

Characterisation of an *Arabidopsis* cDNA encoding a thylakoid lumen protein related to a novel ‘pentapeptide repeat’ family of proteins

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Abstract We have cloned an *Arabidopsis* cDNA encoding a novel thylakoid lumen protein, P17.4, that has been previously isolated from lumen extracts of spinach chloroplasts. The protein is synthesised with a bipartite presequence containing a Sec-type lumen-targeting signal peptide and the precursor protein is imported into the lumen of pea chloroplasts. The encoded protein is homologous to an *Anabaena* protein that is essential for correct glycolipid localisation, and is also related to at least 16 unassigned open reading frames in *Synechocystis*. This family of proteins is characterised by the presence of numerous pentapeptide repeats with the consensus structure AXLXX, and its members are predicted to be located in the cytosol, plasma membrane and periplasm/lumen. P17.4 is therefore the first higher plant member of an extended family of putative cyanobacterial proteins that may serve important roles in lipid transport or assembly.

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Key words: Chloroplast; Membrane biogenesis; Photosynthesis; Protein translocation; Thylakoid lumen

1. Introduction

The thylakoid membranes of chloroplasts and cyanobacteria carry out the light reactions of oxygenic photosynthesis, in which sunlight is used to drive the transfer of electrons from water to NADP⁺ and the resultant proton gradient is harnessed to drive ATP synthesis. These processes are carried out by four major complexes in the thylakoid membrane (photosystems I and II, the cytochrome *b_f* complex and the ATP synthase), together with a small number of ancillary carriers. The complexes themselves have been analysed in great detail and the component proteins have, for the most part, been characterised and cloned. Photosynthetic proteins account for the vast majority of chloroplast membrane protein and the thylakoid membranes of chloroplasts account for approximately 99% of the total chloroplast lipid content.

Although the photosynthetic process of thylakoid membranes have been studied in intricate detail, other functions of this highly abundant biomembrane have been slow to emerge and remarkably few non-photosynthetic thylakoid proteins have been identified. This applies particularly to the soluble lumen phase enclosed by the thylakoid membrane. Several luminal photosynthetic proteins have been well-characterised, of which plastocyanin is probably the best known, and others identified recently include the D1 processing peptidase [1] and polyphenol oxidase [2,3]. However, most luminal proteins have yet to be characterised in any real detail. In

order to gain insight into the resident proteins of this phase, and hence understand more about the functions of the thylakoid network, we recently devised a fractionation procedure which allows the preparation of lumen contents in a highly purified form [4]. At least 25 polypeptides could be identified by SDS-polyacrylamide electrophoresis, of which only six correspond to known lumen photosynthetic proteins. The majority of the remainder are of unknown identity but one has recently been further purified and shown to be a prolyl *cis/trans* isomerase [5]. N-terminal sequence data were obtained for several of the other proteins and in this report we have used these data to identify and characterise an *Arabidopsis* cDNA encoding one of the proteins. We show that the clone encodes a novel 17 kDa luminal protein with significant homology to a new protein family from *Synechocystis* and an *Anabaena* protein that is essential for correct localisation of glycolipids.

2. Materials and methods

An *Arabidopsis* cDNA encoding a 17.4 kDa luminal protein from spinach (P17.4) was identified by searching the expressed sequence tag (EST) database using the N-terminal 16-residue sequence of P17.4 [4]. The cDNA (clone 104N18T7) was obtained from the *Arabidopsis* Biological Resource Center at Ohio State University and both strands were fully sequenced. The EMBL accession number for the cDNA encoding P17.4 is T21992. The encoded precursor protein was synthesised *in vitro* by transcription translation and imported into intact pea chloroplasts as described [6]. Prediction of signal peptide cleavage sites was carried out using the signalP programme [7].

3. Results and discussion

3.1. *Arabidopsis* P17.4 is synthesised as a preprotein with a lumen-targeting, bipartite presequence

The DNA and predicted protein sequence of an *Arabidopsis* cDNA encoding P17.4 have been deposited in the EMBL database. The 16-residue N-terminal sequence determined for spinach P17.4 [4] is highly homologous to residues 78–93 in the *Arabidopsis* sequence, with 12 of the 16 residues identical in this region. The *Arabidopsis* protein is thus synthesised with a 77-residue presequence. As with all known cytosolically synthesised thylakoid lumen proteins, P17.4 is synthesised with a bipartite presequence in which two distinct domains can be recognised. The first domain corresponds to an ‘envelope transit’ peptide which directs translocation into the stroma by an envelope-localised protein translocase. These peptides are usually hydrophilic, overall positively charged and enriched in hydroxylated residues [8], and the first ca. 40 residues of the *Arabidopsis* precursor protein fit these criteria. The second targeting signal specifies translocation across the thylakoid membrane, and these peptides invariably contain a

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Table 1
Synechocystis homologues of P17.4

Gene	AAs	Predicted location
slr1819	331	cytoplasmic
slr0301	169	lumenal/periplasmic
slr0516	166	membrane
slr1152	331	cytoplasmic
slr0577	169	lumenal/periplasmic
slr1851	162	membrane
slr1446	320	cytoplasmic
slr0719	388	membrane
slr0183	259	membrane
slr0967	150	lumenal/periplasmic
slr1697	574	membrane
slr1011	270	cytoplasmic
slr0274	196	membrane
slr0414	286	cytoplasmic
slr1350	398	membrane
slr1519	245	putative membrane

The table lists the open reading frames within the *Synechocystis* genome that are highly homologous to the mature-size *Arabidopsis* P17.4 sequence. The hydrophobicity of each sequence was assessed using the TopPred 2 programme [15] and the encoded proteins are given putative locations in the cytoplasm or membrane according to the prediction obtained. Those proteins containing a single N-terminal hydrophobic region were further assessed for the possible possession of a cleavable signal peptide using the signalP programme [7] and three members have been given a predicted lumen/periplasm location accordingly.

tein is transported by the Sec pathway as predicted from the presequence structure.

3.2. P17.4 is homologous to an *Anabaena* protein required for glycolipid localisation, and to a large family of hypothetical *Synechocystis* proteins containing AXLXX pentapeptide repeats

A search of the databases reveals that P17.4 bears high homology to only one known protein: the product of the *hglK* gene in *Anabaena* sp strain PCC 7120 [14]. Interestingly, it was found that disruption of this gene led to a defect in the localisation of heterocyst-specific glycolipids, and the authors proposed that the *hglK* gene product may be involved in the transport or assembly of these lipids. The HglK protein is much larger than P17.4 (727 amino acids compared with 159 residues for the P17.4 mature protein) and it is the C-terminal half of the protein that is most homologous to P17.4 (Fig. 2A). A very surprising, and thus far unexplained, feature of the HglK protein is the presence of 36 degenerate AXLXX pentapeptide repeats [14].

Although HglK is the only known protein bearing a resemblance to P17.4, several open reading frames within prokaryotic genomes are highly homologous. Among these are a protein encoded by a plasmid in *Erwinia stewartii* (accession number L42525) and, most notably, a large family of 16 hypothetical proteins in *Synechocystis*. Several of these proteins are highly homologous to P17.4 and one such alignment is shown in Fig. 2B. As with the *Anabaena* HglK protein, this *Synechocystis* protein contains multiple copies of the pentapeptide repeat, as do all members of this particular family of hypothetical proteins. Close examination of the mature P17.4 sequence shows that this protein is also a member of this 'pentapeptide repeat' family, containing at least 16 repeat motifs that are shown underlined in Fig. 2B. However, the first and third amino acids of the *Arabidopsis* P17.4 repeats are more variable. Within the 16 repeats, the first position is likewise

occupied in most cases by Ala (9 of 16) but Val is found in four cases, Ser in two and Thr in one. The third position is again invariably hydrophobic but Leu is found in only four cases and Phe is in fact most commonly found (seven cases) with the other repeats containing Met (three cases), Ile (one example) and in one case, Ala. Since this entire region is also conserved in the *Synechocystis* slr1819 sequence, it can be seen that the cyanobacterial motifs also diverge in a few cases from the AXLXX motif, with Val present in some cases at the first position. However, it is striking that the Leu is so highly conserved. As yet, we have no information concerning the function of these repeat units although there is evidence that they are essential for the functioning of the *hglK* gene product [14].

A list of the other *Synechocystis* proteins is shown in Table 1, together with a description of their predicted sizes. Although some of the *Synechocystis* proteins are less homologous, all 16 hypothetical proteins are sequence-related because all display high homology to one another and each contains numerous copies of the characteristic pentapeptide repeat (not shown). We have also made an estimate of the locations of the proteins as deduced from their hydrophobicity profiles combined with an analysis of the N-terminal sequences. Several of the family are strongly predicted to be membrane-bound but others appear to be soluble cytosolic proteins and three are synthesised with N-terminal hydrophobic regions that are strongly predicted to be signal peptides. The latter proteins are thus probably transported into the periplasm or thylakoid lumen (or both) and may be functionally analogous to P17.4. The combined data therefore suggest that this family contains members that are cytosolic, membrane-bound and soluble in the periplasm/lumen. The functions of these proteins are as yet unknown but, given the importance of the *hglK* gene for glycolipid localisation, it may well be the case that at least some members are involved in membrane biogenesis. This possibility remains to be tested but we would point out that the sheer number of the proteins in *Synechocystis* argues for an important physiological function in most, if not all, of the cellular compartments.

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