

# A SecY Homologue Is Required for the Elaboration of the Chloroplast Thylakoid Membrane and for Normal Chloroplast Gene Expression

Laura M. Roy and Alice Barkan

Institute of Molecular Biology, University of Oregon, Eugene, Oregon 97403

**Abstract.** Results of in vitro and genetic studies have provided evidence for four pathways by which proteins are targeted to the chloroplast thylakoid membrane. Although these pathways are initially engaged by distinct substrates and involve some distinct components, an unresolved issue has been whether multiple pathways converge on a common translocation pore in the membrane. A homologue of eubacterial SecY called cpSecY is localized to the thylakoid membrane. Since SecY is a component of a protein-translocating pore in bacteria, cpSecY likely plays an analogous role. To explore the role of cpSecY, we obtained maize mutants with transposon insertions in the corresponding gene.

Null cpSecY mutants exhibit a severe loss of thylakoid membrane, differing in this regard from mutants lacking cpSecA. Therefore, cpSecY function is not limited to a translocation step downstream of cpSecA. The phenotype of cpSecY mutants is also much more pleiotropic than that of double mutants in which both the cpSecA- and  $\Delta$ pH-dependent thylakoid-targeting pathways are disrupted. Therefore, cpSecY function is likely to extend beyond any role it might play in these targeting pathways. CpSecY mutants also exhibit a defect in chloroplast translation, revealing a link between chloroplast membrane biogenesis and chloroplast gene expression.

THE majority of chloroplast-localized proteins are associated with the internal membrane system of the organelle, the thylakoid membrane. Many thylakoid membrane proteins are products of nuclear genes and are synthesized in the cytoplasm; others are products of chloroplast genes and are synthesized in the chloroplast stroma. In either case, newly synthesized proteins must be targeted to their correct position with respect to the membrane. Mechanisms of targeting to the thylakoid lumen have been particularly well studied. All known luminal proteins are nuclear-encoded and are synthesized with a bipartite targeting sequence (for review see Cline and Henry, 1996). The NH<sub>2</sub>-terminal segment is a stromal targeting sequence that directs the protein across the chloroplast envelope. Adjacent to the stromal targeting sequence is a cleavable luminal targeting sequence that resembles the signal sequences that target proteins for translocation across bacterial cytoplasmic membranes. Two energetically and genetically distinct pathways have been described for the translocation of proteins to the thylakoid lumen (for review see Cline and Henry, 1996). One set of luminal proteins is targeted by a mechanism that requires cpSecA, a chloroplast-localized homologue of the bacte-

rial protein SecA (Yuan et al., 1994; Nohara et al., 1995; Voelker and Barkan, 1995; Voelker et al., 1997). Cytochrome *f*, a chloroplast-encoded integral thylakoid protein, is also targeted to the membrane via this pathway (Voelker and Barkan, 1995; Mould et al., 1997; Nohara et al., 1997; Voelker et al., 1997). A second set of luminal proteins is targeted via a pathway that relies upon a *trans*-thylakoidal  $\Delta$ pH in vitro (for review see Cline and Henry, 1996) and that involves the *hcf106* (Voelker and Barkan, 1995) and *tha4* genes (our unpublished results) in vivo. A third pathway has been proposed for several integral membrane proteins that appear to integrate spontaneously in vitro (Kim et al., 1996; Robinson et al., 1996). Finally, the integration of the polytopic membrane protein light-harvesting chlorophyll *alb* binding protein (LHCP),<sup>1</sup> whose targeting signals lie within the mature portion of the protein (Viitanen et al., 1988), defines a fourth pathway in that it requires neither cpSecA (Voelker et al., 1997) nor *hcf106* (Voelker and Barkan, 1995), but does require GTP (Hoffman and Franklin, 1994) and cp54 (Li et al., 1995), a homologue of the signal recognition particle protein SRP54.

1. *Abbreviations used in this paper:* LHCP, light-harvesting chlorophyll *alb* binding protein; OE16, 16-kD subunit of oxygen evolving complex; OE23, 23-kD subunit of oxygen evolving complex; OE33, 33-kD subunit of oxygen evolving complex; PC, plastocyanin; Rubisco, ribulose biphosphate carboxylase; SRP, signal recognition particle; *csy1*, chloroplast secY 1.

Address all correspondence to Alice Barkan, Institute of Molecular Biology, University of Oregon, Eugene, OR 97403. Tel.: (541) 346-5145. Fax: (541) 346-5891.

These previous experiments indicated that each targeting pathway involves some unique components. It is possible, however, that the different targeting machineries also have shared components. For example, two or more of the pathways may converge on a common translocation pore in the membrane, in analogy to the convergence of signal recognition particle (SRP)-dependent and -independent pathways on Sec61-containing translocons in the endoplasmic reticulum membrane (for review see Rapoport et al., 1996). One likely component of the translocon for proteins that engage cpSecA is cpSecY, a chloroplast-localized SecY homologue, since SecY forms a component of the translocon in the bacterial plasma membrane (for review see Rapoport et al., 1996). Genes encoding cpSecY have been discovered in algal chloroplast genomes (for review see Vogel et al., 1996) and in the *Arabidopsis thaliana* and spinach nuclear genomes (Laidler et al., 1995; Berg-hoefler and Kloesgen, 1996). The protein encoded by the *Arabidopsis* cpSecY cDNA is targeted to chloroplast thylakoid membranes in vitro (Laidler et al., 1995). However, functional studies of cpSecY have not been reported.

To gain insight into the roles of cpSecY in vivo, we have used a reverse genetics strategy to obtain maize mutants with transposon insertions in a nuclear gene encoding cpSecY. As shown below, cpSecY mutants exhibit a severe loss of thylakoid membrane. This phenotype is much more severe and global than that of *tha1* mutants, which have exceedingly low levels of cpSecA (Voelker et al., 1997). Therefore, cpSecY function is not limited to a role in translocating proteins that previously engaged cpSecA. Furthermore, cpSecY mutants have a more severe phenotype than double mutants with lesions in both the cpSecA and the  $\Delta$ pH luminal targeting pathways, implicating cpSecY in either the cp54-dependent pathway, the "spontaneous" pathway, and/or in an uncharacterized targeting pathway. An unexpected aspect of the cpSecY mutant phenotype is a global defect in chloroplast translation. Thus, the activity of the chloroplast translation machinery may be linked to the biogenesis of chloroplast membranes.

## Materials and Methods

### Plant Material

Plants with *Mu* insertions in the gene encoding cpSecY (*csy1*) were identified at Pioneer Hi-Bred (Johnston, IA) by using a primer complementary to the *Mu* terminal inverted repeat (Bensen et al., 1995; Meeley and Briggs, 1995) and four primers specific for the maize cpSecY cDNA. Pioneer Hi-Bred generously provided a small number of progeny seed from plants that scored positive in the PCR screen. *csy1*<sup>+</sup> plants were propagated by crossing with inbred lines and subsequent self-pollination to recover homozygous mutant seedlings.

Also used in this study were the targeting mutants *tha1* (Voelker and Barkan, 1995; Voelker et al., 1997), *hcf106* (Barkan et al., 1986; Voelker and Barkan, 1995) and *tha4*. The *tha4* and *hcf106* mutant phenotypes are very similar (i.e., the  $\Delta$ pH thylakoid targeting pathway is specifically disrupted [see Fig. 8, Voelker and Barkan, 1995]) but the two genes are not allelic. The *hcf106-mum3* allele, a deletion derivative of the original *hcf106* mutation (Das and Martienssen, 1995), was used in these experiments; seed was provided by R. Martienssen (Cold Spring Harbor Laboratory, Cold Spring Harbor, NY). Some experiments included the albino mutant *iojap*, which has been described as lacking in plastid ribosomes (Walbot and Coe, 1979), and the albino mutant *w3*, which has a defect in the carotenoid biosynthetic pathway (Fong et al., 1983).

Unless otherwise stated, studies were performed with leaf material obtained from seedlings grown for 9–12 d in a growth chamber (14-h d [400

$\mu$ E/m<sup>2</sup> per s) at 28°C and 10-h nights at 25°C). Samples that were compared in any single experiment were grown and harvested in parallel. Etiolated mutant leaves were obtained as described previously (Voelker et al., 1997).

### Cloning of Maize cpSecY cDNAs

A degenerate primer (5'-GAA[GAT]AT[GAT]AT[AGCT]GGCAT-3') was designed to a conserved amino acid region (met-pro-ile-ile-phe-ser) of cpSecY and then used to prime reverse transcription of maize seedling leaf poly A<sup>+</sup> RNA by following the manufacturer's protocol for Superscript<sup>TM</sup> II reverse transcriptase (GIBCO BRL, Gaithersburg, MD). PCR amplification was performed on the cDNA template using the reverse transcription primer and a second degenerate primer (5'-GT[A/G/C/T]CC[A/G/C/T]TTCAT[A/T/C]AA[C/T]GC-3' encoding val-pro-phe-ile-asn-ala). Each 50- $\mu$ l reaction contained 1  $\mu$ l of the reverse transcription reaction, 2.0 pmol of each primer, 20  $\mu$ M deoxynucleoside triphosphate, 1.5 mM MgCl<sub>2</sub>, 50 mM KCl, and 10 mM Tris-HCl, pH 8.3. The reactions were incubated at 95°C for 5 min, supplemented with 5 U of Taq DNA polymerase, and then incubated according to the following "touchdown" profile: 10 cycles of 94°C/30 s, 54°C/40 s (–1.0°C/cycle), 72°C/60 s; 10 cycles of 94°C/30 s, 44°C/40 s (–0.5°C/cycle), 72°C/60 s; 20 cycles of 94°C/30 s, 40°C/40 s, 72°C/60 s. These reactions were subjected to a secondary amplification involving nested degenerate primers that included a 5' EcoRI site (5'-GGAATTCAT[A/T/C]AA[C/T]GC[A/T/G/C]CA[A/G]AT[A/T/C]GT-3' encoding ile-asn-ala-gln-ile-val) and a 5' BamHI site (5'-CGGGATCCAT[G/A/T]AT[A/G/C/T]GGCAT[A/T/C/G]AC[A/C/T/G]CC-3' encoding gly-val-met-pro-ile-ile), according to the following conditions: 20 cycles of 95°C/30 s, 48°C/40 s (–0.5°C/cycle), 72°C/60 s; 20 cycles of 95°C/30 s, 39°C/40 s, 72°C/60 s). The amplification product was gel-purified and then subcloned into a Bluescript SK<sup>+</sup> plasmid. The isolated insert was used to screen a cDNA library constructed from mRNA isolated from the leaves of 2-wk-old greenhouse-grown maize, inbred line B73.

### Sequence Alignments

Sequences of SecY homologues from *Arabidopsis thaliana* (EMBL/GenBank/DBJ accession number U37247), *Spinacia oleracea* (accession number Z54351), *Escherichia coli* (accession number 134413), *Synechococcus* PCC7942 (accession number 401077), cryptomonas (accession number X62348), and *Pyrenomonas salina* (accession number X74773) were compared to the maize *csy1* cDNA sequence (accession number AF039304). Alignments were calculated using ClustalW 1.7 (Thompson et al., 1994) and BoxShade (Bioinformatics Group, ISREC, Lausanne, Switzerland), with default parameters.

### Isolation and Analysis of Plant DNA and mRNA

DNA was extracted and then analyzed as previously described (Voelker et al., 1997). The genomic Southern blot was probed with a partial *csy1* cDNA, corresponding to amino acids 205–387 (see Fig. 1) that was radiolabeled by the random hexamer priming method. RNA was extracted and analyzed by Northern hybridization as described in Jenkins et al. (1997). The association of chloroplast mRNAs with polysomes was assayed as described previously (Barkan, 1993). The *csy1* transcript was detected with the full-length *csy1* cDNA radiolabeled by random hexamer priming. The 16S rRNA and LHCP mRNAs were detected with the radiolabeled DNA probes described in Barkan (1993). The *rbcs5* mRNA was detected with a radiolabeled DNA probe prepared from the cDNA described in Nelson et al. (1984). The *atpF* probe was described in Jenkins et al. (1997). The *rbcl* mRNA was detected with a radiolabeled RNA probe prepared by in vitro transcription of a plasmid containing a 570-bp PstI fragment of the maize *rbcl* gene.

### Extraction and Analysis of Protein and Chlorophyll

Leaf proteins were extracted, analyzed on immunoblots, radiolabeled in vivo, and then immunoprecipitated by using the antibodies and methods described previously (Voelker and Barkan, 1995). The relative chlorophyll content of mutant and normal leaves was determined by excising a 35-mg segment from the tip of 10-d-old seedlings, grinding in a mortar and pestle in the presence of 2 ml of 80% acetone, centrifuging at 12,000 g for 1 min, and then measuring the absorbance of the supernatant at 652 nm.

### Electron Microscopy

Leaf tissue was fixed overnight at room temperature in 4% glutaraldehyde in 0.05 M potassium phosphate, pH 7.0. After three washes in phos-

phate buffer, the tissue was postfixed in 2% osmium tetroxide in the same buffer. Samples were dehydrated in an ethanol series and then embedded in Spurr's resin. Sections were stained for 50 min in 5% uranyl acetate and for 10 min in Reynolds lead citrate (Reynolds, 1963). The tissue preparation, sectioning, and electron microscopy was performed by J. Selker (University of Oregon Electron Microscopy Facility, Eugene, OR).

## Results

### The Maize Genome Contains Several Genes That Are Closely Related to *secY*

We took advantage of the Trait Utility System for Corn developed at Pioneer Hi-Bred (Meeley and Briggs, 1995) to obtain plants with *Mu* transposons in a nuclear gene encoding cpSecY. A prerequisite for this approach was knowledge of the nucleotide sequence of the maize cpSecY gene(s). To obtain maize cDNAs encoding cpSecY, the deduced *Arabidopsis*, spinach, *Cryptomonas*, and *Pyrenomonas* cpSecY proteins were compared to identify regions of high conservation. A degenerate primer designed against one conserved region was used to prime reverse transcription of maize leaf polyA+ RNA. A second degenerate primer was then used in conjunction with the first to amplify a PCR product. The amplified cDNA fragment was of the expected size (550 bp) and was related in sequence to known cpSecY genes (data not shown). When the PCR fragment was used to screen a maize leaf cDNA library, several full-length cDNA clones (2.1 kbp) were obtained whose DNA sequence revealed a continuous open reading frame encoding a protein with high identity to plastid and eubacterial SecY proteins (Fig. 1).

*Arabidopsis* and spinach cpSecY are synthesized with an NH<sub>2</sub>-terminal extension of ~120 amino acids, relative to the bacterial and plastid-encoded SecY homologues (Fig. 1). This region has characteristics typical of chloroplast targeting signals (for review see Cline and Henry, 1996). The maize cDNAs encode a protein with an NH<sub>2</sub>-terminal extension of a similar length and with features of chloroplast targeting signals. Downstream of this region, the deduced maize protein is highly similar to other chloroplast-localized SecY homologues (e.g., 94% similar/86% identical to *Arabidopsis* cpSecY). As shown below, the phenotypes associated with mutations in the gene encoding the maize cDNA are restricted to the chloroplast. Together, these observations provide strong evidence that the cDNA encodes maize cpSecY. The gene encoding this cDNA was named *csy1*.

Genomic Southern blots were probed at high stringency with a cDNA fragment encoding a highly conserved region of cpSecY (Fig. 2). Multiple bands were detected when the genomic DNA was digested with enzymes that do not cut within the genomic sequence corresponding to the probe. These results revealed the existence of at least two and perhaps three genes in the maize genome that are closely related to the cDNA clone. To estimate the relative contributions of these different genes to the pool of cpSecY in leaf tissue, the same probe was used to identify clones in a maize leaf cDNA library. 18 independent cDNA clones were obtained. The nucleotide sequences of the 3' untranslated regions were identical for all 18 clones, although their sites of polyadenylation varied (Fig. 3 c). A probe made from this 3' untranslated region detected only

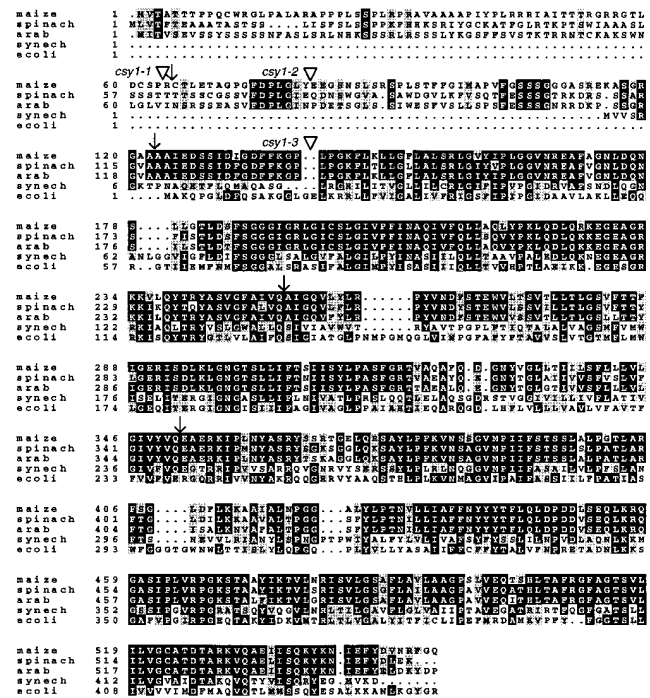
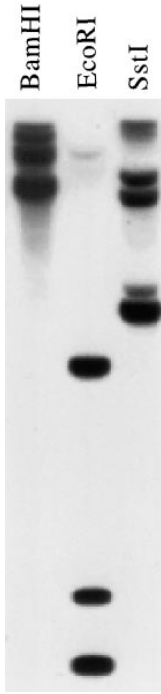


Figure 1. Deduced amino acid sequence of a maize cDNA encoding a SecY homologue. The *Arabidopsis* (*Arab*) and spinach proteins are chloroplast localized (Laidler et al., 1995; Berghoefer and Kloesgen, 1996). The *Synechococcus* (*syn*) and *E. coli* sequences are representative of bacterial SecY. Each triangle indicates the site of the *Mu* transposon insertion in the mutant allele named to its left. Positions of introns are indicated by arrows.

one band on genomic Southern blots (data not shown), indicating that all of the cDNAs were derived from the same gene. These results strongly suggest that the *csy1* gene contributes the vast majority of cpSecY in leaf tissue.

### *Mu* Insertions Cause Severe Defects in the Expression of *csy1*

The progeny of maize plants with *Mu* transposons in *csy1* were tentatively identified at Pioneer Hi-Bred in a PCR screen involving *csy1*-specific primers used in conjunction with a primer corresponding to the *Mu* terminal inverted repeat (Bensen et al., 1995; Meeley and Briggs, 1995). We confirmed that three of these putative alleles had *Mu* insertions in *csy1*. To define the precise position of each insertion, the PCR products resulting from amplification with a *Mu* primer and a *csy1* primer were cloned and then their nucleotide sequences were determined. Fig. 3 shows a partial map of the *csy1* gene, including the location of the *Mu* insertion in each mutant allele. *csy1-1* contains a *Mu* insertion at a 5' splice junction, in sequences encoding the presumed chloroplast targeting signal. *csy1-2* has a *Mu* insertion in sequences encoding the targeting signal, and *csy1-3* has a *Mu* insertion in sequences encoding the highly conserved portion of cpSecY. As is typical of nuclear-encoded chloroplast proteins, an intron lies between sequences encoding the presumed targeting sequence and sequences encoding the portion of the protein that was derived from the ancestral bacterial gene (refer to Fig. 1).



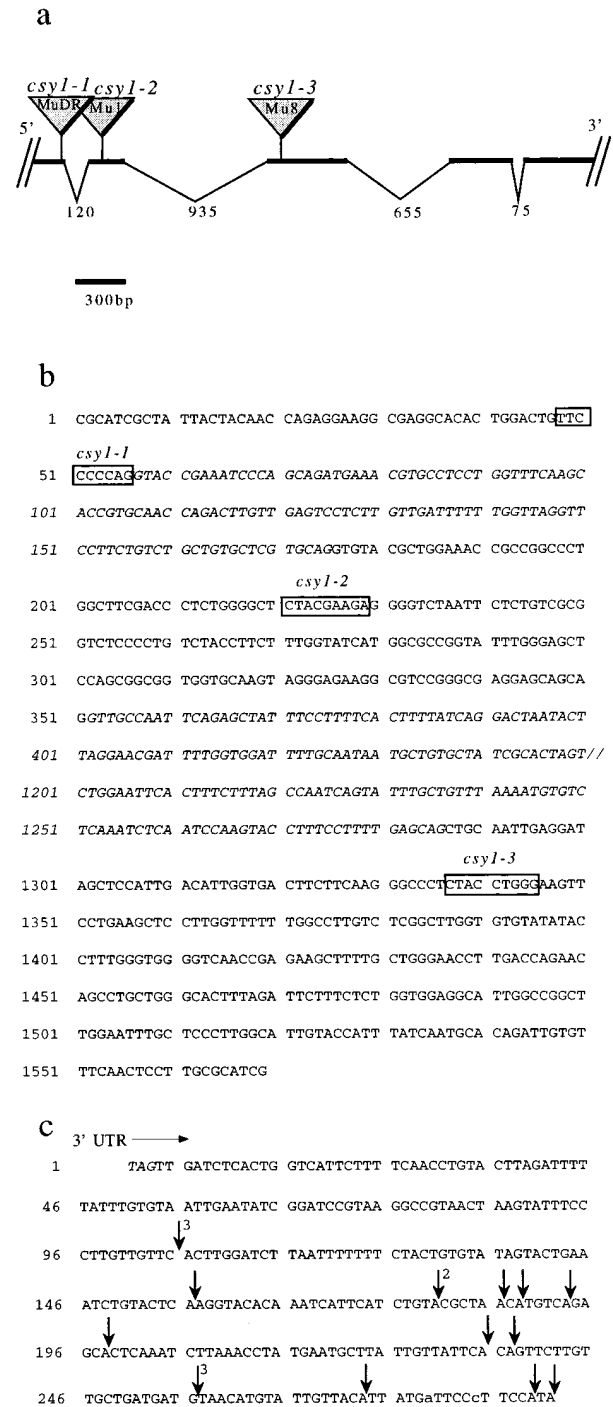
**Figure 2.** Genomic Southern blot of DNA from the maize inbred line Tx303 probed with the *csy1* cDNA. The probe consisted of the cDNA fragment encoding amino acids 208–381. BamHI and EcoRI do not recognize sites inside the genomic region corresponding to the probe. The filter was washed at high stringency (65°C in 0.3× SSC).

The insertions in *csy1* are genetically linked to the pigment-deficient phenotype shown in Fig. 4. This pigment deficiency did not segregate from *csy1* in >50 meioses. Homozygous mutants are nearly albino, with a distinct yellow-green tint. Heterozygous plants appear entirely normal. All three *csy1* mutant alleles are similar in phenotype, confirming that the pigment-deficiency results from the *Mu* insertions in *csy1*. The chlorophyll content of *csy1* mutant seedlings grown under relatively low-intensity light (400 μE/m<sup>2</sup> per s) is ~10% of that of their normal siblings grown in parallel, when normalized to leaf wet weight (data not shown). In contrast, *th1* and *hcf106* mutants, with lesions in the cpSecA and ΔpH thylakoid targeting pathways, respectively, accumulate near normal amounts of chlorophyll (Martienssen et al., 1987, and our unpublished observations).

During the development of their first three leaves, *csy1* mutant plants grow at the same rate as their wild-type siblings (data not shown). The early normal growth rate indicates that mitochondrial function is not significantly disrupted by these mutations and supports the notion that *csy1* function is limited to the chloroplast compartment. The rate of growth of the mutant seedlings began to lag as seed reserves became depleted and they died after the development of four leaves, as is typical of mutations that disrupt photosynthesis in maize (Miles, 1982).

A single *csy1* mRNA of ~2.1 kb was detected in RNA prepared from wild-type leaf tissue (Fig. 5). This mRNA

**Figure 3.** Map and partial nucleotide sequence of *csy1*. (a) Partial genomic map of *csy1*. Sites of *Mu* insertions are indicated. The identity of the *Mu* family member at each site was deduced based upon the sequence of its terminal-inverted repeat (Chandler and Hardeman, 1992). The insertion sites are shown in the context of the encoded amino acid sequence in Fig. 1. The size of each intron in nucleotides is indicated. (b) Partial nucleotide sequence of



*csy1*. The sequence shown begins 135 nucleotides downstream of the start codon and ends within the third exon. Locations of *Mu* insertions are shown by boxes marking the nine base pairs that were duplicated upon insertion. Intron sequence is in italics. Only a portion of the sequence of the second intron is shown, with the missing sequence represented by //. The complete nucleotide sequence of the *csy1* cDNA and more extensive genomic sequence information has been deposited (EMBL/GenBank/DBJ accession numbers AF039304 and AF039305). (c) Sites of polyadenylation in 18 independent leaf *csy1* cDNA clones. The sequence shown begins at the stop codon. The number of cDNA clones with the same polyadenylation site is indicated in superscript. Clones with identical 3' ends differed at their 5' ends, indicating that they represent independent cDNAs.



**Figure 4.** Pigment deficiency of *csy1* mutant seedlings. A *csy1* mutant (left), a wild-type sibling (middle), and an *iojap* mutant (right) were photographed when they were 2-wk old. The *iojap* mutant is shown to illustrate a true albino mutant. All three *csy1* alleles are associated with similar pigmentation defects.

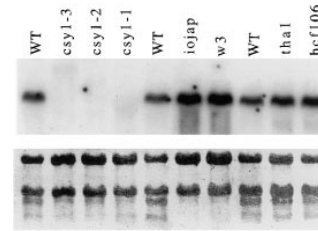
was not detectable in any of the mutant alleles. A low abundance aberrant transcript of ~3.5 kb in *csy1-3* is of the expected size of a chimeric transcript that includes the *Mu8* sequences. Two albino mutants, *iojap* and *w3*, whose albinism results from different primary defects (Robertson et al., 1978; Walbot and Coe, 1979; Han et al., 1992), and two mutants disrupted in thylakoid protein targeting (*tha1* and *hcf106*), were analyzed in parallel. All of these contain normal levels of *csy1* mRNA, indicating that the loss of the mRNA in the *csy1* mutants is not a consequence of their pigment deficiency or of a defect in thylakoid protein targeting. The lack of detectable *csy1* mRNA in the mutants indicates that the mutations are either null or nearly so.

### Chloroplasts in *csy1* Mutants Have Little Thylakoid Membrane

The ultrastructure of *csy1* mutant chloroplasts was examined by transmission electron microscopy (Fig. 6). Both the bundle sheath and mesophyll chloroplasts in mutant leaves contain little internal membrane. Small internal vesicles were frequently observed (Fig. 6 *b* and data not shown). Mitochondrial morphology is not altered in the mutants (Fig. 6, *a-c*).

Etioplasts in *csy1* mutants are also abnormal in structure (Fig. 6 *c*). Whereas normal etioplasts contain a crystalline prolamellar body with radiating membranes, *csy1* mutant etioplasts contain a condensed but noncrystalline prolamellar body that resembles structures observed in maize mutants defective in chlorophyll biosynthesis (Mascia and Robertson, 1978). However, unlike the etioplasts in chlorophyll-biosynthetic mutants, *csy1* mutant etioplasts lack the radiating membranes, indicating that their ultrastructural defects are not simply a consequence of the absence of chlorophyll.

The accumulation of representative proteins from each major thylakoid membrane complex was assayed by immunoblot (Fig. 7 *a*). Lesions in either the  $\Delta$ pH or cpSecA luminal targeting pathway do not significantly reduce the content of the chloroplast ATP synthase or of LHCP (Voelker and Barkan, 1995). In *csy1* mutant leaves, however, the abundance of these proteins is ~10-fold lower



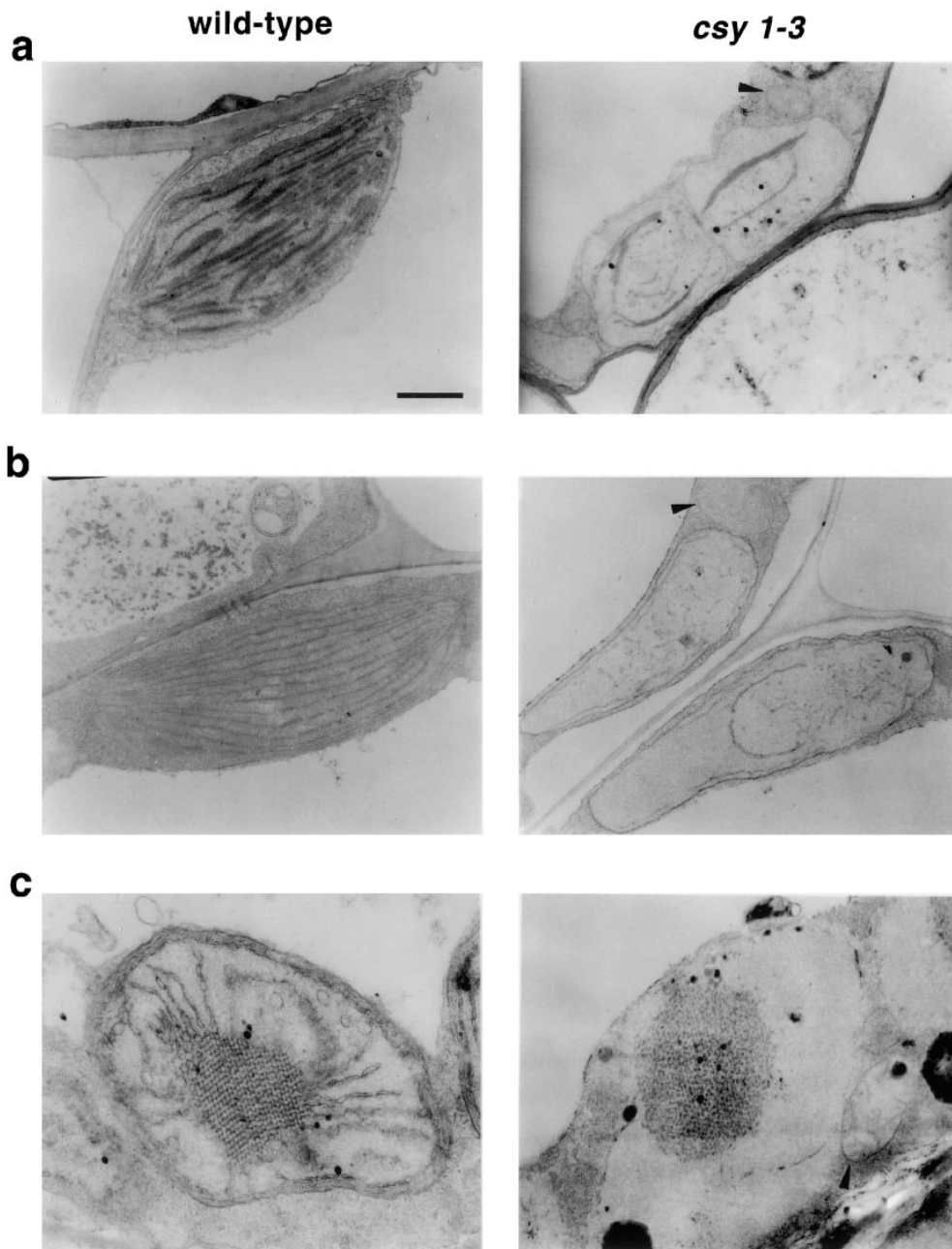
**Figure 5.** RNA gel blot hybridization showing *csy1* mRNA in seedling leaf tissue. Total leaf RNA (10  $\mu$ g) was gel fractionated and then hybridized with a radiolabeled *csy1* cDNA probe (top panel). The same filter was stained with methylene blue to visualize rRNAs (bottom panel). Other mutants were analyzed to illustrate that the *csy1* mRNA defects do not result from defects in thylakoid protein targeting (*tha1* and *hcf106*), as a consequence of pigment deficiencies (*w3* and *iojap*), or as a consequence of an early block in chloroplast development (*iojap*).

than normal. The *psaD* and *petD* gene products and the D1 protein are reduced to <10% of their normal levels in *csy1* mutants. The abundance of these proteins likely reflects the abundance of other subunits of the photosystem I core complex, the cytochrome *b<sub>6</sub>f* complex, and the photosystem II core complex, respectively, since subunits of these complexes are generally reduced coordinately in mutants in which the availability of any single subunit is limiting (for review see Barkan et al., 1995).

The accumulation of luminal proteins in *csy1* mutant leaves was measured to assess the status of the cpSecA and  $\Delta$ pH luminal targeting pathways (Fig. 7 *c*). A disruption in luminal targeting is reflected by a decrease in the abundance of the mature protein coupled with an increase in the abundance of its stromal precursor (Voelker and Barkan, 1995). In *csy1* mutants, the mature form of three substrates of the cpSecA pathway (the 33-kD subunit of the oxygen evolving complex [OE33], plastocyanin [PC], and Psaf) accumulate to <10% of their normal levels, and proteins corresponding in size to their stromal intermediates accumulate to unusually high levels. Two substrates of the  $\Delta$ pH pathway, the 23- and 16-kD subunits of the oxygen evolving complex (OE23 and OE16, respectively), behave analogously. These properties are consistent with the possibility that both the  $\Delta$ pH and cpSecA pathways involve a translocon with a cpSecY subunit. The loss of LHCP is likewise consistent with the idea that cpSecY is required for the integration of proteins that engage cp54. However, it is possible that one or more of these defects is a consequence of the reduction in membrane content, or results because cpSecY is required to target components of other translocons to the membrane.

### cpSecY Mutants Are More Severe in Phenotype Than Mutants Simultaneously Disrupted in Both the cpSecA and $\Delta$ pH Targeting Pathways

Every aspect of the mutant phenotype investigated was more global and severe in *csy1* mutants than in *tha1* and *hcf106* mutants, suggesting that cpSecY function is not limited to either the cpSecA or the  $\Delta$ pH thylakoid targeting pathway. It remained possible, however, that cpSecY functions in both pathways. If so, and if it has no additional functions, the *csy1* mutant phenotype should resemble the phenotype resulting from simultaneous disruption



**Figure 6.** Ultrastructure of *csy1* mutant chloroplasts and etioplasts. (a) Mesophyll chloroplasts. (b) Bundle sheath chloroplasts. (c) Etioplasts. Arrowheads point to mitochondria. Chloroplast ultrastructure was similar in all three mutant alleles (data not shown). Bar, 1  $\mu\text{m}$ .

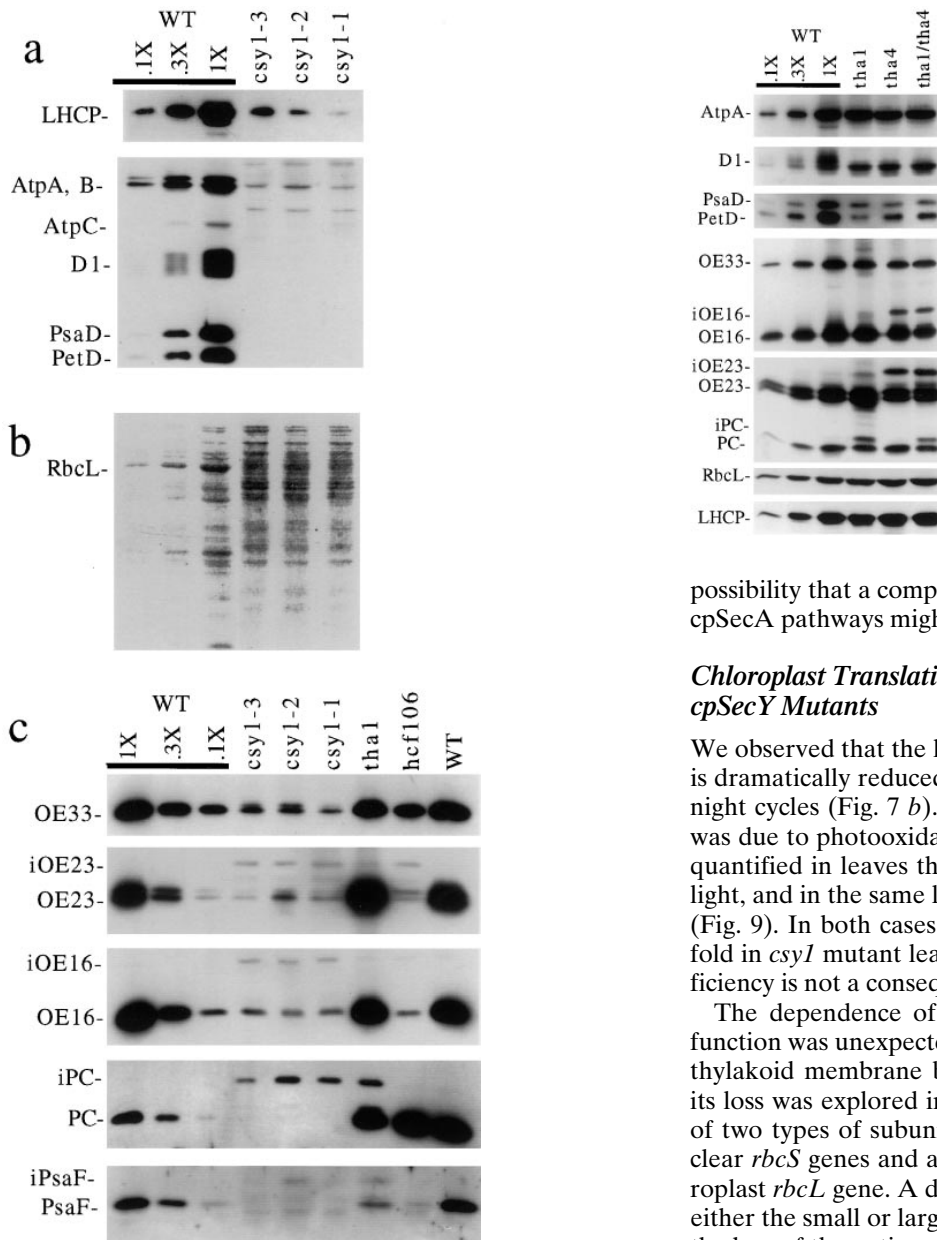
of both the cpSecA and  $\Delta\text{pH}$  pathways. To address this possibility, *tha4/tha1* double mutants were constructed. The *tha1* gene encodes cpSecA and the mutation causes at least a 40-fold decrease in cpSecA accumulation (Voelker et al., 1997). The *tha4* mutation disrupts the  $\Delta\text{pH}$  luminal targeting pathway (our unpublished results and Fig. 8) and its phenotype strongly resembles that of the *hcf106* mutation (Voelker and Barkan, 1995).

Fig. 8 shows that *tha1/tha4* double mutants have a much less severe phenotype than *csy1* mutants. For example, LHCP and RbcL, the large subunit of ribulose biphosphate carboxylase (Rubisco), accumulate normally in the double mutants but are severely reduced in *csy1* mutants (refer to Fig. 7). The accumulation of the luminal proteins OE16 and OE23 (and their stromal precursors) is similar in *tha1/tha4* double mutants to that in *tha4* single mutants.

The accumulation of the luminal proteins OE33 and PC (and their stromal precursors) is similar in *tha1/tha4* double mutants to that in *tha1* single mutants. Components of the ATP synthase, photosystem I, photosystem II, and the cytochrome *b<sub>6</sub>f* complex also accumulate to similar levels in the double and single mutants (Fig. 8, AtpA, PsaD, D1, and PetD, respectively). Finally, the chlorophyll content of double mutant seedlings is  $\sim 50\%$  of that in their normal siblings, and is similar to that of *tha1* and *tha4* single mutants (data not shown).

The fact that the *csy1* mutant phenotype is much more global than that of *tha1/tha4* double mutants suggests that *csy1* gene function is not limited to the cpSecA and  $\Delta\text{pH}$  targeting pathways. This finding is consistent with the possibility that cpSecY is part of the translocon for integrating proteins via the cp54 pathway and/or for the proposed





**Figure 7.** Immunoblot analysis of leaf proteins in *csy1* mutants. 5  $\mu$ g of total leaf protein (or the indicated dilutions of the wild-type [WT] sample) were fractionated by SDS-PAGE and then transferred to nitrocellulose. (a) Accumulation of integral and membrane extrinsic thylakoid proteins. LHCP, PsaD, D1, and PetD were each detected with monospecific antisera. AtpA, AtpB, and AtpC were detected with sera raised against the CF<sub>1</sub> complex. (b) The protein on the blot used in a was visualized by staining with Ponceau S. RbcL is the large subunit of Rubisco. (c) Accumulation of luminal proteins and their precursors. Proteins were detected with the indicated monospecific antisera. iOE23, iOE16, iPC, and iPsaF are of the size predicted for the stromal intermediates of OE23, OE16, PC, and PsaF, respectively. The stromal intermediate of OE33 migrates only slightly slower than the mature form, and can be seen as a doublet in the *csy1* mutant samples.

spontaneous pathway (for review see Cline and Henry, 1996). However, because the *tha4* mutation does not result in the complete loss of *tha4* gene product (Walker, M., and A. Barkan, unpublished results), we cannot eliminate the

**Figure 8.** Immunoblot analysis of leaf proteins in *tha1/tha4* double mutants. The procedures and antibodies were the same as those described in Fig. 7.

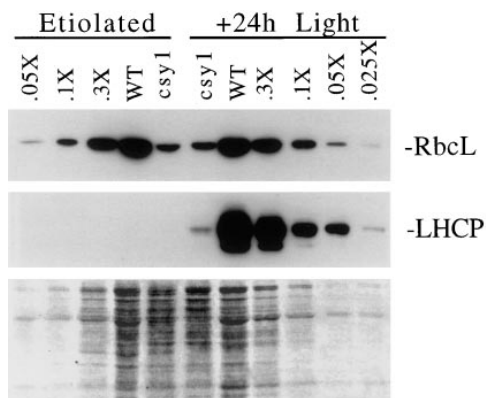
possibility that a complete disruption of both the  $\Delta$ pH and cpSecA pathways might mimic the *csy1* phenotype.

### Chloroplast Translation Is Disrupted in cpSecY Mutants

We observed that the level of the large subunit of Rubisco is dramatically reduced in *csy1* mutants grown under day-night cycles (Fig. 7 b). To address the possibility that this was due to photooxidative damage, the RbcL subunit was quantified in leaves that had developed in the absence of light, and in the same leaves after 24 h of exposure to light (Fig. 9). In both cases, RbcL abundance was reduced 10-fold in *csy1* mutant leaves, indicating that the Rubisco deficiency is not a consequence of photooxidative damage.

The dependence of Rubisco accumulation on cpSecY function was unexpected, given cpSecY's presumed role in thylakoid membrane biogenesis. Therefore, the basis for its loss was explored in some detail. Rubisco is composed of two types of subunits, a small subunit encoded by nuclear *rbcS* genes and a large subunit encoded by the chloroplast *rbcL* gene. A defect in the synthesis or targeting of either the small or large subunit is sufficient to account for the loss of the entire enzyme, since the two subunits accumulate stoichiometrically when the synthesis of one subunit is blocked (Schmidt and Mishkind, 1983; Spreitzer et al., 1985). We considered the possibilities that: (a) *csy1* is required for normal rates of import of proteins into the chloroplast; (b) *csy1* is required for normal accumulation of Rubisco mRNAs; and (c) *csy1* is required for normal chloroplast translation.

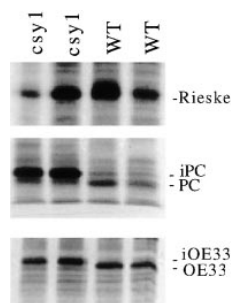
Nuclear-encoded proteins that are imported into the chloroplast engage a common translocation machinery that spans the inner and outer envelope membranes (for review see Schnell, 1995; Cline and Henry, 1996). A defect in import is expected to be reflected by a reduced rate of cleavage of stromal targeting sequences. Rates of processing of three nuclear-encoded chloroplast proteins were monitored by pulse labeling leaf proteins *in vivo*, immunoprecipitation, and then gel fractionation (Fig. 10). There were no detectable differences between wild-type and *csy1* mutant samples in the rates of accumulation of imported forms of the Rieske protein, PC, and OE33. PC and OE33



**Figure 9.** Immunoblot showing loss of the large subunit of Rubisco in both dark- and light-grown *csy1* mutant leaves. Protein was extracted from etiolated *csy1-2* and wild-type seedling leaf tips. The plants were then exposed to light for 24 h and protein was extracted from leaves of the same plants. Total leaf proteins (5  $\mu$ g) or the indicated dilutions of the wild-type samples were fractionated by SDS-PAGE, transferred to nitrocellulose, and then probed with antiserum specific for either RbcL or LHCP. LHCP was analyzed to illustrate that the etiolated plants had not been exposed to light, since it is known that LHCP does not accumulate in leaves grown in the absence of light and that it accumulates rapidly after exposure to light (Nelson et al., 1984). (Bottom) Results of staining the filter with Ponceau S, to illustrate total bound proteins.

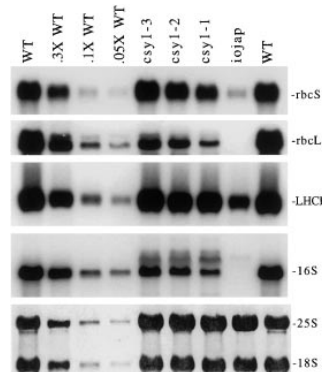
accumulated in their stromal intermediate form rather than in their mature luminal forms in *csy1* mutants, a predicted consequence of a disruption in the cpSecA-dependent luminal targeting pathway. These results provide strong evidence that the *csy1* gene does not function either directly or indirectly in the import of proteins into the chloroplast. They also show that the low levels of PC and OE33 in *csy1* mutants result from an increase in their rates of degradation rather than a decrease in their rates of synthesis.

Mutants in which chloroplast development is blocked at an early stage contain reduced levels of LHCP mRNA and of several other nuclear-encoded mRNAs encoding chloroplast proteins (for review see Taylor, 1989). This has given rise to the notion that the transcription of certain nuclear genes is dependent upon a signal from chloroplasts. To determine whether a disruption of this hypothetical



**Figure 10.** Immunoprecipitation of chloroplast proteins from in vivo-labeled leaf extracts. The results of identical analyses of two mutant and two wild-type seedlings are shown. Seedling leaves were radiolabeled with  $^{35}$ [S]methionine/cysteine for 60 min. Protein containing 300,000 cpm was subject to immunoprecipitation with antisera specific for the chloroplast Rieseke protein, PC, or OE33. Precipitated proteins were fractionated by SDS-PAGE and then the radioactive

proteins were detected by direct autoradiography. iPC and iOE33 are proteins of the size predicted for the stromal intermediates of PC and OE33, respectively. The *csy1-2* allele was used for this experiment.



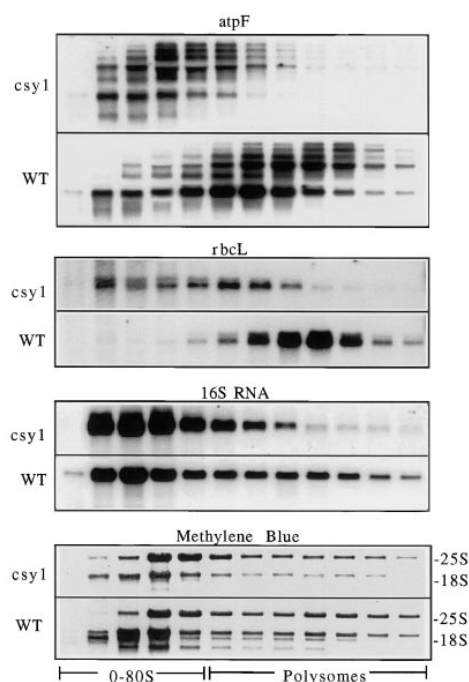
**Figure 11.** RNA gel-blot hybridizations of nuclear and chloroplast transcripts in *csy1* mutants. 5  $\mu$ g (or the indicated dilutions of the wild-type sample) of total leaf RNA were gel fractionated and then hybridized with radiolabeled probes specific for the indicated transcripts. Two identical blots were prepared and then probed sequentially for LHCP and *rbcL* mRNAs, or *rbcS* mRNA and 16S rRNA. (Bottom) Results of staining one of the filters with methylene blue to visualize the bound rRNAs.

signal by *csy1* mutations could be responsible for the Rubisco deficiency, the LHCP and *rbcS* mRNAs were quantified by Northern hybridization (Fig. 11). Both mRNAs accumulate to normal levels in *csy1* mutants. Therefore, the signaling between nucleus and chloroplast appears to be intact in *csy1* mutants and the loss of Rubisco cannot be attributed to a defect in the accumulation of the *rbcS* mRNA.

The abundance of the chloroplast *rbcL* mRNA is reduced four- to fivefold in *csy1* mutants (Fig. 11). Since this was not sufficient to account for the 10-fold decrease in Rubisco protein, the status of *rbcL* mRNA translation was assessed by examining its association with polysomes. Fig. 12 shows that whereas essentially all of the *rbcL* mRNA is associated with polysomes in normal chloroplasts, only a small proportion is polysome associated in *csy1* mutants. The analogous result was obtained for the chloroplast *atpF* mRNAs (Fig. 12) and *psbA* mRNA (data not shown). These findings indicate a decreased rate of translation initiation or a stalling of ribosomes at a site early on the mRNAs examined. Previously it was noted that the abundance of the *rbcL* mRNA is reduced fourfold in maize mutants with defects in chloroplast polysome assembly (Barkan, 1993), and it was proposed that this resulted from increased access of ribonucleases to sites that are ordinarily protected by ribosomes. It seems likely, therefore, that the *rbcL* mRNA deficiency in *csy1* mutants is a consequence of its lack of association with ribosomes rather than an independent manifestation of the *csy1* defect.

To obtain a more global picture of the status of the chloroplast translation machinery, 16S rRNA was quantified (Fig. 11) and its association with polysomes was examined (Fig. 12). The content of 16S rRNA is reduced approximately fivefold in *csy1* mutants, presumably reflecting a similar decrease in the content of 30S ribosomes. Phosphor-Imager quantification revealed little difference between mutant and wild-type samples in the proportion of 16S rRNA found in polysomes, although the average polysome size is reduced in the mutants. The distribution of 16S rRNA in the gradient was less dramatically altered than that of the *rbcL*, *atpF*, and *psbA* mRNAs, suggesting that either the translation of only a subset of chloroplast mRNAs is disrupted in *csy1* mutants, or that the translational block is global but that the timing of the block (i.e., initiation, early or late in elongation) varies between mRNAs.





**Figure 12.** Analysis of chloroplast polysomes in *csy1* mutants. Total leaf extracts prepared under conditions that maintain the integrity of polysomes were fractionated in sucrose gradients. An equal proportion of each fraction was analyzed by RNA gel-blot hybridization with the probes indicated. The methylene blue-stained filters show the distribution of cytosolic rRNAs in the gradients.

## Discussion

### *cpSecY* Function Is Not Limited to the Translocation of Proteins That Engage *cpSecA*

In bacteria, secreted proteins that interact with SecA pass through a translocon consisting of SecY, SecE, and SecG (for review see Rapoport et al., 1996). It seems likely that cpSecY is, by analogy, a component of the thylakoid translocon for proteins that engage cpSecA. The *csy1* mutant phenotype is consistent with this notion. The rate of processing of two cpSecA substrates, PC and OE33, is reduced dramatically in *csy1* mutants (refer to Fig. 10). In addition, a >10-fold decrease in the accumulation of mature PC and PsaF is accompanied by an increase in the accumulation of their stromal intermediates (refer to Fig. 7). However, we cannot eliminate the possibility that the poor translocation of cpSecA substrates is a consequence of the lack of target membrane. Furthermore, if the signal peptidase itself requires cpSecY to cross the membrane, then the rate of signal-sequence cleavage may not be a reliable indicator of a translocation defect.

It is clear from our results, however, that regardless of whether cpSecY functions in the translocation of proteins that engage cpSecA, it must also play other roles. Previously, we found that *thai* mutants contain little, if any, cpSecA, but nonetheless contain normal amounts of thylakoid membrane (Voelker et al., 1997). *thai* mutants also differ from *csy1* mutants in that they contain normal levels of LHCP and Rubisco (Voelker and Barkan, 1995).

Therefore, cpSecY function must extend beyond translocating proteins that previously interacted with cpSecA. Our results are consistent with the possibility that cpSecY is an essential component of the thylakoid Sec translocon, whereas cpSecA is merely a facilitator. Alternatively, it seems plausible that cpSecY mediates the translocation of  $\Delta$ pH-pathway substrates. Indeed, *csy1* mutants exhibit a reduced accumulation of mature  $\Delta$ pH substrates and an increase in the abundance of their stromal precursors. The defects in the  $\Delta$ pH pathway cannot be attributed to a loss of *tha4* and *hcf106* gene products because both accumulate to normal levels in *csy1* mutants (data not shown). On the other hand, the magnitude of the defect in the  $\Delta$ pH pathway as reflected by the immunoblot assay appears less severe than that in the cpSecA pathway (refer to Fig. 7), suggesting that cpSecY may not function directly in this pathway. The notion that defects in  $\Delta$ pH substrate processing are only an indirect effect of the absence of cpSecY is supported by the recent discovery of a bacterial secretory mechanism related to the thylakoid- $\Delta$ pH pathway that does not require SecY in vivo (Santini et al., 1998).

To address the possibility that cpSecY function does not extend beyond the  $\Delta$ pH and cpSecA luminal targeting pathways, we compared the phenotype of cpSecY mutants with that of *thai1tha4* double mutants (Fig. 8). The double mutants differ from *csy1* mutants in that they have a much higher content of chlorophyll and of the core thylakoid membrane complexes, and they are not deficient for LHCP or Rubisco. The much more severe phenotype of *csy1* mutants suggests that cpSecY performs some function in addition to any role it might play in the SecA and  $\Delta$ pH pathways. Sec61p, the endoplasmic reticulum homologue of SecY, functions in the translocation of proteins that engage SRP54 (for review see Rapoport et al., 1996). It seems likely, by analogy, that cpSecY mediates the integration of proteins like LHCP that engage cp54. It is also plausible that cpSecY is required for the integration of proteins typified by CF<sub>0</sub>II, which differ from SecA,  $\Delta$ pH, and cp54 substrates in that they are capable of integration into protease-treated thylakoid membranes (Robinson et al., 1996). Since cpSecY is predicted to be an integral membrane protein with only limited exposure to the stroma (Laidler et al., 1995; Berghoefer and Kloesgen, 1996), it may be resistant to proteases.

### Does *cpSecY* Function in the Integration of Proteins into the Inner Envelope Membrane?

Membrane proteins that are synthesized in the cytosol and imported into the chloroplast stroma have two potential destinations: the thylakoid membrane and the inner envelope membrane. Two routes to the inner envelope can be envisioned for such proteins, a “stop transfer” route in which the protein becomes arrested in the inner envelope during import, and a “conservative sorting” route in which the protein is imported into the stroma and then reexported to the inner envelope by an ancestral secretory system. Recent evidence suggests that TIC110, a nuclear-encoded inner envelope protein, passes through the stroma before its insertion into the envelope membrane, consistent with a conservative sorting mechanism (Luebeck et al., 1997). Should a conservative sorting mechanism exist,

a chloroplast-localized SecY homologue could well be involved. However, the *csyI* gene is clearly not critical for the targeting of all inner envelope proteins because *csyI* mutants are not defective for chloroplast import (refer to Fig. 10), a process that requires several integral components of the inner envelope, including TIC110. In addition, the accumulation of TIC110 protein is unaltered in *csyI* mutants (data not shown). It remains possible that the product of the *csyI* gene functions in the targeting of just a subset of proteins to the inner envelope, or that another of the closely related genes detected by Southern hybridization (refer to Fig. 2) could encode an envelope-localized cpSecY isoform.

### **Role of cpSecY in the Elaboration of the Thylakoid Membrane**

The loss of thylakoid membrane in *csyI*<sup>-</sup> chloroplasts indicates that cpSecY plays a role in membrane accumulation itself. In contrast, the absence of cpSecA in *thal* mutants does not interfere with the elaboration of the thylakoid membrane (Voelker et al., 1997). Many steps in chloroplast-lipid synthesis are localized to the inner envelope membrane (for review see Joyard et al., 1991). Thus, a loss of thylakoid membrane could result if cpSecY functions in integrating lipid biosynthetic enzymes into the inner envelope. Alternatively, cpSecY may play a role in the transfer of lipids from the inner envelope to the thylakoid membrane. There is evidence that thylakoid membranes initially arise by invagination of the inner envelope membrane, followed by vesicle formation and vesicle fusion (for review see Hooper et al., 1994; Cline and Henry, 1996). A stromal protein has been identified that promotes the fusion in vitro of vesicles isolated from chromoplasts (Hugueney et al., 1995). If a similar mechanism exists in chloroplasts, integral membrane proteins are likely to be required to capture these vesicles. cpSecY might be needed for the integration of such proteins. It is interesting in this regard that many of the plastids lacking cpSecY contain small membrane vesicles (refer to Fig. 6 and data not shown), as would be expected if vesicle fusion were disrupted.

### **Role of cpSecY in Chloroplast Translation**

An unanticipated aspect of the *csyI* mutant phenotype is the loss of the stromal enzyme Rubisco. This is not a consequence of the chlorophyll deficiency per se, since several maize mutants with more severe chlorophyll deficiencies accumulate near normal levels of Rubisco (Harpster et al., 1984). Likewise, the Rubisco deficiency is not a consequence of photooxidative damage since it is equally severe in dark- and light-grown plastids (refer to Fig. 9). The *rbcS* mRNA accumulates to normal levels in *csyI* mutants (refer to Fig. 11) and the import of nuclear-encoded proteins into the mutant chloroplasts appears normal (refer to Fig. 10). These negative results led us to investigate the possibility that *csyI* mutants have a defect in chloroplast translation. In fact, *csyI*<sup>-</sup> plastids exhibit a dramatic decrease in the average number of ribosomes associated with the *rbcL* mRNA and with several other chloroplast mRNAs examined (refer to Fig. 12 and data not shown). The fact that the distribution of 16S rRNA in the polysome gradients is not dramatically affected in *csyI* mutants (refer to Fig. 12) sug-

gests that the mutation does not cause a global decrease in the translation initiation rate. Thus, the translation defect in *csyI* mutants differs from that in *cps* mutants, which are defective in chloroplast polysome assembly (Barkan, 1993). The small amount of pre-16S rRNA in *csyI* mutants (refer to Fig. 11) and in all *cps* mutants (Barkan, 1993) is likely the result, rather than the cause, of the translational defects (see Discussion in Barkan, 1993).

This translation defect is intriguing in that it suggests a link between chloroplast membrane biogenesis and chloroplast gene expression. One possibility is that the absence of cpSecY results in the prolonged stalling of ribosomes that are translating membrane proteins, thereby reducing the concentration of free ribosomes available for initiation. A proportion of chloroplast ribosomes is associated with thylakoid membranes (Chua et al., 1973; Margulies and Michaels, 1975; Minami and Watanabe, 1984; Breidenbach et al., 1988), and chloroplast ribosomes pause during the translation of several membrane proteins (Kim et al., 1991; Stollar et al., 1994). It is plausible that the release of some pauses is facilitated by interactions between the ribosome and a proteinaceous receptor in the thylakoid membrane. This would be analogous to the release of SRP-mediated ribosome pausing by interactions with the SRP receptor in the endoplasmic reticulum membrane (for review see Walter and Johnson, 1994). An alternative possibility arises from the recent finding that several chloroplast RNA-binding proteins are associated with internal chloroplast membranes (Zerges and Rochaix, 1998). If translational activators are among these proteins, then the absence or mislocalization of these activators in *csyI* mutants could cause defects in translation. Consistent with this notion are the observations that several translational activators in yeast mitochondria are tightly associated with the mitochondrial inner membrane (for review see Gillham et al., 1994), and that polysomes containing the *rbcL* mRNA have been found in association with thylakoid membranes (Minami and Watanabe, 1984; Breidenbach et al., 1988; Klein et al., 1988).

In summary, the data presented here provide strong evidence that cpSecY functions in more than just the cpSecA-dependent movement of proteins across the thylakoid membrane. Our results suggest further that cpSecY functions in some aspect of thylakoid membrane biogenesis other than translocating proteins to the thylakoid lumen. Finally, analysis of *csyI* mutants has revealed a link between chloroplast translation and the biogenesis of internal chloroplast membranes. A more detailed description of cpSecY function will require biochemical studies and the use of leaky and/or conditional mutations in genes encoding cpSecY.

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