RESEARCH PAPER

Two members of the Arabidopsis CLC (chloride channel) family, AtCLCe and AtCLCf, are associated with thylakoid and Golgi membranes, respectively

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Abstract

Though numerous pieces of evidence point to major physiological roles for anion channels in plants, progress in the understanding of their biological functions is limited by the small number of genes identified so far. Seven chloride channel (CLC) members could be identified in the Arabidopsis genome, amongst which AtCLCe and AtCLCf are both more closely related to bacterial CLCs than the other plant CLCs. It is shown here that AtCLCe is targeted to the thylakoid membranes in chloroplasts and, in agreement with this subcellular localization, that the clce mutants display a phenotype related to photosynthesis activity. The AtCLCf protein is localized in Golgi membranes and functionally complements the yeast gef1 mutant disrupted in the single CLC gene encoding a Golgi-associated protein.

Key words: Arabidopsis, CLC chloride channels, Golgi membranes, thylakoids.

Introduction

Anion channels play important roles in plant physiology, but the major limitation to investigating their biological functions originates from our poor knowledge of their molecular identity. Only genes belonging to the CLC (chloride channel) family are known; they were first described in tobacco (Lurin et al., 1996). In the Arabidopsis genome, seven CLC genes could be identified (AtCLCa–AtCLCg), and four of them (AtCLCa, -b, -c, and -d) have been cloned so far (Hechenberger et al., 1996; Geelen et al., 2000). The intracellular localization of these four proteins was deduced from expression studies of green fluorescent protein (GFP) fusion proteins in yeast (Hechenberger et al., 1996), but direct evidence for their subcellular localization and their transport activity in plant cells was lacking. The physiological characterization of Arabidopsis mutants suggested the involvement of AtCLCa (Geelen et al., 2000) and AtCLCf (Harada et al., 2004) in the regulation of nitrate levels in planta. Very recently, the tonoplast localization of AtCLCa and its role as a nitrate/proton antiporter in that membrane was demonstrated, in agreement with its physiological function in planta (De Angeli et al., 2006).

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It is shown here that fusion proteins of AtCLCe and AtCLCf with fluorescent proteins (GFP/DsRed2) are targeted to chloroplasts and Golgi vesicles, respectively, in both onion epidermal cells and Arabidopsis protoplasts. Furthermore, western blot analyses revealed the presence of AtCLCe in the thylakoid membranes. Functional data, a photosynthesis-related phenotype for clce mutants, and the functional complementation of the yeast gef1 mutant by AtCLCf are also reported. These results indicate the functionality of the proteins and suggest putative roles in agreement with their subcellular localization.

Materials and methods

Yeast strains, plant material, and culture conditions

All yeast strains were isogenic to the W303 (ura3-1 can1-100 leu2-3, 112trp1-1 his3-11, 15) strain. Two strains disrupted by the insertion of the His synthesis gene in the Sc:CLC gene were used in functional complementation tests, the haploid RGY86 and the diploid RGY192 strains (Gaxiola et al., 1998). The strains were kindly provided by R. Gaxiola (University of Connecticut, Storrs, CT, USA).

Experiments were performed using Arabidopsis thaliana, acces-
sions Columbia (Col) or Wassilewskija (WS). Sterilized seeds were grown in vitro on standard culture medium ABIS as described by Geelen et al. (2000).

Transient expression experiments of GFP fusion proteins were performed on yellow onion bulbs (Allium cepa) bought in the local market, or on protoplasts isolated from Arabidopsis cell suspensions prepared according to the procedure described in Thomine et al. (2003).

Yeast transformation and complementation tests

The positive control AtCLCd was cloned in the yeast expression vector pRS1024 carrying ampicillin resistance and LEU2 markers (Gaxiola et al., 1998). The cDNAs of AtCLCe and AtCLCf were cloned in pDR195, a yeast expression vector modified according to the Gateway system (Invitrogen), with ampicillin resistance and URA3 as selection marker genes. Yeast transformation was performed using the lithium acetate method (Clontech). Complementa-
tion tests were performed in two different discriminating growth conditions as described in Gaxiola et al. (1998), basically: (i) a low iron-containing medium (+ 0.6 mM ferrozine, Fluka) at pH 5.8, and (ii) minimal growth media at pH 7 supplemented or not with copper (0.1 mM CuSO4). Each construct was tested in both the haploid RGY86 and the diploid RGY192 yeast strains.

Transient expression of GFP/DsRed2 fusions in onion epidermal cells

The plasmid pSmRSGFP, expressing GFP under the control of the cauliflower mosaic virus 35S promoter, was used for transient expression in onion epidermal cells. It encoded a soluble highly fluo-
rescent variant of jellyfish GFP optimized for use in higher plants (Haseloff et al., 1997; Davis and Vierstra, 1998). AtCLCe or AtCLCf cDNAs were cloned upstream of the GFP and in-frame with GFP in pSmRSGFP using the BamHI site. Different constructs were used to perform co-expression experiments. The SKL22 sequence (Mollier et al., 2002), a peroxisomal targeting sequence, and SYTP, a threonyl tRNA synthetase pre-sequence, were fused to DsRed2 in the pOL vector (a gift of I Small, URGV, CNRS-INRA, Evry, France), and used as markers of peroxisomes and plastids/mitochondria, respectively. Biolistic bombardments were performed with a PSD-1000/He instrument (Bio-Rad). Acceleration of gold microcarriers (1.6 μm) coated with 1.25 μg of pure plasmid DNA (purified with Qiagen mini-prep kits) was used to transform onion epidermal cells. Bombardment parameters were as follows: vacuum, 28 inches Hg; distance to target, 6 cm; helium pressure, 650 psi. Onion scales were left for 12–24 h in the dark at 21 °C, and then epidermal tissues were removed and layered in water on glass slides for microscopy.

Transient expression of GFP/DsRed2 fusions in Arabidopsis protoplasts

Protoplasts were isolated from Arabidopsis cell suspensions, and GFP fusions were transiently expressed in the protoplasts by polyethylene glycol (PEG)-mediated transformation as described in Thomine et al. (2003). AtCLCf cDNA was introduced in the pOL-DsRed2 vector. Two Golgi markers, α-1.2 Man99-GFP (the first 99 amino acids of α-1,2 mannosidase I) (Saint-Jore-Dupas et al., 2006) and α-1.4 FucT:GFP (α-1,4 fucosyltransferase; unpublished), were kindly provided by V. Gomord (UMR 6037, Rouen University); they were used in co-expression experiments with AtCLCf:DsRed2 fusions.

GFP fluorescence visualization

Confocal microscopy was carried out using a confocal laser-scanning microscope (Leica, confocal system TCS SP2). GFP and YFP (yellow fluorescent protein) fluorochromes were excited by an argon laser at 488 nm and 514 nm, respectively; DsRed2 was excited by a helium–neon laser at 543 nm, and chlorophyll by a helium–neon red laser at 633 nm. Fluorescence was collected between 500 nm and 535 nm for GFP and between 570 nm and 637 nm for DsRed2. In Arabidopsis protoplasts, chlorophyll fluorescence was collected between 675 nm and 750 nm.

SDS-PAGE and western blot analyses

Chloroplast subfractions had been prepared from Arabidopsis cell suspensions (Ferro et al., 2002) and immunologically characterized (Seigneurin-Berny et al., 2006); they were kindly provided by N. Rolland (CEA/CNRS/UJF/INRA, Grenoble, France). Western blots were performed after SDS-PAGE of chloroplast subfractions (20 μg per fraction), using a purified rabbit antibody raised against AtCLCe synthetic peptides (Eurogentec) at a 1:1000 dilution for 3 h. The rat anti-CLC3 (Sigma) was used at a 1:300 dilution for 3 h.

Screening for T-DNA insertion mutants by PCR

A primary PCR screen was performed on pooled genomic DNA from 45 312 independently isolated T-DNA-transformed Wassilewskija (WS) lines (Bechtold et al., 1993; Bouchez et al., 1993). Twelve pairs of primers were used, each including an AtCLCe gene-specific primer distributed all over the gene and a T-DNA-specific primer for the left and right borders of the T-DNA. A series of PCR screenings on hyperpools (768 lines per pool), superpools (364 lines per pool), pools (48 lines per pool), and finally 48 independent lines subsequently led to the identification of the line of interest, clce-1. Searches in several mutant collections led to the identification of another allele, clce-2, in the SALK collection (SALK 010237; Columbia accession, Col). Each selected homozygous line was back-crossed using wild-type pollen. New homozygous mutant lines were then produced for their phenotypic characterization. During that work, no clef mutant was identified in the course of the PCR screening of DNA pools.

Fluorescence measurements

The fluorescence induction kinetics were measured with a home-
built set-up described in Rappaport et al. (2007). Briefly, leaves were cut and illuminated by continuous light (5300 μE m–2 s–1 light intensity) provided by electro luminescent diodes peaking at
AtCLCe and AtCL Cf proteins are closely related to prokaryotic CLC channels

AtCLCe (chromosome IV) and AtCL Cf (chromosome I) genes show different gene structures, with six and eight exons, respectively. Southern blot analyses revealed that these genes are present as single copies in the genome of two different Arabidopsis accessions, Col and WS (data not shown). AtCLCe encodes a polypeptide of 709 amino acids with a calculated molecular mass of 75.4 kDa (At4g35440); AtCL Cf potentially encodes two proteins, a short one with a 62.5 kDa calculated molecular mass (586 amino acids; At1g55620.1) and a longer one with a calculated molecular mass of 83.5 kDa (781 amino acids; At1g55620.2).

The AtCLCe and AtCL Cf proteins are highly hydrophobic as they showed up to 12 membrane-spanning domains, in agreement with the crystallographic structure of two bacterial CLCs resolved by high resolution X-ray and revealing the existence of 18 α-helices (Mindell et al., 2001; Dutzler et al., 2002). In the phylogenetic tree based on protein sequence comparisons (Fig. 1), most plant CLCs, including those of Arabidopsis, belong to a eukaryotic branch, except AtCLCe which defines a distinct subfamily with AtCL Cf and some tomato and rice CLCs, all closely related to bacterial CLC proteins. The AtCLCe protein sequence displays 41% amino acid identity with AtCL Cf, and 24–34% identity with animal and other plant CLC proteins. The large CLC family includes anion channels such as the torpedo-fish CLC0 (Miller and White, 1980) and the mammalian CLC1 (Steinmeyer et al., 1999), but also proton-coupled chloride transporters such as the bacterial CLCec1 (Accardi and Miller, 2004) and the mammalian CLC-4 and CLC-5 (Picollo and Puich, 2005). A refined analysis of structure/transport mechanism relationships pointed out the key role of two glutamate residues in the Cl−-binding region, E148 and E203, the latter being a hallmark that distinguishes antiporters from channels (Miller, 2006). In plants, demonstration of the antiporter activity of AtCL Ca as a nitrate/proton exchanger occurred very recently (De Angeli et al., 2006). The AtCL Ca sequence displays the two glutamate residues as well as AtCL Cb, -c, -d, and -g, while both AtCL Ce and AtCL Cf possess only E148, suggesting a different transport mechanism.

The AtCL Ce protein resides in chloroplasts

The ARAMEMNON database (http://aramemnon.botanik.uni-koeln.de/) did not give strong predictions for the subcellular localization of AtCL Ce either in chloroplasts (0.49) or in mitochondria (0.39). Nevertheless, a close analysis of the N-terminal region (MAATPLCAALRSPVSSRFF) indicated a global positive charge, enrichment in serine and in proline, and the presence of an alanine in the second position; altogether, these features are in favour of a plastid localization. The maximum cleavage site was predicted between the amino acids S17 and R18.

Transient expression in onion epidermal cells of a control construct containing GFP alone resulted in fluorescence throughout the cytosol and within the nucleus (Fig. 2A), as already reported by Haseloff et al. (1997). AtCL Ce: GFP expression resulted in a dotted fluorescence pattern, restricted to mobile organelles located in the cytosol with a diameter of about 3 μm (Fig. 2B), suggesting a plastid localization (leucoplasts in the case of onion epidermal cells; Carde, 1984). Co-expression of AtCL Ce:GFP with SYTP: Ds-Red, which is targeted to both mitochondria and plastids, revealed that green and red fluorescence co-localized to a high degree in plastids (Fig. 2C–E). In addition, upon transient expression of the AtCL Ce:GFP construct in protoplasts from Arabidopsis cell suspensions, a co-localization of the fusion protein (green fluorescence) and chlorophyll (red fluorescence) was observed (Fig. 2F–H), demonstrating the chloroplast targeting of AtCL Ce. To localize AtCL Ce further, western blot analyses were performed on chloroplast subfractions using both anti-CL C3, an antibody directed against the rat CLC3, and a purified IgG raised against two C-terminal regions of the AtCL Ce protein. The patterns obtained with anti-CL C3 could not be interpreted as it cross-reacted with peptides in chloroplasts, envelope, and stroma fractions (data not shown), but blots with anti-CL C3 IgGs confirmed the chloroplast localization and showed that AtCL Ce resides specifically in thylakoid membranes (Fig. 2I). Interestingly, from transcriptomics data available in Genevestigator tools (http://www.genevestigator.ethz.ch/at/), AtCL Ce expression is higher in green tissues compared with roots. In seedlings, AtCL Ce is expressed almost four times more in cotyledons than in roots, and at an intermediate level in hypocotyls (anatomy data sets). In other respects, looking for functional clues, the mutant gene-chip data sets show up-regulation of AtCL Ce in lec1 (~2-fold), a leaf developmental mutant, and in a gun1 gun5 double mutant (~3-fold) altered in plastid signalling pathways during de- etiolation. In contrast, AtCL Ce expression is down-regulated (~2-fold) in the pho3 mutant, in which carbon metabolism is affected. Altogether, these data would support a relationship between green tissue thylakoid localization of AtCL Ce and a potential function in chloroplasts. This hypothesis was further investigated searching for a photosynthetic phenotype in clce mutants.

Mutant clce plants display altered photosynthetic activity

clce-1 and clce-2 homozygous mutant plants exhibit developmental and morphological traits similar to those of
wild-type plants grown either in vitro or in the greenhouse (data not shown). Looking for a specific phenotypic trait of clce mutants which could be related to the chloroplast localization of the protein, chlorophyll fluorescence was measured to assess the photosynthetic activity in vivo.

High light intensities induced a strongly polyphasic fluorescence time-course, as previously observed (Delosme, 1967), when the photosynthetic rate, i.e. the reduction rate of QA, is faster than the QA reoxidation rate (Fig. 3). The first initial rising phase reflects essentially the reduction of QA (reviewed in Schreiber, 2002) and, for a given light intensity, its rate is determined by the photosystem II (PSII) antenna size and the photochemical properties of PSII. The half-time of this phase was similar in clce-1, clce-2, and wild-type plants, indicating that the light trapping efficiency of PSII was unaffected by the mutation. Consistent with this, the PSII quantum yields, as determined by the ratio ($F_m - F_0$)/$F_m$ (Genty et al., 1989), where $F_0$ and $F_m$ respectively, stand for the fluorescence yield when all PSIIa are photochemically active and inactive, were similar (0.82 ± 0.3, 0.80 ± 0.4 for the wild type and mutants, respectively). Interestingly, the clce-1 and clce-2 mutants displayed a marked phenotype, with the fluorescence increase component, developing in the 10 ms time range, being significantly slowed down, whereas the subsequent phase, occurring in the 100 ms time range, remained unaltered (Fig. 3). As discussed in Schansker et al. (2005), these two phases reflect the
disappearance of a quencher concomitant with the reduction of the plastoquinone pool, for the first one, and with the reduction of the soluble PSI electron acceptors, for the subsequent one. It is unlikely that AtCLCe activity has direct consequences on the plastoquinone pool per se.

It is thus proposed that the alterations in the kinetics of fluorescence changes induced by the illumination of dark-adapted clce mutant leaves originate from indirect effects, such as changes in the ionic strength or osmotic properties of the lumen resulting from an impaired anionic permeability of the thylakoid membrane. A CLC-type protein might contribute to anion channel activities previously reported on thylakoid membranes by Schönknecht et al. (1988) in the higher plant Peperomia metallica, and by

Fig. 2. The AtCLCe protein is targeted to chloroplasts and resides in thylakoid membranes. (A–H) Confocal microscopy analysis of the transient expression of various protein fusions in onion epidermal cells 24 h after bombardement (GFP fusions, A and B, co-expression of GFP and DsRed2 fusions, C–E) and in PEG-transformed Arabidopsis protoplasts (GFP fusions, F–H). (A and B) Results from the horizontal projection of different images. (C–H) Unique sections. The white bar indicates the scale in one dimension (µm). (A) Control GFP; (B, C, and F) AtCLCe:GFP fusion; (D) SYTP:DsRed2 fusion, a chloroplast/mitochondria marker; (G) chlorophyll fluorescence; E and H show merges of C (AtCLCe:GFP)/D (SYTP:DsRed2) and F (AtCLCe:GFP)/G (chlorophyll) images, respectively. (I) Western blot analysis of four different fractions, chloroplasts (Chl), envelope (Env), thylakoid (Thy), and stroma (Str). The IgG raised against AtCLCe immunogenic peptides revealed a specific signal in thylakoid membranes, at the expected size.
The light intensity was 5300 \( \mu \text{E m}^{-2} \text{s}^{-1} \). The leaves from plants dark-adapted for 2 h were cut and immediately used for the experiments. Fluorescence changes were measured on leaves harvested from \( \text{clce-1} \) and \( \text{clce-2} \) mutants and wild-type plants, WS and Col, respectively. Wild-type genotypes present the same pattern. The traces are representative of five different experiments performed with leaves from five different plants.

Potossin and Schönknecht in the alga \( \text{Nitellopsis obtusa} \) (1995, 1996), but no direct evidence for such anion currents in \( \text{Arabidopsis} \) photosynthetic thylakoid membranes has been provided so far. Such modifications in the intra-thylakoid ionic status could modify the overall architecture of the thylakoid, and hence the reduction of the plastoquinone pool, since the formation of the grana architecture of the thylakoid, and hence the reduction of intra-thylakoid ionic status could modify the overall plastoquinone pool, since the formation of the grana stacks is known to depend on the ionic strength.

The \( \text{AtCLCf} \) protein is targeted to Golgi vesicles

\( \text{AtCLCf:GFP} \) fluorescence in onion epidermal cells was localized in numerous small (1 \( \mu \)m diameter) organelles showing a Brownian movement throughout the cytosol (Fig. 4A). The size of these structures allowed plastids to be excluded, but could correspond to several types of organelles, mitochondria, peroxisomes, or Golgi vesicles. Co-expression of \( \text{AtCLCf:GFP} \) with \( \text{SYTP:Ds-Red} \) (Fig. 4B) led to the exclusion of mitochondrial and chloroplastic locations, as green and red fluorences did not overlap. Similarly, the green fluorescence of \( \text{AtCLCf:GFP} \) did not match the distribution of the red fluorescence displayed by a fusion protein targeted to peroxisomes, \( \text{SKL22:Ds-Red} \) (Fig. 4C). Treatment of onion epidermal cells expressing \( \text{AtCLCf:GFP} \) with brefeldin-A (BFA), a potent inhibitor of endocellular traffic, induced aggregation and redistribution of the green fluorescence (data not shown), suggesting that the \( \text{AtCLCf:GFP} \) protein was associated with a BFA-sensitive compartment. This observation is in agreement with a Golgi targeting of the \( \text{AtCLCf} \) protein.

To analyse further the subcellular compartmentation of \( \text{AtCLCf} \), transient co-expression of \( \text{AtCLCf:DsRed2} \) fusions with fluorescent markers of \( \text{cis} \)- and \( \text{trans} \)-Golgi cisternae was achieved in protoplasts from \( \text{Arabidopsis} \) cell suspensions. In tobacco leaves transformed by agroinfection, the \( \alpha \)-1,2 mannosidase I is localized in both endoplasmic reticulum (ER) and \( \text{cis} \)-Golgi subcompartments (Saint-Jore-Dupas et al., 2006). In the present biological system, \( \text{Man99-GFP} \) also showed a dual compartmentation (Fig. 4D). Red fluorescence of \( \text{AtCLCf:DsRed2} \) (Fig. 4E) and green fluorescence of \( \text{Man99-GFP} \) (Fig. 4D) completely co-localized in \( \text{cis} \)-Golgi vesicles, but not in the ER (Fig. 4F), suggesting a targeting of \( \text{AtCLCf} \) to the early Golgi compartment. Co-expression of \( \text{AtCLCf} \) with a protein targeted to \( \text{trans} \)-Golgi cisternae (Saint-Jore-Dupas et al., 2006) was also performed. A partial co-localization was observed between a \( \text{trans} \)-Golgi marker, \( \text{FucT:GFP} \) (V Gomord and M-C Kiefer-Meyer, personal communication) (Fig. 4G), and \( \text{AtCLCf:DsRed2} \) (Fig. 4H, I). Similar fluorescence patterns were obtained when co-expressing \( \text{AtCLCf:GFP} \) and \( \text{ST:DsRed2} \) (the first 52 amino acids of the sialyl transferase) fusions (data not shown). These results obtained in a homologous expression system confirm that \( \text{AtCLCf} \) would reside in Golgi vesicles. A refined analysis revealed that the fusion protein is mainly targeted to the early \( \text{cis} \)-Golgi subcompartment, and to some extent to \( \text{trans} \)-Golgi cisternae. This result contrasts with a report on the chloroplast localization in spinach of a putative CLC channel sharing sequence similarity with \( \text{AtCLCf} \) (Teardo et al., 2005). In this study, biochemical and mass spectrometry analyses relied on the use of a combination of heterologous tools that may lead to confusing results. The authors themselves do not exclude the hypothesis of the simultaneous presence of CLCf and/or CLCe in other subcellular compartments in spinach.

The present confocal microscopy analyses revealed that in \( \text{Arabidopsis} \), \( \text{AtCLCe} \) and \( \text{AtCLCf} \) reside in different membrane systems, i.e. thylakoid and Golgi membranes, respectively, suggesting that these proteins might play distinct biological roles.

AtCLCf functionally complements the yeast gef1 mutant

The yeast CLC protein, ScCLC, has been localized in the Golgi apparatus, most of the protein residing in its medial portion (Gaxiola et al., 1998; Schwappach et al., 1998) but also in the late- or post-Golgi vesicles (Gaxiola et al., 1999). Disruption of the ScCLC gene in yeast (gef1 mutant) led to a growth defect on iron-limited medium.
containing non-fermentable carbon sources (Greene et al., 1993). AtCLCd was first demonstrated to restore iron-limited growth of the yeast mutant gef1 (Hechenberger et al., 1996). Later on, Gaxiola et al. (1998) showed that other phenotypes of the gef1 mutant, the pH-induced phenotype and salt sensitivity, were also suppressed by complementation with AtCLCd and AtCLCc, but not by AtCLCa. The ability of AtCLCc and AtCLCe functionally to complement growth defects of the mutant strains RGY86 (haploid) and RGY192 (diploid) either on a low-iron medium or on a high pH medium was tested. Figure 5 illustrates the results obtained in the haploid strain RGY86 gef1. Similar pictures were obtained for the diploid strain (data not shown). No growth was observed in discriminating growth conditions when yeast strains were transformed with the corresponding empty vectors pDR195 and pRS1024 (data not shown). The gef1 mutant is unable to grow on non-fermentable carbon sources in the absence of high iron concentrations (Fig. 5A, B). This growth defect is suppressed in gef1 strains that express the Arabidopsis CLCf, but not AtCLCe (Fig. 5A, B). The gef1 mutant also failed to grow on the minimal media SD or SGE, buffered at pH 7 (Fig. 5D, F), though they grew well on a minimal medium YPD (Fig. 5C); this phenotype was also rescued by the expression of AtCLCd (Fig. 5D, F). The addition of copper allows growth of all genotypes (Fig. 5E–G), probably because it restores in gef1 background the high affinity iron uptake through the Fet3–Ccc2 complex (Gaxiola et al., 1998). In all discriminating growth conditions, expression of AtCLCe did not complement the two yeast mutant strains, probably as a consequence of its chloroplast localization (Fig. 5). In all cases, AtCLCf expression in the gef1 background was able to restore growth of the mutant strains although at a lower rate compared with the wild-type strain or with the positive control represented by the gef1 strain expressing AtCLCd (Fig. 5).

Recently, AtCLCd has been shown to co-localize in the trans-Golgi network with VHA-a1, a subunit of the proton-translocating V-type ATPase (Von der Fecht-Bartenbach et al., 2007). These data suggest that AtCLCd would be involved in the transport of a counter-anion for compensating acidification of the luminal pH in the trans-Golgi network required during endocytic and secretory

Fig. 4. The AtCLCf protein is targeted to the Golgi membranes. Confocal microscopy analysis of the transient expression of various protein fusions in onion epidermal cells 24 h after bombardment. (A) GFP fusions; (B, C) co-expression of GFP and DsRed2 fusions; (D–I) co-expression of GFP and DsRed2 fusions in PEG-transformed Arabidopsis protoplasts. (A) AtCLCf:GFP fusions; (B) merge of AtCLCf:GFP and co-expressed fusions of a chloroplast/mitochondria marker SYTP:DsRed2; (C) co-expression of AtCLCf:GFP and fusions of a peroxisome marker SKL22:DsRed2; (F) merge of (E) AtCLCf:DsRed2/(D) α-1,2 Man99:GFP; (I) merge of (H) (AtCLCf:DsRed2)/(G) α-1,4 FucT:GFP. The white bar indicates the scale in one dimension (µm).
processes (Dettmer et al., 2006). The present data illustrate that AtCLCf is functional in yeast cells and provide evidence for its ability to complement the pH-dependent growth phenotype of the gef1 mutant. AtCLCf might thus play a role in delivering anions to facilitate the luminal acidification of the cis-Golgi subcompartment.

Concluding remarks

Here the molecular cloning and preliminary biological characterization of AtCLCe and AtCLCf, two new Arabidopsis genes encoding putative plant CLC anion channels, are reported. In contrast to AtCLCa, these two proteins are located in membrane systems which are not easily amenable to electrophysiology techniques, and this hampers progress to demonstrate their genuine channel or transporter activity. However, the identification of the subcellular localization of the proteins in thylakoid or Golgi membranes, together with mutant analysis and yeast complementation assays, provides the first clues as to their cellular functions. Such functions appear specific to plant cells (AtCLCe and photosynthesis) or similar to those described for yeast and some of the mammalian CLCs (AtCLCf and acidification of Golgi or endosomal vesicles) (reviewed in Jentsch, 2007).

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