in all amplifications. The final product was cut with NdeI and BamHI and ligated to pRSET open with the same enzymes to yield pRSET-HSP. The engineered mutation was confirmed by sequencing both strands. For bacterial expression, E. coli BL21(DE3) cells (Novagen, Madison, WI) transformed with pRSET-HSP were grown at 37°C and 250 rpm to an A600 of 1.0, then 1 mM IPTG was added, and growth was continued for up to 4 h.

Protein Purification and Immunodetection

CsHSP17.5 was purified from chestnut seeds as described previously (Collada et al., 1997). SDS-PAGE fractionation and protein immunoblotting were carried out as described by Garcia-Casado et al. (1998) using a 1:500 dilution of monospecific polyclonal antibodies to CsHSP17.5 (Collada et al., 1997).

Immunoelectron Microscopy

For immunoelectron microscopy experiments, immature chestnut cotyledons were fixed with 4% (v/v) paraformaldehyde in PBS immediately after collection. Low-temperature embedding and mounting of ultrathin sections were carried out as described previously (Rodriguez-Cerezo et al., 1997). Sections were blocked for 60 min in 30 mM Tris-HCl, pH 7.5, 0.1% (w/v) BSA, and 1% (w/v) gelatin and then incubated for 2 h with a 1:50 dilution of monospecific antibodies against CsHSP17.5. Sections were washed, colloidal gold labeled (10-nm particles), and stained with uranyl acetate and lead citrate (Rodriguez-Cerezo et al., 1997).

Cell-Viability Experiments

For heat-shock experiments cell cultures were grown at 37°C to an A600 of 1.0 and then diluted once with fresh Luria-Bertani medium supplemented with ampicillin at 100 μg/mL and IPTG to a final concentration of 1 mM. Two hours after induction, cultures were diluted to 6 × 10^5 cells mL^-1, and 1 mL samples were shifted to 50°C. Aliquots (100 μL) were taken at 0, 30, and 60 min, and serial dilutions were plated in triplicate onto Luria-Bertani plus ampicillin plates. Cell viability was estimated by counting the number of colony-forming units after incubation of the plates overnight at 37°C. For cold treatments, appropriate dilutions from induced cultures were plated onto Luria-Bertani agar supplemented with ampicillin and 1 mM IPTG. Plates were then incubated at 4°C for different periods and cell viability was estimated as described above. For both treatments (heat and cold), the means of three experiments were determined from at least two independent transformants (with SD being less than 5% in all cases).

Thermostability of Soluble Proteins in E. coli

The effect of heat shock on protein stability was analyzed in recombinant IPTG-induced E. coli cells according to the method of Muchowski and Clark (1998). At various times during the 50°C treatments, aliquots were centrifuged to collect cells. The pellets were washed once with 20 mM Tris-HCl, pH 7.5, and 1 mM EDTA and extracted with the same buffer. Crude cell lysates were then centrifuged at 16,000g for 30 min, and the supernatants were analyzed by SDS-PAGE.

RESULTS

Characterization of a cDNA Encoding Chestnut Seed sHSP

An abundant protein was recently purified from mature chestnut cotyledons that shows homology to class I sHSPs from plant sources (Collada et al., 1997). To isolate its cDNA, a library from late-mature chestnut cotyledons (1.1 × 10^8 plaque-forming units) was screened at moderate stringency with a class I sunflower HSP cDNA. Five positive clones were randomly selected and their inserts sequenced. All inserts correspond to a single nucleotide sequence (Fig. 1), which includes a 462-bp reading frame flanked by 5'- and 3'-noncoding sequences of 54 and 217 nucleotides, respectively. A putative polyadenylation signal, AATAAA, is located 158 nucleotides upstream of the poly(A^-) tail. The encoded polypeptide, which is 154 residues long, has been designated CsHSP17.5. Its predicted Mᵣ (17,482) and pi (5.95) are similar to those experimentally determined for chestnut seed sHSP (Collada et al., 1997). Moreover, CsHSP17.5 includes the two known internal peptides of the seed protein with agreement at every residue (Fig. 1). Heterologous expression in E. coli and western and northern analyses further supported the correspondence between these proteins (see below).

Database searches revealed that CsHSP17.5 had the highest amino acid sequence similarity to class I cytosolic sHSPs. Aside from the two motifs conserved in all plant sHSPs (Vierling, 1991; Waters et al., 1996), CsHSP17.5 contains an N-terminal region (residues 10–24) characteristic of class I cytosolic proteins. Like other sHSPs, CsHSP17.5 also contains an N-terminal region (residues 10–24) characteristic of class I cytosolic proteins. Like other sHSPs, CsHSP17.5 also shows a weak but significant similarity to mammalian eye-lens α-crystallins. Molecular chaperone activity has been demonstrated for members of both protein groups (Jakob et al., 1993; Lee et al., 1995).

Subcellular Location of CsHSP17.5

In mature chestnut seeds the greatest amount of CsHSP17.5 is localized in the cotyledons (Collada et al., 1997). Because good sections are difficult to obtain from mature seed tissue, cotyledonary cells of the late-mature stage were subjected to immunoelectron microscopic analysis. For these experiments monospecific antibodies against purified CsHSP17.5 were prepared as described...
Figure 2. Immunocytochemical localization of CsHSP17.5 in late-mature (approximately 4 weeks before shedding) cotyledonary cells of chestnut. A low-magnification view (A) shows the anatomy of a typical, untreated cell at this stage of seed development. Before colloidal gold labeling (10-nm particles), similar sections were incubated with preimmune serum (B) or with rabbit monospecific antibodies to CsHSP17.5 (C and D). In the latter case, gold label was found exclusively in the cytosol, with no significant labeling of starch granules (C), apoplastic space, vacuoles, or cell walls (D). No significant labeling was observed in sections treated with preimmune serum (B). a, Apoplastic space; cw, cell wall; cy, cytoplasm; s, starch granules. Bars in A = 2 μm; bars in B to D = 200 nm.

Heterologous Expression in E. coli Cells

To analyze its possible in vivo function under stress conditions, the complete coding sequence for seed CsHSP17.5 was introduced into E. coli using the pRSET expression vector. The vector alone was also introduced into E. coli as a control. Under normal culture conditions, similar growth rates were observed for both types of recombinant cells (pRSET-HSP and pRSET) and for untransformed wild-type cells (Fig. 3A). After IPTG addition, SDS-PAGE analysis showed the overproduction of recombinant CsHSP17.5 (apparent M, approximately 20,000) in extracts from pRSET-HSP cells but not in extracts from cells carrying the control plasmid. The recombinant protein reached maximal expression levels 2 to 4 h after induction and had the same apparent size as seed sHSP, as shown by SDS-PAGE (Fig. 3B).

Heat- and Cold-Stress Experiments

The effect of recombinant CsHSP17.5 on cell survival was evaluated in cultures subjected to heat stress. Two hours after IPTG addition, cultures were diluted to 6 × 10^6 cells mL^-1 and then transferred to 50°C, a temperature that is known to cause cell autolysis. At various times after the temperature shift, cell viability was measured by counting colony-forming units in serially diluted culture aliquots. Whereas cell viability decreased rapidly in both pRSET and pRSET-HSP cultures upon heat shock (Fig. 4A), the measured survival rates were significantly higher in cells overexpressing CsHSP17.5 (approximately 2-fold after 60 min at 50°C). Similar differences were observed when untransformed wild-type cells were used for comparison (not shown). Because in vitro molecular chaperone activity has been demonstrated previously for seed sHSP (Collada et al., 1997), we investigated whether the expression of recombinant CsHSP17.5 had any effect on the thermostability of bacterial soluble proteins. At various times after the shift to 50°C, culture aliquots were taken and the cells pelleted, washed, and extracted as described in “Materials and Methods.” SDS-PAGE analysis of these extracts showed that, although many soluble proteins precipitated or were rapidly degraded in control cells during the heat shock, this effect was delayed and quantitatively less pronounced in pRSET-HSP cells (Fig. 4B). Such an increase in protein thermostability is associated with a rapid insolubilization of recombinant CsHSP17.5. The inclusion of 6 M urea and 2% (w/v) SDS in the extraction buffer (not shown) ruled out degradation of CsHSP17.5 during the 50°C treatment.

Figure 3. Heterologous expression of CsHSP17.5 in E. coli BL21(DE3). A, Growth of wild-type (○) and transformed pRSET (x) and pRSET-HSP (●) cells at 37°C. At the times indicated, 1-mL culture aliquots were taken and the A600 was determined. Before induction (time 0) and at different times after IPTG addition (1, 2, 3, and 4 h), samples were taken from pRSET (control) and pRSET-HSP cultures, and crude cell lysates were prepared. Proteins (25 μg) were then fractionated by SDS-PAGE and immunoblotted with monospecific antibodies to purified seed sHSP (lane 7).

Figure 4. Heat- and cold-stress survival of CsHSP17.5-expressing E. coli cells. A, Cell viability was measured in cultures subjected to heat shock at 50°C (●) or cold shock at 0°C (○). B, SDS-PAGE analysis of protein thermostability in cultures subjected to heat shock at 50°C. The inclusion of 6 M urea and 2% (w/v) SDS in the extraction buffer (not shown) ruled out degradation of CsHSP17.5 during the 50°C treatment.
We also tested whether recombinant CsHSP17.5 might be relevant for cell viability at chilling temperatures, as hypothesized for heat-induced tomato sHSPs (Sabehat et al., 1996, 1998; Kadyrzhanova et al., 1998). For these experiments aliquots from IPTG-induced cultures were plated and kept at 4°C. At different times cell viability was measured by counting colony-forming units in triplicate plates. As shown in Figure 5, both control pRSET cells and cells overexpressing CsHSP17.5 lost viability upon storage in the cold, although at significantly different rates. The control cells died with a half-life of 5 to 6 d, and after 10 d at 4°C only about 10% remained alive; conversely, approximately 60% of the pRSET-HSP cells survived after the same period.

Figure 4. Protective effect of recombinant CsHSP17.5 on cell viability and protein stability upon heat stress in vivo. A, Viability of E. coli transformants for pRSET-HSP (hatched bars) and pRSET (white bars) constructs subjected to 50°C treatments. At the times indicated after the temperature shift, culture samples were taken, serially diluted, and plated onto Luria-Bertani plus ampicillin plates. Cell viability is plotted as the percentage of colony forming units relative to the starting number of colonies at time 0. Means of three independent experiments are shown (SD was less than 5%). B, SDS-PAGE analysis of bacterial soluble proteins during heat shock at 50°C. Based on the results described above, culture samples corresponding to similar amounts of viable cells were taken at 0, 15, 30, 45, and 60 min after heat shock. Cells (pRSET-HSP and pRSET) were pelleted and soluble proteins extracted as described in “Materials and Methods.” The position of recombinant CsHSP17.5 is marked (●).

Figure 5. Effect of recombinent CsHSP17.5 on cell viability at 4°C. Cultures grown normally and induced with IPTG for 2 h at 37°C were diluted and plated onto Luria-Bertani agar supplemented with 100 μg/ml ampicillin and 1 mM IPTG. Plates were then kept at 4°C. At the times indicated after temperature downshift, plates were transferred to 37°C and cell viability was estimated as in Figure 4A. Means of three independent experiments are shown (SD was less than 5%). White bars, pRSET cells; hatched bars, pRSET-HSP cells.

Figure 6. Induction of Cs hsp17.5 homologs in 18-week-old chestnut plantlets subjected to heat or cold stress. RNA was extracted and analyzed by northern hybridization using as a probe the first 662 bp of Cs hsp17.5 cDNA (EcoRI-Sall fragment). For heat-stress experiments, plantlets were treated at either 32°C or 40°C for 8 h. Cold-stress experiments were carried out at 4°C for up to 4 weeks. The amount of RNA loaded per lane was approximately 3 μg (seeds) or 10 μg. As a control, the same filters were hybridized with a barley 18S ribosomal probe (shown here for the cold treatment). All hybridizing bands were approximately 730 nucleotides in size. R, Roots; S, stems; L, leaves.
Expression of \textit{Cs hsp17.5} Homologs in Chestnut Plantlets

A single hybridizing band was detected when total RNA from late-mature chestnut cotyledons was probed at moderate stringency with the cDNA for \textit{CsHSP17.5} (Fig. 6). Under the same conditions a weak band of similar size could be observed in RNA from stems, but not from roots or leaves, of nonstressed chestnut plantlets. However, when plantlets of the same age were subjected to thermal stress, increased transcript abundance was observed for \textit{Cs hsp17.5} homologs in all organs analyzed. Expression was higher at 40°C than at 32°C in all cases and appeared to be especially pronounced in stems.

On the other hand, induction of \textit{Cs hsp17.5} homologs could be observed in chestnut plantlets kept at 4°C for up to 4 weeks (Fig. 6). Although the highest transcript levels were also found in stems, the time course of induction was much slower than under heat-stress conditions. It is noteworthy that no hybridizing bands could be detected under the same experimental conditions in leaves of cold-stressed plants. Likewise, no transcripts were detected when total RNA from leaves, stems, or roots of salt-stressed chestnut plantlets was hybridized with the same probe (not shown).

The patterns observed in control samples (leaves, stems, or roots) at the beginning of the cold shock (shown in Fig. 6) were consistent throughout the experiment and several months later. Hybridization of the same filters with an 18S ribosomal probe was performed to verify that similar amounts of RNA were loaded in each lane.

\section*{DISCUSSION}

It has been shown that mature chestnut cotyledons accumulate a highly abundant shSHP under normal growing conditions. This protein (\textit{CsHSP17.5}) forms oligomeric complexes under nondenaturing conditions and possesses molecular chaperone activity (Collada et al., 1997). As a first step toward analyzing its possible function in vivo, we have isolated and characterized a full-length cDNA clone that encodes \textit{CsHSP17.5} (Fig. 1). The predicted polypeptide (154 amino acids) includes the two internal peptidic sequences previously determined for the chestnut seed shSHP.

Moreover, monospecific antibodies to the purified seed protein recognized recombinant \textit{CsHSP17.5} when expressed in \textit{E. coli} (Fig. 3B). The highest sequence identity in databank searches was found with class I cytosolic shSHPs, including a characteristic N-terminal motif (residues 10–24) that was absent from class II shSHPs. Aside from cytosolic members (class I and II), the plant shSHP family includes proteins localized to the chloroplasts, ER, and mitochondria (Boston et al., 1996; Waters et al., 1996). Immunoelectron microscopic studies of chestnut cotyledonary cells revealed an overall cytoplasmic localization for \textit{CsHSP17.5} (Fig. 2), as predicted by sequence analysis.

In spite of their abundance and unique multiplicity, little is known at present about the specific role of plant shSHPs (Boston et al., 1996). Recent overexpression experiments involving \r{17}crystallins and shSHPs of animal origin (for review, see Buchner, 1996), as well as a rice shSHP-glutathione S-transferase fusion (Yeh et al., 1997), have illustrated the ability of these proteins to confer thermostolerance. To investigate the possible function of \textit{CsHSP17.5} in vivo, we introduced its coding sequence into \textit{E. coli} using the pRSET expression vector. It is known that this organism does not synthesize class I shSHPs in response to heat stress (Buchner, 1996; Yeh et al., 1997). In our study the coding sequence of \textit{Cs hsp17.5} cDNA was engineered so that no vector-encoded amino acids were present in the recombinant protein.

As shown in Figure 4A, we found that overexpression of \textit{CsHSP17.5} in \textit{E. coli} was correlated with maintenance of viability under heat-stress conditions. Furthermore, SDS-PAGE analysis of cell lysates suggested that the protective effect of \textit{CsHSP17.5} is associated with an increase in the thermostability of soluble proteins (Fig. 4B). Whether such stabilization is due directly to the chaperone function of the shSHP or to interactions with other \textit{E. coli} proteins (e.g. heat-induced chaperones) remains to be determined. Like other shSHPs, \textit{CsHSP17.5} can bind nonnative proteins in vitro and promote their renaturation in an ATP-independent manner (Collada et al., 1997; M.-A. Guevara, C. Aragoncillo, and L. Gomez, unpublished results). On the other hand, in vitro refolding experiments, as well as studies with transgenic Arabidopsis cell cultures, support the notion that shSHPs act in concert with other HSPs during the refolding process (Ehmsperger et al., 1997; Forreiter et al., 1997; Lee et al., 1997). The increased thermotolerance of pRSET-HSP cells could also be related to a hypothetical effect of \textit{CsHSP17.5} on membrane stability under heat-shock conditions. Supportive evidence of membrane stabilization has been obtained for GroEL/ES chaperones (Török et al., 1997) and a chloroplastic shSHP from \textit{Synechocystis} PCC 6803 (Hovrath et al., 1998). Likewise, heat-induced membrane association was recently described for a prokaryotic shSHP homologous to \textit{CsHSP17.5} (Jobin et al., 1997).

It is well established that exposure of plants to moderately high temperatures induces thermostolerance (Vierling, 1991). Less expected was the finding that prestorage heat treatments increase the chilling tolerance of a number of marketable fruits and vegetables (e.g. Lurie and Klein, 1991; McCollum et al., 1995; Sabehat et al., 1996). In agreement with these observations, experimental evidence is presented here that recombinant \textit{CsHSP17.5} is important in maintaining cell viability at low temperatures. As shown in Figure 5, pRSET-HSP cells overexpressing the chestnut protein died more slowly upon storage at 4°C than control cells. Such an effect on bacterial viability has been demonstrated so far only for TF (trigger factor), an abundant \textit{E. coli} protein up-regulated by low temperatures (Kandror and Goldberg, 1997). Molecular chaperone activity has been reported for both TF and \textit{CsHSP17.5}.

Although the precise reasons why bacterial cells die at 4°C have not yet been discovered, the protective effect of \textit{CsHSP17.5} might be due to the maintenance of proteins in a functional conformation, as in the case of heat stress. That role would be especially relevant at low temperatures, at which ribosomal function is inhibited and the solubility and folding properties of many proteins are substantially altered. Chaperones of the HSP70 family that are induced
by low but not high temperatures were recently described in yeast and E. coli (Craig et al., 1993; Thieringer et al., 1998). Aside from representing an example of improved chilling tolerance mediated by a nonbacterial protein, our results with pRESET-HSP cells subjected to cold stress demonstrate a novel in vivo role for sHSPs. This role had been hypothesized based on the correlation between the acquisition of chilling tolerance in heat-treated tomato fruits and the expression of sHSPs and other heat-induced proteins (Sabehat et al., 1996, 1998; Kadyrzhanova et al., 1998).

The recalcitrant seeds of chestnut, in which CsHSP17.5 accumulates abundantly, have one of the highest moisture contents known at shedding (typically >50%). Compared with orthodox seeds, high-moisture seeds are more sensitive to certain types of environmental stresses. Found naturally in a wide area of southern Europe, the seeds of chestnut must endure extreme temperatures both during maturation (August to mid-October) and during the winter, shortly after shedding. Temperatures frequently range from above 35°C in summer to below 0°C in winter. The results reported here support a role for CsHSP17.5 in protecting seed tissues against the damaging effects of both thermal extremes. This notion is reinforced by the finding that transcripts hybridizing with the Cs lsp17.5 cDNA are induced in vegetative organs of chestnut plantlets subjected to either heat or cold stress, but not salt stress (Fig. 6). In these experiments the strongest inductions were observed in stems, in which constitutive expression also occurs. Although the thermostabilizing function of sHSPs was recently extended to lipid membranes (Jobin et al., 1997; Horváth et al., 1998), the exact biochemical mechanisms by which sHSPs, and probably other HSPs, attenuate both heat- and cold-induced cell damage remain to be determined. The differential induction kinetics observed in chestnut plantlets at high and low temperatures (Fig. 6) may reflect fundamental differences in cellular requirements at each thermal extreme.

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