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Category: Cell Biology
Pea chloroplast DnaJ-J8 and Toc12 are encoded by the same gene and localized in the stroma

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This work was supported by grants from the National Science Council (NSC 99-2321-B001-015, H-mL) and Academia Sinica (H-mL) of Taiwan.

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The author(s) responsible for distribution of materials integral to the findings presented in this article in accordance with the Journal policy described in the Instructions for Authors (http://www.plantphysiol.org) is: Hsou-min Li (mbhmli@gate.sinica.edu.tw).
ABSTRACT
Toc12 is a novel J-domain-containing protein identified in pea chloroplasts. It was shown to be an integral outer-membrane protein localizing in the intermembrane space of the chloroplast envelope. Furthermore, Toc12 was shown to associate with an intermembrane space Hsp70, suggesting that Toc12 is important for protein translocation across the chloroplast envelope. Toc12 shares a high degree of sequence similarity to Arabidopsis DnaJ-J8, which has been suggested to be a soluble protein of the chloroplast stroma. Here we isolated genes encoding DnaJ-J8 from pea and found that Toc12 is a truncated clone of one of the pea DnaJ-J8s. Protein import analyses indicate that Toc12 and DnaJ-J8s possess a cleavable transit peptide and are localized in the stroma. Arabidopsis mutants with T-DNA insertions in the DnaJ-J8 gene show no defect in chloroplast protein import. Implications of these results in the energetics and mechanisms of chloroplast protein import are discussed.
INTRODUCTION

Most chloroplast proteins are encoded by the nuclear genome and synthesized in the cytosol as higher molecular weight precursors with an N-terminal extension known as the transit peptide. Precursor proteins are imported into chloroplasts through a translocon complex located at the chloroplast envelope. Translocon components associated with the outer membrane are called Toc (translocon of the outer envelope membrane of chloroplast) proteins and those associated with the inner membrane are called Tic (translocon of the inner envelope membrane of chloroplast) proteins. Cleavage of the transit peptide from the precursor by a specific stromal processing peptidase during translocation results in the production of the lower molecular weight mature protein. Various translocon components have been assigned functions in the basic steps of the import process (for reviews, see Inaba and Schnell, 2008; Jarvis, 2008; Li and Chiu, 2010). For example Toc159 (the number indicates the calculated molecular mass of the protein) and Toc34 are receptors for the transit peptides and Toc75 is the protein-translocating channel across the outer membrane. Toc64, on the other hand, has a dual function: it serves as a docking site for the cytosolic Hsp90 through its cytosolic domain, and as a scaffold for translocon components located in the intermembrane space through its intermembrane-space domain (Qbadou et al., 2007).

Protein import into chloroplasts involves at least two distinct ATP consuming steps. The first step is called “early import intermediate” or “docking”, in which less than 100 µM ATP is required and precursors are translocated across the outer membrane and come into contact with translocon components in the inner membrane (Olsen et al., 1989; Kouranov and Schnell, 1997; Inaba et al., 2003; Inoue and Akita, 2008). It has been shown that the ATP is used in the intermembrane space (Olsen and Keegstra, 1992), most likely by a yet unidentified intermembrane-space Hsp70 called imsHsp70 or Hsp70-IAP (ims for “intermembrane space” and IAP for “import intermediate associated protein”; Marshall et al., 1990; Schnell et al., 1994; Qbadou et al., 2007). The second ATP consuming step is the complete translocation of precursors across the two envelope membranes into the stroma. This step requires about 1 mM ATP. The ATP is most likely used by the stromal Hsp93 and cpHsc70 associated with the translocon to drive protein translocation into the stroma (Nielsen et al., 1997; Shi and Theg, 2010; Su and Li, 2010).
Hsp70 family proteins are involved in many cellular processes including protein folding, protein translocation across membranes and regulation of protein degradation. Hsp70 proteins are often recruited to perform a certain function by specifically localized J-domain contain proteins. The J-domain containing proteins interact with Hsp70 when Hsp70 is bound to ATP and stimulates ATP hydrolysis by Hsp70. The specific J-domain containing co-chaperone that recruits the stromal cpHsc70 to the inner envelope membrane to assist in protein translocation has not been identified. The specific J-domain containing co-chaperone for 3msHsp70 for its function in protein import into chloroplasts is proposed to be a protein named Toc12 (Becker et al., 2004).

Toc12 was identified as a novel J-domain containing protein from pea chloroplasts. It belongs to the type III J-domain proteins containing only the J domain without the glycine- and phenylalanine-rich domain (G/F domain) and the zinc-finger domain originally found in *E. coli* DnaJ. It has been shown that the protein is synthesized at its mature size of 103 amino acids without a cleavable transit peptide. After import, the protein has been shown to anchor in the outer membrane by its N-terminal part, which has been suggested to form a β-barrel type domain. Its C-terminal part, composed of the J domain, has been shown to localize in the intermembrane space. Toc12 has been shown to associate with 3msHsp70. Toc12 and 3msHsp70 interact with the intermembrane space domain of Toc64, which in turn associates with another intermembrane space translocon component, Tic22. It is proposed that the Toc12-3msHsp70-Toc64-Tic22 complex mediates protein translocation across the intermembrane space through specific precursor binding and ATP hydrolysis (Becker et al., 2004; Qbadou et al., 2007). However, the existence of 3msHsp70 has only been shown on immunoblots by its reactivity to the monoclonal antibody SPA820 raised against human Hsp70. Its encoding gene has never been identified. The Arabidopsis Hsp70 gene family has 14 members. Only two of them are localized in chloroplasts and both have been shown to locate in the stroma (Ratnayake et al., 2008; Su and Li, 2008). A recent study has further shown that the major protein recognized by the SPA820 antibody in pea chloroplasts is located in the stroma, indicating that 3msHsp70 is most likely a stromal protein (Ratnayake et al., 2008).

Most translocon components were originally identified from pea chloroplasts. While all translocon components identified from pea have easily recognizable
Arabidopsis homologs, Toc12 seems to be an exception. The Arabidopsis gene suggested to be the pea TOC12 homolog, At1g80920 (Inoue, 2007; Jarvis, 2008), encodes a protein that is much larger than pea Toc12 and is annotated as J8 (referred to as AtJ8 herein). The entire pea Toc12 has a high sequence similarity to the N-terminal two thirds of AtJ8. AtJ8 contains an extra C-terminal domain of 60-amino-acid that is highly conserved among J8 proteins from other higher plants. However, in contrast to pea Toc12, AtJ8 is predicted to locate in the stroma (Miernyk, 2001 and www.arabidopsis.org). Indeed a fusion protein consisting of the first 80 amino acids of AtJ8 fused at the N terminus of GFP was imported into the chloroplast stroma and approximately 46 amino acids from the N terminus was processed after import (Lee et al., 2008), indicating that the first 46 amino acids of AtJ8 function as a cleavable stroma-targeting transit peptide. A T-DNA insertion in the AtJ8 gene that causes the truncation of the last three amino acids results in no visible phenotype. However detailed analyses indicate that the mutant has a lower CO₂ assimilation and Rubisco activity than the wild type (Chen et al., 2010).

We are interested in identifying J-domain-containing proteins interacting with stromal Hsp70. As part of the initial effort, we investigated the sub-organelar location of J8 and examined the relationship between Toc12 and J8. We found that, in pea, there are at least two genes encoding J8, which we named PsJ8a and PsJ8b. TOC12 represents part of PsJ8b. Toc12, AtJ8 and the two PsJ8 proteins could be imported into chloroplasts and processed to stromally localized soluble mature proteins. Four alleles of AtJ8 mutants were analyzed but none of them showed any defect in the import of various chloroplast precursor proteins.
RESULTS

Toc12 is transcribed from one of the two genes encoding PsJ8

Through multiple sequence alignment we found that pea Toc12 is extremely similar to
the N-terminal two thirds of J8 homologs from various legume species (Fig. 1). For
example, Toc12 is 78.8% identical to the N-terminal two-thirds of *Medicago truncatula*
J8. However, all J8s have an extra C-terminal domain about 60 amino acids in length.
Part of this domain is highly conserved from legumes to Arabidopsis (Fig. 1). Instead of
this C-terminal domain, Toc12 has a short tail of five amino acids not found in any other
species.

To isolate J8-encoding genes from pea, we took advantage of the conservation of
the C-terminal domain. In J8s of *Medicago*, castor bean and soybean, the last 18 bases
before the stop codon have identical DNA sequences (Supplemental Fig. S1). Using this
conserved stretch of sequence to design a reverse primer, part of the 5’ UTR and the first
11 bases of pea *TOC12* ORF to design a forward primer (Supplemental Table S1, primers
PsJ8-R2 and PsJ8-F1, respectively) and first-strand cDNA reverse-transcribed from total
pea RNA as templates, we obtained an approximately 450-bp cDNA fragment by PCR
amplification. Sequencing revealed that the PCR product was actually a mixture of two
different cDNAs (Supplemental Fig. S2), which we designated as *PsJ8a* and *PsJ8b*.
Clone specific primers were then designed to isolate their corresponding genomic clones
and to perform 3’ RACE to identify the authentic cDNA sequence encoding the C-
terminal end and 3’ UTR.

Sequence comparison revealed that *PsJ8a* has nucleotide differences from *PsJ8b*
throughout the cDNA sequence and resulted in 12 amino-acid changes, including a two-
amino-acid insertion near the N terminus of PsJ8b (the alanine and glycine of the 7th and
8th residues in J8b, Fig. 2A, arrowhead). Most strikingly, however, *PsJ8b* contains in the
second intron a 1229-bp insertion that is absent in *PsJ8a* (Fig. 2B, red line).

Toc12 is identical to the N-terminal two thirds of PsJ8b in both protein (Fig. 2A)
and cDNA (Supplemental Fig. S2) sequences except in two places. Residues 34 and 36
are both threonine in Toc12 but arginine in PsJ8a and PsJ8b (Fig. 2A, red box). The last
six amino acids and the 3’ UTR of Toc12 are encoded by the sequence corresponding to
the beginning of the second intron of *PsJ8b*, including part of the 1229-bp insertion
unique to \textit{PsJ8b} (Fig. 2B and 2C). This result suggests that \textit{TOC12} RNA arises from partially or alternatively spliced \textit{PsJ8b}. The difference of arginine vs. threonine at residues 34 and 36 may be attributed to different pea varieties used. Using a primer located immediately after the stop codon of Toc12 in the second intron, we could indeed amplify a cDNA with the C-terminal sequence of Toc12 but residues 34 and 36 were still arginines. Using site-directed mutagenesis, we created a cDNA clone encoding the exact protein sequence of the published Toc12 by changing residues 34 and 36 to threonine.

\textbf{AtJ8 and PsJ8s are localized in the stroma}

We first investigated the sub-organellar location of AtJ8 and the two PsJ8s using in vitro protein import and fractionation. [\textsuperscript{35}S]Met-labeled J8 proteins were incubated with isolated pea chloroplasts under import conditions. AtJ8 was synthesized as an approximately 21-kD protein (Fig. 3A, AtJ8 panel, lane 1). After import into chloroplasts, a smaller protein of approximately 16 kD was produced. Chloroplasts after import were further treated with thermolysin or trypsin. Thermolysin can only degrade proteins exposed on the cytosolic side of the outer envelope membrane, while trypsin can digest proteins in the intermembrane, as it can penetrate the outer but not the inner envelope membrane (Cline et al., 1984; Jackson et al., 1998). The effectiveness of the trypsin treatment was shown by the degradation of the outer-membrane protein Toc75 and the resistance of Tic110 (Fig. 3A, lanes 5 and 6), which is an inner-membrane protein with a large stromal domain. The 21-kD AtJ8 was degraded by thermolysin and trypsin but the 16-kD protein produced after import was resistant to both proteases, indicating that the 16-kD protein was inside the inner envelope membrane. Thus it is likely that the 21-kD in-vitro-translated protein was the precursor form and the 16-kD protein was the imported mature form after removal of the transit peptide. A difference of 5-kD agrees well with the ChloroP prediction and the experimental result (Lee et al., 2008) that the first 46 amino acids functions as a cleavable transit peptide.

Trypsin-treated chloroplasts were further lysed hypotonically and separated into membrane and soluble fractions. Most of the 16-kD mature AtJ8 was found in the soluble fraction and a small fraction was associated with the membranes (lanes 7 and 8). However, mature AtJ8 was entirely observed in the soluble fraction when chloroplasts
were treated by alkaline extraction (lanes 9 and 10). These results indicate that the 16-kD mature AtJ8 protein is a soluble stromal protein with a small portion peripherally associated with membranes. Control proteins the small subunit of ADP-glucose pyrophosphorylase (APS) and Tic110 were localized to the soluble and membrane fraction, respectively.

The two PsJ8s were also translated as 21-kD proteins but a protein with a size slightly larger than 16 kD was also produced, most likely due to internal initiations from one of the downstream methionines (Fig. 3A, lane 1 of the PsJ8 panels). To check whether this smaller protein could be imported into chloroplasts, we abolished translation initiation from the first methionine of PsJ8b by mutating the initiation ATG codon into ATT and generated the mutant clone PsJ8b-M1x. In vitro transcription/translation of PsJ8b-M1x indeed produced a protein that is the same size as the internal initiation product synthesized from PsJ8b (Fig. 3B, lanes 1 and 2). Incubation of PsJ8b-M1x with chloroplasts under import conditions resulted in PsJ8b-M1x associating with chloroplasts but the proteins remained entirely thermolysin sensitive (Fig. 3B, lanes 4 and 5), indicating that PsJ8b-M1x was only sticking to the chloroplast surface and could not be imported.

Import of the two PsJ8s into chloroplasts produced essentially the same results as those from AtJ8 import. For both PsJ8 proteins, a 16-kD mature protein resistant to both thermolysin and trypsin was produced after import. When trypsin treated chloroplasts were lysed hypotonically, most of the imported 16-kD mature proteins were in the soluble fraction and a small fraction was associated with the membrane fraction (Fig. 3B, lanes 7 and 8). When chloroplasts were lysed by alkaline extraction, all of the imported 16-kD mature PsJ8s were in the soluble fraction (Fig. 3B, lanes 9 to 10). Results in Fig. 3 indicate that AtJ8 and the two PsJ8s are synthesized as higher molecular weight precursors with a cleavable transit peptide and that the imported mature proteins are soluble and localized in the stroma.

**Toc12 is processed to a 7-kD mature protein localized in the stroma**

Toc12 is essentially identical to the N-terminal two thirds of PsJ8b. Therefore if PsJ8b has a cleavable transit peptide, Toc12 should possess an identical transit peptide except
residues 34 and 36 are threonine in Toc12 but arginine in PsJ8b. Yet, PsJ8b is a soluble protein in the stroma and Toc12 was shown to be an integral outer-membrane protein. The transit peptide region of PsJ8b is the region predicted to be a β-barrel type membrane anchor in Toc12. To determine whether the sequence differences in PsJ8b and Toc12 have resulted in their different localizations, we reinvestigated the localization of Toc12. [35S]Met-labeled Toc12 was incubated with isolated pea chloroplasts under import conditions. Chloroplasts after import were treated with thermolysin. It was reported that Toc12 was completely thermolysin resistant after import due to its localization in the intermembrane space (Becker et al., 2004). However, we found that about 30% of Toc12 was already thermolysin resistant even before import (Fig. 4, lanes 1 to 5). After incubation with chloroplasts, the same percentage of Toc12 was thermolysin resistant. The percentage of Toc12 that was thermolysin resistant remained unchanged even after the thermolysin concentration was increased to 400 µg/ml or after the addition of 0.1% Triton X-100 in the digestion to permeabilize the chloroplasts (Fig. 4, lanes 6 to 10). Under these conditions, Toc159, an outer membrane protein with a large cytosolic domain, was degraded by the lowest concentration of thermolysin without Triton X-100 addition. In comparison, Tic110 became accessible to thermolysin only after Triton X-100 addition, confirming the effectiveness of the thermolysin and Triton X-100 treatments (Fig. 4, the immunoblots panels). These results suggest that some Toc12 molecules are intrinsically thermolysin resistant probably due to their tight folding or aggregation, not due to protection by the outer membrane. As a result, thermolysin is not a suitable protease for determining the location of Toc12.

We noticed that if Toc12 has a transit peptide like PsJ8b and the processing of Toc12 occurs at the corresponding site to the one predicted for AtJ8 (Fig. 1, arrow), the mature Toc12 produced after removal of the transit peptide would contain no methionine residues and thus would not be visible in autoradiograph. We therefore added two methionine residues at the C terminus of Toc12 by site-directed mutagenesis and named this protein Toc12MM. When [35S]Met-labeled Toc12MM was incubated with isolated pea chloroplasts under import conditions, a 7-kD protein was produced (Fig. 5, lane 5). This 7-kD protein was produced only when 5 mM ATP was added to the import reaction and was not produced when no ATP was added (Fig. 5, lane 6), suggesting that it was the
mature protein produced after import into chloroplasts. However, possibly due to its tight folding and tendency to aggregate, the import efficiency of Toc12MM was very low. Under identical conditions, no band was visible in the same region of the Toc12 import samples either with or without ATP addition (Fig. 5, lanes 2 and 3). In a control reaction using the precursor to the small subunit of RuBP carboxylase (prRBCS), mature RBCS was only produced when ATP was added, confirming the effectiveness of ATP addition (Figure 5, lane 8).

To identify the location of the 7-kD protein produced after import of Toc12MM, chloroplasts after import were treated with trypsin. As shown in Fig. 6A, after import into chloroplasts, the 7-kD protein produced after import was trypsin resistant, indicating that it was inside the inner membrane. When a high amount of in-vitro-translated Toc12 before import was analyzed, an internal initial product very similar in size to the 7-kD mature protein could be observed (Fig. 6A, lane 1). This internal initiation product was completely trypsin sensitive, further supporting that the 7-kD trypsin-resistant mature protein was produced after importing into chloroplasts. The same import samples were analyzed by immunoblotting using antibodies against Toc75 and Tic110. Toc75 was trypsin sensitive and Tic110 was trypsin resistant, indicating that trypsin had indeed penetrated the outer but not the inner membrane.

To further analyze whether Toc12MM and the 7-kD mature protein are integral membrane proteins or soluble proteins, chloroplasts after import of Toc12MM were lysed and separated into membrane and soluble fractions. Import of prRBCS and Toc12 was performed as controls. A lower concentration of ATP was present in the prRBCS reaction in order to produce a higher amount of prRBCS associated with the envelope membranes to serve as an envelope-protein control. When chloroplasts were lysed hypotonically, the 7-kD protein produced after import of Toc12MM was present both in the membrane and the soluble fractions. However, when chloroplasts were treated by alkaline extraction, all the 7-kD mature protein was in the soluble fraction. Most Toc12MM and Toc12 fractionated with the membranes when chloroplasts were lysed hypotonically but more than half of Toc12MM and Toc12 were in the soluble fractionation when chloroplasts were lysed by alkaline extraction (Fig. 6B, lane 4). This membrane association pattern of Toc12MM and Toc12 was similar to that of prRBCS bound to the chloroplast envelope,
partially translocated through the translocon (Fig. 6B, prRBCS panel). In comparison, Toc75, a protein with a β-barrel type membrane anchor, was not affected by the alkaline extraction and was entirely in the membrane fraction. These results suggest that the 7-kD protein produced after import of Toc12MM is a soluble protein with a fraction peripherally associated with membranes. Together with its trypsin resistance, our results indicate that this 7-kD protein is located in the stroma.

It has been shown that a commercially available anti-DnaJ antibody can also recognize Toc12 (Becker et al., 2004). We used the same antibody in order to know the relative position of endogenous Toc12 to PsJ8. We used the antibody on immunoblots of purified outer membrane proteins, and total chloroplast soluble and membrane proteins. However, perhaps due to different immunoblotting conditions, we failed to detect specific proteins in these fractions (Supplemental Fig. S4). A few bands were detected after a longer exposure but they were also detected by the anti-Toc75 antibody and on a control blot probed with the secondary antibody alone.

**Mutations in AtJ8 result in no import defect**

To further investigate the possible function of J8 in chloroplast protein import, we obtained four Arabidopsis mutants with T-DNA insertions in the *AtJ8* gene (Fig. 7A). The *j8-1* mutant (Salk_024617) is the same allele as previously reported (Chen et al., 2010). The *j8-1*, *j8-2* (Gabi _922G05) and *j8-4* (WiscDxLoxHs005M07G) alleles are in the Col ecotype and the *j8-3* (Flag_349D04) allele is in the Ws4 ecotype. The *j8-1* and *j8-2* alleles have a T-DNA insertion close to the C terminus of AtJ8 (Fig. 7A and Fig. 1). The *j8-3* and *j8-4* alleles have a T-DNA insertion in the promoter region 80 and 85 bp upstream of the transcription start site, respectively. RT-PCR using primers on one side of the T-DNA insertion sites (F4 + R6, Fig. 7A) indicated that the *j8-1*, *j8-3* and *j8-4* alleles had a very small amount of *AtJ8* RNA left and the *j8-2* allele still had a substantial amount of *AtJ8* RNA (Fig. 7B, top panel). When using a pair of primers located on two sides the T-DNA insertion sites of *j8-1* and *j8-2* (F4 + R5, Fig. 7A), no *AtJ8* transcript could be detected in *j8-1* or *j8-2* (Fig. 7B). Primers were further designed to quantify the amount of *AtJ8* RNA in the mutants using real-time quantitative RT-PCR. When using a pair of primers located on one side of the T-DNA insertion sites (Q-F1 + Q-R1, Fig. 7A),
the j8-1, j8-3 and j8-4 alleles had about 26%, 21% and 16% of AtJ8 transcript left, respectively, compared to their corresponding wild types. The j8-2 allele still had about 89% of AtJ8 transcript left (Fig. 7C). However, all four mutants appear indistinguishable from the wild types (Fig. 7D). They also have the same chlorophyll and carotenoid contents as those in the wild type (Supplemental Fig. S3), similar to the reported data for j8-1 (Chen et al., 2010).

Chloroplasts were isolated from 14-day-old seedlings of the four j8 mutants and their corresponding wild types. They were incubated with chloroplast precursor proteins under import conditions. Three precursors were tested: prRBCS, the precursor to the nucleus-encoded chloroplast 50S ribosomal subunit L11 (prL11), and the precursor to the thylakoid lumen protein plastocyanin (prPC). As shown in Fig. 8, for all three precursors tested, the import efficiency of the four mutants was similar to that of their corresponding wild types.

DISCUSSION
We show here that the pea TOC12 gene is part of the PsJ8b gene. Furthermore, all four proteins, AtJ8, PsJ8a, PsJ8b and Toc12, are synthesized as higher molecular weight precursors with cleavable transit peptides. After import into chloroplasts, the processed mature proteins are localized in the stroma. A fraction of the imported mature proteins was associated with membranes but was extracted to the soluble fraction when chloroplasts were treated with alkaline extraction. This tendency to associate with membranes is similar to the other two stromal J proteins investigated, Arabidopsis J11 (ATJ11, Chen et al., 2010) and pea PCJ1 (Schlicher and Soll, 1997).

Several experiments were performed to investigate the localization of Toc12 when it was first identified (Becker et al., 2004). In-vitro-translated, [35S]Met-labeled Toc12 was imported into isolated chloroplasts and shown to be thermolysin resistant after import, suggesting that Toc12 was located inside the outer membrane. However, the in-vitro-translated [35S]Met-Toc12 before import was not tested for its thermolysin resistance. We show here that a fraction of Toc12 was intrinsically thermolysin resistant even before import. The amount of Toc12 that was thermolysin resistant did not increase...
after import. The fraction of Toc12 that was thermolysin resistant remained resistant even in the presence of Triton X-100. Therefore, thermolysin resistance cannot be used as an indication that Toc12 was inside the outer membrane. In the original report, a fusion protein of Toc12 fused at the N terminus of GFP (Toc12-GFP) was transiently expressed in tobacco protoplasts. No microscopy data was provided for the location of Toc12-GFP. Chloroplasts isolated from the transfected protoplasts were analyzed by immunoblots and Toc12-GFP was shown to be trypsin sensitive, suggesting that Toc12-GFP was outside the inner membrane. However, in the same experiment, Tic110 was also trypsin sensitive. Although there are still some debates about the topology of Tic110, all models agree that the large majority of the Tic110 polypeptide is localized in the stroma (Jackson et al., 1998; Balsera et al., 2009). Therefore it is possible that in the original experiment, trypsin was not properly quenched and had gained access into the stroma. Nonetheless, Toc12 was identified by ESI MS/MS spectrometry from isolated pea chloroplast outer membrane proteins. Although only one peptide was identified, the peptide was located close to the C terminus of Toc12 and contains two amino acids unique to Toc12. An antibody generated against the entire length of Toc12, including the transit peptide region, recognized a single protein around 14 kD on immunoblots when isolated pea chloroplast outer membrane vesicles were analyzed (Becker et al., 2004). Therefore it is possible that TOC12 RNA is an alternatively spliced form of PsJ8b, leading to the generation of Toc12 protein in pea. If this is the case, it seems to be unique to pea. No EST corresponding to a similar alternatively-spliced form of J8 has been found in Arabidopsis or in other species. Whole genome tiling arrays of Arabidopsis also indicate that the second intron is not transcribed (http://signal.salk.edu/cgi-bin/atta, Yamada et al., 2003). During the revision of this manuscript, a review article was published (Schwenkert et al., 2010) and indicated that a thorough analysis of a pea EST database revealed that the pea Toc12 should be the same length as that of all the J8 proteins from other species (Fig. 2 of Schwenkert et al., 2010). The revised Toc12 assembled from the pea EST database is essentially the same as our PsJ8b, which we show here is a stromally localized soluble protein (Fig. 3A).

J-domain containing proteins are important cochaperones for the Hsp70 family proteins. The J domain interacts with Hsp70 when Hsp70 is bound to ATP and stimulates
ATP hydrolysis by Hsp70. Specific J-domain proteins often recruit Hsp70 to perform specific functions. It has recently been shown that chloroplast stromal Hsp70s are important for protein import into chloroplasts (Shi and Theg, 2010; Su and Li, 2010). Because J8 is localized in the stroma, we investigated whether J8 is involved in protein import by analyzing the j8 mutant phenotype. We analyzed four T-DNA insertion mutant alleles. Two of the alleles, j8-3 and j8-4, exhibited a significant decrease (~80%) in AtJ8 transcript. The other two alleles, j8-1 and j8-2, do not possess any full-length AtJ8 transcript. However, all four mutants did not show any defect in importing various chloroplast precursor proteins, suggesting that J8 is unlikely to be involved in protein import. A recent report indicated that the j8-1 mutant has a lower CO2 assimilation and a reduced Rubisco activity (Chen et al., 2010), suggesting that J8 may be a cochaperone for the activity of Hsp70 in folding and assembly of enzymes in the carbon fixation reaction. However since the mutants we analyzed might still produce a low level of truncated J8 proteins, we cannot exclude the possibility that the highly conserved C-terminal region is not required for activity of J8 in protein import.

It has been generally assumed that the ATP required for the binding/early-import-intermediate step is consumed by the intermembrane space Hsp70 imsHsp70. Now that imsHsp70 and its cochaperone Toc12 have been shown to be most likely localized in the stroma, the only currently known intermembrane space components of the translocon are Tic22 and the intermembrane space domain of Toc64. Neither of these two components possesses any ATPase activity. Therefore, the proteins that use the ATP in the binding step remain unknown. It needs to be pointed out that, although it has been generally assumed that an ATPase like an Hsp70 facilitates the formation of the early import intermediates, it has been shown that GTP can also supports the formation of early import intermediates without being converted to ATP (Olsen and Keegstra, 1992). When supplied with 2.5 mM GTP as the sole energy source at room temperature, the transit peptide plus a part of the mature region of precursors can be transported across the outer membrane. Interestingly, this transport is inhibited by ATPγS (Inoue and Akita, 2008). Therefore it is possible that the translocon component that supports the formation of early import intermediates may be a novel intermembrane-space NTPase with a broad nucleotide specificity that can use ATP or GTP, but with a preference for ATP. This
inference makes the Hsp70 family proteins unlikely candidates as the component driving transport across the outer membrane. However, it is also possible that multiple NTPases, including the Hsp70 family proteins, are involved in the formation of early import intermediates. In summary, the linkage between the Toc and Tic complexes and the process of precursor translocation across the outer membrane and the intermembrane space remain to be elucidated.

MATERIALS AND METHODS

Isolation of J8 and Toc12 cDNA and genomic clones
First-strand cDNAs were synthesized using the M-MLV reverse transcriptase (Invitrogen) with RNA isolated from pea (Pisum sativum cv. Little Marvel) or Arabidopsis leaves. Primers used to amplify the PsJ8 cDNA and genomic DNA, and the AtJ8 cDNA were listed in Supplemental Tables S1 and S2. PCR-amplified fragments were subcloned into vectors pGEM-T or pSP72 (Promega). Site-directed mutagenesis to create Toc12 and Toc12MM was performed on plasmids containing the coding sequence for PsJ8b using the QuikChange site-directed mutagenesis kit (Stratagene) and primers listed in Supplemental Table S1. 3’RACE was performed according to the manufacture’s protocol of 5’/3’ RACE Kit (Roche).

Chloroplast protein import and post-import treatments
Arabidopsis seedlings were grown on 0.3% Gelrite-solidified Murashige and Skoog (MS) medium containing Gamborg’s B5 vitamin and 2% sucrose. Plants were grown in growth chambers under a 16-h photoperiod at 22°C with a light intensity about 80 μmol m⁻² s⁻¹. For growing pea seedlings (Pisum sativum cv Little Marvel; De Bruyn Seed Store), imbibed seeds were grown on vermiculite under a 12-h photoperiod at 20°C with a light intensity about 150 μmol m⁻² s⁻¹. Chloroplasts were isolated from 9-day-old pea seedlings or 14-day-old Arabidopsis seedlings as described (Perry et al., 1991), except the grinding buffer for Arabidopsis chloroplast isolation was modified to 50 mM HEPES-KOH (pH 8.0), 330 mM sorbitol, 2 mM EDTA, and 0.5% bovine serum albumin. Synthesis of [³⁵S]Met-labeled precursors through in vitro transcription and in vitro translation using
wheat germ lysate (Promega), protein import into chloroplasts and treatment of chloroplasts with thermolysin after import were performed as described (Perry et al., 1991). Reticulocyte lysates cannot be used for translating Toc12 because Toc12 migrated at the same position as hemoglobin that is sometime produced in reticulocyte lysates. In-vitro-translated Toc12 tends to aggregate and was therefore centrifuged at 20,000g for 15 min to remove large aggregates before used for further experiments. For import under ATP depleted conditions, ATP was removed from the in-vitro-translated precursors by gel filtration as described (Perry et al., 1991) and import was performed under a green-safe light. Hypotonic lysis of chloroplasts after import was performed by resuspending reisolated intact chloroplasts in 25 mM Hepe-KOH (pH 8.0) and 4 mM MgCl2. Alkaline-extraction was performed by resuspending reisolated intact chloroplasts after import in 0.1 M Na2CO3 (pH 11.5) and incubated for 30 min at 4°C. Hypotonically-lysed and alkaline-extracted samples were then separated into membranes and soluble fractions by ultracentrifugation at 100,000g for 45 min. Soluble fractions were precipitated in 10% trichloroacetic acid, washed with ice-cold acetone and dissolved in SDS-PAGE sample buffer. Trypsin treatment of chloroplasts after import was performed as described (Jackson et al., 1998). Immunobloting was performed as described (Su and Li, 2010) using specific primary antibodies and an alkaline phosphatase-conjugated secondary antibody, and visualized by the NBT-BCIP colorimetric system, except for Supplemental Fig. S4, which was visualized using the horseradish peroxidase–conjugated secondary antibodies and the ImmobilonWestern Chemiluminescent HRP system (Millipore), visualized with the UVP BioSpectrum 600 Image System (Ultra Violet Products). Antibodies against Arabidopsis Tic110 and Toc159 and pea Toc75 were generated as described (Tu et al., 2004). The antibody against E. coli DnaJ (ADI-SPA-410) was purchased from Stressgen (Enzo).

**Isolation of T-DNA-tagged Arabidopsis j8 mutants**

Arabidopsis mutants with T-DNA insertion in AtJ8 genomic region were obtained from various sources. The j8-1 (Salk_024617, Alonso et al., 2003) and j8-4 mutants (WiscDxLoxHs005M07G, Nishal et al., 2005) were obtained from the Arabidopsis Biological Resource Center (http://abrc.osu.edu/). The j8-2 mutant (Gabi _922G05,
Rosso et al., 2003) was obtained from Nottingham Arabidopsis Stock Centre (http://arabidopsis.info/). The j8-3 mutant (Flag_349D04, Samson et al., 2002) was obtained from French National Institute for Agricultural Research (http://www.inra.fr/).

Primers for identification of T-DNA insertions in *AtJ8* genomic region are: AtJ8geno-F2, AtJ8geno-R2, and various left border primers. Left border primers used are Salk-LBa1 for *j8-1*, Gabi-LB for *j8-2*, Flag-LB4 for *j8-3* and Wisc-L4 for *j8-4*. Primer sequences are listed in Supplemental Table S1. T-DNA insertion sites were confirmed by sequencing.

**RT-PCR, quantitative RT-PCR and chlorophyll content analyses of *j8* mutants**

Total RNA was isolated using TRIzol reagent (Invitrogen) from Arabidopsis leaves. First-strand cDNAs were synthesized using total RNA, M-MLV reverse transcriptase (Invitrogen), and a poly (dT) primer. For RT-PCR (Fig 7B), cycle numbers for amplification were 20 for *UBQ10*, and 25 for *AtJ8*. Real-time quantitative RT-PCR was performed with the LightCycler System (Roche) and the LightCycler-FastStart DNA Master SYBR Green I kit (Roche). For each PCR reaction, 10 to 50 ng of cDNA and 0.5 µM of primer pairs were used. The initial denaturing step of 10 min was followed by 40 cycles of 95°C for 10 s, 60°C for 5 s, and 72°C for 1 s per 25 bp of the expected product. Standard curve based on serial dilutions of the wild-type cDNA for each primer pair was included. The absence of nonspecific products was checked by melting curve analysis. Quantification was performed using the LightCycler Relative Quantification software version 1.0. Normalization was done using the transcript level of *UBQ10*. Primers used to amplify each transcript are listed in Supplemental Tables S1 and S2. Chlorophyll and carotenoid contents were determined by the method of Lichtenthaler (1987).

Sequence data from this article can be found in the GenBank/EMBL data libraries under accession numbers AY357119 (pea Toc12), BT051272 (J8 of *Medicago truncatula*), XM_002512691 (J8 of castor bean *Ricinus communis*), BT094680 (J8 of soybean Glycine max), NP_178207 or At1g80920 (AtJ8 of *Arabidopsis thaliana*), HM565932 (PsJ8a cDNA of pea *Pisum sativum*), HM565933 (PsJ8b cDNA), HM565934 (PsJ8a genomic DNA), and HM565935 (PsJ8b genomic DNA).
ACKNOWLEDGMENTS

We thank Dr. Jychian Chen for the anti-APS antibody, Dr. Heiko Kuhn for English editing, Dr. Yi-shin Su for the purified pea chloroplast outer-membrane vesicles and Drs. Bo Liu and Yuh-Ru Lee for helping us purchasing the anti-DnaJ antibody. This work was supported by grants from the National Science Council (NSC 99-2321-B001-015, H-mL) and Academia Sinica (H-mL) of Taiwan.
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FIGURE LEGENDS

Figure 1. Toc12 is highly similar to the N-terminal two thirds of J8 from various legume species.
Sequence alignment of pea Toc12 and J8 from *Medicago* (MtJ8), soybean (GmJ8), castor bean (RcJ8), and *Arabidopsis* (AtJ8). The predicted transit peptide processing site of AtJ8 is indicated with an arrow. The conserved tripeptide HPD in the J domain is underlined. T-DNA insertion sites of *Arabidopsis* *j8-1* and *j8-2* mutants are indicated with open arrowheads.

Figure 2. Deduced amino acid sequences and genomic structure of the two *PsJ8* genes.
(A) Toc12 is part of PsJ8b. Sequence alignment of PsJ8a, PsJ8b and Toc12. The additional alanine and glycine residues present in PsJ8b is indicated with an arrowhead. Threonines at residues 34 and 36 of Toc12 substituted with arginines in PsJ8a and PsJ8b are boxed in red. Stop codon is marked as an asterisk.
(B) *TOC12* is a partially spliced or alternatively spliced RNA of *PsJ8b*. Exons and introns are shown as black arrows and blue lines, respectively. A unique 1229-bp insertion in the second intron of *PsJ8b* is shown as a red line. *TOC12* cDNA is represented by the pink line. Sequences of the beginning and end of the boxed region are shown in (C).
(C) Detailed sequences of the beginning and end of the boxed region marked in (B). Nucleotide sequences (same color code as in B) and deduced amino acid sequences of *PsJ8b* and *TOC12* are shown.

Figure 3. AtJ8, PsJ8a and PsJ8b are synthesized as higher molecular weight precursors with cleavable transit peptides and imported into the chloroplast stroma.
(A) Import of AtJ8, PsJ8a and PsJ8b into isolated pea chloroplasts. In-vitro-translated *[35S]Met-labeled precursor proteins (Ivt) and chloroplasts after import of the precursor proteins (Chpt) were treated with thermolysin or trypsin. Trypsin-treated chloroplasts were lysed hypotonically (hypo) or treated by alkaline extraction (alka), and then separated into membrane (M) and soluble (S) fractions. Samples were analyzed by SDS-
PAGE followed by fluorography (for AtJ8 and PsJ8s) or immunoblotting (for Tic110, Toc75 and APS). pr, precursor form; m, mature form. The Ivt lanes contained 1% of the in-vitro-translated proteins used for the import reactions shown in the Chpt lanes. (B) PsJ8b-M1x could not be imported into chloroplasts. In-vitro-translated [35S]Met-PsJ8b-M1x (Ivt, lanes 2 and 3) and chloroplasts after import of PsJ8b-M1x (Chpt, lanes 4 and 5) were treated with thermolysin. All samples were analyzed by SDS-PAGE and fluorography. In-vitro-translated PsJ8b (lane 1) was also analyzed for size comparison.

**Figure 4. A fraction of Toc12 is intrinsically resistant to thermolysin digestion.**
In-vitro-translated [35S]Met-Toc12 before import (Ivt) and chloroplasts after import of Toc12 (Chpt) were treated with 200, 300, or 400 µg/ml of thermolysin or with 200 µg/ml thermolysin plus 0.1% Triton X-100. Samples were analyzed by SDS-PAGE followed by fluorography (for Toc12) or immunoblotting (for Toc159 and Tic110). Lanes 1 to 5 (Ivt) contained 3.6% of the in-vitro-translated Toc12 used for chloroplast import shown in lanes 6 to 10. Toc159 is easily degraded into various fragments during chloroplast isolation. The full-length Toc159 is indicated by the arrow and the major 86-kD degradation fragment is indicated with the asterisk.

**Figure 5. Toc12MM can be processed into a 7-kD protein after import into chloroplasts.**
In-vitro-translated [35S]Met-Toc12, [35S]Met-Toc12MM and [35S]Met-prRBCS were incubated with isolated pea chloroplasts in the presence (lanes 2, 5, and 8) or absence (lanes 3, 6, and 9) of 5 mM ATP. In-vitro-translated proteins (Ivt) and chloroplasts after the import reactions (Chpt) were analyzed by SDS-PAGE and fluorography. Lane 1, 4 and 7 represent 0.8%, 0.24% and 0.24% of the in-vitro-translated proteins used for the corresponding import experiments, respectively.

**Figure 6. The 7-kD protein produced after import of Toc12MM is localized in the stroma.**
(A) In-vitro-translated [35S]Met-Toc12MM (Ivt) and chloroplasts after import of Toc12MM (Chpt) were treated with trypsin. Samples were analyzed with SDS-PAGE
followed by fluorography (for Toc12MM) or immunoblotting (for Toc75 and Tic110). The Iv lanes contained 0.4% of the in-vitro-translated Toc12MM used for the import reactions shown in the Chpt lanes.

(B) Fractionation of Toc12MM, Toc12 and prRBCS after import into chloroplasts. Chloroplasts after import of Toc12MM, Toc12 and prRBCS were lysed hypotonically or treated by alkaline extraction and then separated into membrane (M) and soluble (S) fractions. Samples were analyzed by SDS-PAGE followed by fluorography (for imported proteins) or immunoblotting (for Toc75). For Toc12MM and Toc12 import reactions, 5 mM ATP was added. For the prRBCS import reaction, no additional ATP was supplied. Import reactions were performed under light and the in-vitro-translated protein products contained a low amount of ATP from the in vitro translation system.

**Figure 7. Characterizations of four Arabidopsis j8 mutants.**

(A) Schematic representation of T-DNA insertion sites of the j8 mutants. Filled and open boxes represent the translated and untranslated exon regions, respectively. LB, left border of the T-DNA. Positions of primers used for RT-PCR shown in (B) and real-time quantitative RT-PCR shown in (C) are indicated by arrows. Primers R6 and Q-R1 span the junction between the second and the third exons.

(B) RT-PCR analyses of AtJ8 transcripts in the j8 mutants. The primers used are indicated in parentheses on the right. Parental ecotype of the j8-1, j8-2 and j8-4 alleles are Col. The j8-3 allele is in the Ws4 ecotype. The amount of UBQ10 transcripts was analyzed as a control.

(C) Real-time quantitative RT-PCR analyses of the AtJ8 transcript levels in the j8 mutants. The AtJ8 transcript level was first normalized to the level of UBQ10 in each sample and the expression level in wild type was then set as 100%. Primers for AtJ8 detection are Q-F1 and Q-R1 as shown in (A). Data are means ± S. D., n=2.

(D) Phenotype of the j8 mutants. Arabidopsis seedlings were grown on MS medium under 16 h light/8 h dark for 14 days.

**Figure 8. The Arabidopsis j8 mutants show no detectable defect in chloroplast protein import.**
(A) In-vitro-translated \(^{35}\text{S}\)Met-prRBCS, -prL11 and -prPC were imported into chloroplasts isolated from the \(j8\) mutants and their corresponding wild types. Samples were analyzed by SDS-PAGE. The gel was first stained with Coomassie blue and then dried for fluorography. The amount of endogenous RBCS as revealed by Coomassie blue staining is shown below the fluorograph of the same gel. Iv, in-vitro-translated precursor protein.

(B) Quantification of imported mature proteins shown in (A). The amount of imported mature protein was first normalized to the amount of endogenous RBCS of the same sample. The import efficiency of the wild type was then set as 100%. Data are means ± S. D., \(n=3-6\).

Supplemental Data

Supplemental Figure S1. Alignment of \(J8\) cDNA sequences from \(Medicago\) (\(MtJ8\)), soybean (\(GmJ8\)) and castor bean (\(RcJ8\)). Position of the \(PsJ8-R2\) primer used for initial amplifications of \(PsJ8\) from pea is shown as a blue line with an arrowhead indicating the direction of the primer. Stop codons of \(J8\) cDNAs are marked.

Supplemental Figure S2. Sequence alignment of \(PsJ8a\) and \(PsJ8b\) cDNA and \(TOC12\).
Sequences of \(PsJ8a\) and \(PsJ8b\) from the start codon to the poly-A tail are shown. Stop codons of \(PsJ8s\) and \(TOC12\) are indicated in red and blue, respectively. The junction between the second and the third exons is also indicated.

Supplemental Figure S3. Pigment contents of Arabidopsis \(j8\) mutants.
Chlorophyll and carotenoid contents of 14-day-old wild type and \(j8\) mutants grown on MS medium were measured. Data are means ± S. D., \(n=8\).

Supplemental Figure S4. The antibody against \(E.\ coli\ DnaJ\) failed to detect specific proteins in various fractions from pea chloroplasts.
Total membrane (M) and soluble (S) fractions (35 \(\mu g\) of protiens in each lane), and the purified outer (O) and inner (I) envelope membranes (12.5 \(\mu g\) of proteins in each lane)
from pea chloroplasts were analyzed by SDS-PAGE followed by immunoblotting with rabbit antibodies against *E. coli* DnaJ (Stressgen, SPA-410), pea Toc75, and Arabidopsis Tic110, or with a rabbit non-immune serum. Proteins were visualized using the horseradish peroxidase–conjugated secondary antibodies and the ImmobilonWestern Chemiluminescent HRP system (Millipore), with the UVP BioSpectrum 600 Image System (Ultra Violet Products).

(A) Shorter exposure time of blots.

(B) Longer exposure time of blots. The 20-kD band recognized by all rabbit antibodies in the outer membrane is marked by an asterisk.

Supplemental Table S1. Sequences of primers used in this study.

Supplemental Table S2. Primers used for amplification of PsJ8s and Toc12 cDNA and genomic DNA and site-directed mutagenesis.
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