Functional analysis of plastid-encoded genes.
Application of reverse genetics on *Nicotiana tabacum*.

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CURRICULUM VITAE....................................................................................................................................................... 151
ABBREVIATIONS

A  absorbance
amp  ampicillin
APS  ammonium persulfat
ATP  adenosine 5’-triphosphate
BAP  6-benzylaminopurin
bp  base pairs
BSA  bovine serum albumin
cDNA  complementary DNA
chl  chlorophyll
DMSO  dimethyl sulfoxide
DNase  deoxyribonuclease I
DNA  deoxyribonucleic acid
DTT  dithiotreitol
EDTA  ethylenediamine tetraacetic acid
ELIP  early light-inducible proteins
EtBr  ethidium bromide
EtOH  ethanol
g  gram
h  hour
HEPES  N-(2-hydroxyethyl)piperazine N’-(2-ethane sulfonic acid)
IAA  iodoacetamide
IgG  immunoglobulin G
IPTG  isopropyl β-D-thiogalactoside
IR  inverted repeat
kb  kilobases
kDa  kilo Dalton
LB  Luria broth
LHC  light harvesting complex
LSC  large single-copy region
MES  2-N-morpholinoethanesulfonacid
min  minute
MOPS  4-morpholinopropansulfonacid
NAA \(\alpha\)-naphthalene acetic acid
nm nanometer
NPQ nonphotochemical quenching
NTP nucleoside triphosphate
OD optical density
OEC oxygen-evolving complex
ORF open reading frame
PAA polyacrylamide
PAGE polyacrylamide gel electrophoresis
PBS phosphate-buffered saline
PCR polymerase chain reaction
PEG polyethylene glycol
PNK polynucleotid kinase
PSI photosystem I
PSII photosystem II
Pt plastid
RC reaction center
rpm rotations per minute
RNase ribonuclease
RT room temperature
PFGE pulsed-field gel electrophoresis
PFD photon flux densities
RuBisCo ribulosebisphosphat carboxylase/oxygenase
SDS sodium dodecylsulfate
SDS-PAGE sodium dodecylsulfate polyacrylamide gel electrophoresis
sec second
SSC standard saline citrate
SSC small single-copy region
TBE Tris-borate-EDTA buffer
TCA trichloroacetic acid
TE Tris-EDTA
TEMED N,N,N’ ,N’-tetramethylethylene diamine
Tris Tris(hydroxymethyl)aminomethane
Tween 20 polyoxyethylensorbitanmonolaurat
| Amino acids | A, Ala  | Alanine       | M, Met | Methionine  |
|            | C, Cys  | Cysteine      | N, Asn | Asparagine  |
|            | D, Asp  | Aspartic acid | P, Pro | Proline     |
|            | E, Glu  | Glutamic acid | Q, Gln | Glutamine   |
|            | F, Phe  | Phenylalanine | R, Arg | Arginine    |
|            | G, Gly  | Glycine       | S, Ser | Serine      |
|            | H, His  | Histidine     | T, Thr | Threonine   |
|            | I, Ile  | Isoleucine    | V, Val | Valine      |
|            | K, Lys  | Lysine        | W, Trp | Tryptophan  |
|            | L, Leu  | Leucine       | Y, Tyr | Tyrosine    |

| Bases      | A: Adenine   |
|           | C: Cytosine  |
|           | G: Guanine   |
|           | T: Thymine   |
|           | U: Uracil    |
Plastids are essential organelles for the plant function and comprise one of the primary features that distinguish plants from other eukaryotes. In multicellular plants, all forms of plastids are derived from undifferentiated proplastids, which during cell differentiation form a variety of cell-specific organelles, including amyloplasts, chromoplasts, leucoplasts, elaioplasts and chloroplasts. Chlorophyll-containing plastids, chloroplasts, are considered to be the most complex type of plastids, containing six different compartments (outer and inner membranes with the periplasmatic space, thylakoid membranes, lumen and stroma), and performing processes vital for plant cell function. The biochemical key process of chloroplasts is photosynthesis, which supplies the cell with fixed carbon and energy in the forms of ATP and NADPH. Chloroplasts contain the entire enzymatic machinery for the process of photosynthesis, including both thylakoid membrane system and stroma. The organelle possesses further biogenetic potential, such as fatty acid or amino acid biosynthesis (Galili 1995, Ohlrogge and Browse 1995).

Thylakoid membranes consist of approximately 50% each lipid and proteins. Most of the polypeptides complement is comprised of five major protein complexes which span the lipid bilayer and are associated with both peripheral and soluble polypeptides in stroma and lumen, respectively. The protein complexes are designated photosystem II with the water oxidation system, photosystem I, antenna complexes LHCI and LHCII associated with both photosystems, cytochrome $b_{6f}$ complex, ATP synthase and NAD(P)H dehydrogenase (reviewed in Herrmann et al. 1991, Wollman et al. 1999). These assemblies catalyze the photosynthetic light reaction, i.e. they utilize light energy to drive electron transport from water to NADP$^+$ with coupled proton translocation across the thylakoid membrane that generates a proton motive force for synthesis of ATP.

**Photosystem II (PSII)** mediates electron transfer from water to plastoquinone as a result of the light-induced charge separation between the primary chlorophyll donor P680 and a pheophytin acceptor molecule. In higher plants and green algae, the PSII core is associated with several light harvesting antenna units. The core complex of PSII comprises six polypeptide subunits: D1 and D2 proteins, which bear two tyrosine donors $Y_Z$ and $Y_D$, cytochrome $b_{599}$ (PsbE and PsbF), two low molecular weight subunits PsbI and PsbW and two inner-antenna CP43 and CP47. The reaction center also contains six chlorophyll $a$ molecules,
two pheophytin a molecules, plastoquinone, non-heme iron and two β-carotenes. At its lumenal face PSII harbors the oxygen-evolving complex (OEC, also designated oxygen evolving enhancer, OEE) which contributes to catalyze water oxidation. In plants, OEC contains three peripheral subunits, O, P, and Q, also called OE33, OE23 and OE16. The OEC bears the Mn-cluster which is essential for catalytic activity. Studies of the structure of dimeric supercomplexes of photosystem II, that in the dimeric form associate light-harvesting antenna, by electron microscopy and single-particle image analysis presented arrangement of peripheral antenna proteins and the destabilization of the dimeric structure upon removal of 33 kDa protein. Removal of the extrinsic 23 and 16 kDa proteins induced changes in the position of the peripheral antenna proteins, in particular, movement of CP29 was recorded (Boekema et al. 2000). The mechanism of the PSII dimer-stabilizing effect of OEC33 proteins remains unclear, although the possibility that the 33 kDa protein partially shields the central region of the PSII core dimer was proposed (Boekema et al. 2000). Advanced x-ray structure studies of photosystem II crystals obtained from cyanobacteria (Vasil’ev et al. 2001, Zouni et al. 2001) provided insight into the protein composition and dynamics of the complex. The membrane-integral part of PSII monomer was shown to contain 36 transmembrane α-helices, of which twenty-two were assigned to D1, D2, CP43 and CP47, two further to the α and β subunits of cytochrome b_{559}, and another five α-helices were assigned to PsbH, PsbI, PsbL, PsbK and PsbX proteins. Seven additional transmembrane α-helices, one in proximity of CP47, six located close to CP43 and one close to cytochrome b_{559}, have been found in the electron density map but could not be assigned (Zouni et al. 2001). Analysis of excited-state dynamics in photosystem II from the x-ray crystal structure revealed a crucial role of CP43- and CP47-associated chlorophyll molecules in energy transfer from antenna to the reaction center (RC) complex (Vasil’ev et al. 2001). Altogether the PSII consists of more than 25 polypeptide subunit species, 200 - 250 chlorophyll molecules per P680 and a chlorophyll a/b ratio of 2 - 3 (Green and Durnford 1996, Barber et al. 1997, Hankamer et al. 1999, Wollman et al. 1999, Barber and Kühlbrandt 1999, Messinger 2000).

**Photosystem I (PSI)** is formed by a core complex with an attached light harvesting complex (LHCl). It drives the electron transfer from plastocyanin at the lumenal side to ferredoxin located in the chloroplast stroma. The electron transfer within PSI occurs via the primary electron donor P700, which is a chlorophyll a dimer, and five electron acceptors, a chlorophyll a monomer (A0), vitamin K (A1) and the three 4Fe-4S clusters, F_X, F_A, F_B (A2-A4). The integral core of PSI is formed by PsaA and PsaB and several small transmembrane
subunits. PsaG, PsaH, and PsaK proteins have been shown to be involved in the interaction between LHCl and PSI core (Jensen et al. 2000), PsaI and PsaJ subunits appear to be responsible for the organization of peripheral proteins, PsaL stabilizes the PSI trimer (Jordan et al. 2001), and the PsaM subunit may be involved in cyclic electron flow around photosystem I (reviewed by Chitnis 1996). The peripheral subunit PsaE, localized at the stromal face of photosystem I, forms a ferredoxin-docking pocket (Rousseau et al. 1993, Jordan et al. 2001) and, as discussed below, plays a role in cyclic electron flow around photosystem I (Yu et al. 1993). The docking pocket for plastocyanin is formed by luminal-face exposed PsaF (Chitnis 1996, Jordan et al. 2001). Crystal structure of cyanobacterial photosystem I and PSI-LHCI complexes established that the complex is made of up to 17 subunits and 127 cofactors comprising 96 chlorophylls, 2 phylloquinones, 3 Fe₄S₄ clusters, 22 carotenoids, 4 lipids, a putative Ca²⁺ ion and 201 water molecules. The chlorophyll a/b ratio is more than 5 (Mullet et al. 1980ab, Xu et al. 1994, Wollman et al. 1999, Jordan et al. 2001).

**Antenna proteins (LHC).** The common ancestor of the LHC proteins is believed to be a cyanobacterial high-light inducible protein with one membrane-spanning helix (Dolganov et al. 1995). After endosymbiotic chloroplast formation, the Lhc gene progenitor was transferred to the nuclear genome, where subsequent duplication and deletion events took place, eventually leading to the large Lhc gene family that exists today (Green and Kühlbrandt 1995, Herrmann 1997, Jansson 1999). The LHC-encoding genes are all of nuclear origin. The protein precursors contain an N-terminal transit peptide that is cleaved off during import into the chloroplast, after which the mature proteins are inserted into the thylakoid membrane with one to four membrane-spanning helices (Cline and Henry 1996). The CP29, CP26, CP24 apoproteins are encoded by the nuclear genes Lhcb4, Lhcb5 and Lhcb6 and after integration into thylakoid membranes form a minor antenna complex associated with the core of PSII. The LHCI antenna (genes: Lhca1-4) associate exclusively with PSI, and LHCII (Lhcb1-3) antenna mainly associate with PSII. The latter may also migrate to PSI as a mobile antenna pool capable of light-energy distribution between the two photosystems upon a mechanism regulated by phosphorylation/dephosphorylation of LHCII proteins (Allen et al. 1981, Allen 1992). Two additional groups of proteins with homology to the LHC family, PsbS (Funk et al. 1995) and early light-inducible proteins (ELIPs; Adamska 1997), have been identified in higher plants. All LHC-related proteins have been shown to bind chlorophylls and carotenoids (Peter and Thornber 1991, Kühlbrandt et al. 1994, Funk et al. 1995, Green and Kühlbrandt 1995, Jansson 1999, Xiao-Ping et al. 2000).
The **cytochrome b$_6$f** complex, as the mitochondrial cytochrome $bc_1$ complex, belongs to the $bc$-type cytochroms, which couple electron transfer and proton translocation across the membrane. The cytochrome $b$_6$f$ complex transfers electrons from plastoquinol to a soluble electron carrier located in the thylakoid lumen, plastocyanin. The electron transfer is accompanied by forming a proton gradient across the thylakoid membrane, which is used by the ATP synthase to generate ATP. The cytochrome $b$_6$f$ complex is formed by the Rieske protein (containing [2Fe-2S] cluster, PetC), the cytochromes $f$ (PetA) and $b_6$ (PetB), subunit IV (PetD), and several small subunit species: PetG, PetL, PetM, PetN including the newly described subunit V encoded by the nuclear gene petO (Staehelin and Arntzen 1983, Vener et al. 1997, Wollman et al. 1999, Hamel et al. 2000). The PetL subunit has been shown to play a role in stability of the cytochrome $b$_6$f$ dimer (Breyton et al. 1997). The existence of cytochrome $b$_6$f$ dimers was also proven by electron crystallography of Chlamydomonas complexes (Bron et al. 1999). The mechanism for the $bc$-type complexes is a “Q-cycle” proposed by Mitchell (1975) and modified by Crofts et al. (1983). According to this model, two electron paths exist within the complex, a high potential pathway composed of the [2Fe-2S] cluster of Rieske protein and the $c$-type heme of cytochrome $f$, and a low potential chain, composed of the two hemes of the $b$-type cytochrome, designated $b_L$ and $b_H$. Two quinol/quinone binding sites are located on opposite sites of the membrane: the Q$_o$ on the luminal face of the membrane, and the Q$_i$ site on the stromal side of the membrane. Binding of the plastoquinol to the Q$_o$ site comprises a two-electron transfer, one electron reducing the Rieske protein, which in turn reduces the $c$-type cytochrome, and the other electron reducing two $b$-type cytochrome hemes. These events are associated with the release of two protons into the luminal space. A second cytochrome $b$_6$f$ turnover reduces another plastocyanine molecule and places both $b_L$ and $b_H$ hemes of cytochrome $b_6$ in a reduced state. Oxidation of the $b$-type cytochrome hemes occurs through a two-step reduction of a plastoquinone at the Q$_i$ site coupled with an uptake of two protons from the stromal space. This mechanism contributes to generating the proton gradient that drives ATP synthesis. An unique feature of cytochrome $f$ is the existence of an internal chain of five water molecules, which may serve as a “proton wire” for protons to be translocated across the membrane (Martinez et al. 1996). Beside its role in photosynthetic electron transfer, the cytochrome complex is also recruited for redox sensing and signal transduction in chloroplasts (summarized in Wollman 2001).
INTRODUCTION

Binding of the plastoquinol at the Q_o site and subsequent reduction of cytochrome b_{6f} complex triggers kinase activation, which results in LHCII complex phosphorylation and its movement from PSII to PSI in the so-called state 1/state 2 transition mechanism (Gal et al. 1987, Bennett et al. 1988, Vener et al. 1997, Zito et al. 1999). The contribution of Rieske protein in this process was proposed. Oscillation of the Rieske protein between the proximal (close to Q_o) and distal (close to cytochrome f) site were observed in the Q-cycle process (Zhang et al. 1998). The switch of the Rieske protein into the proximal position was proposed to trigger kinase activity (Vener et al. 1997, Zito et al. 1999). Cytochrome b_{6f} complex through its interaction with the plastoquinone pool acts as a redox-sensor, thus playing a major role in acclimatization processes of plants.

The chloroplast ATP synthase belongs to the family of F_{1}-type ATPases that is also present in bacteria and mitochondria (Nelson 1992). It generates ATP from ADP and inorganic phosphate using energy derived from a trans-thylakoid electrochemical proton gradient. The ATP synthase contains a membrane-embedded proton channel, CF_{0}, and a catalytic sector with ATP synthase activity, CF_{1}, located at the stromal face of thylakoids. The entire ATP synthase complex comprises four transmembrane I - IV subunits of the CF_{0} sector (encoded by the genes atpF, atpG, AtpH and atpI), and five stromal subunits of CF_{1} with the stochiometry \( \alpha_{3}, \beta_{3}, \gamma_{1}, \delta_{1}, \varepsilon_{1} \) (encoded by the genes atpA, atpB, AtpC, AtpD and atpE). The supramolecular organization of the ATP synthase has recently been resolved by x-ray crystallography (Engelbrecht and Junge 1997, Stock et al. 1999, Menz et al. 2001). The catalytic sites are in the \( \beta \) subunits at the \( \alpha/\beta \) subunit interface (Abrahams et al. 1994). The ATP formation is known as the “binding change mechanism” (Boyer 1993), in which rotation of the central stalk (composed of subunits \( \gamma, \delta, \) and \( \varepsilon \)) is accompanied by conformational changes in the three \( \beta \) subunits, which cycle sequentially through structural states corresponding to low, medium, and high nucleotide affinities. These three states are probably associated with the release of the product (ATP), binding of substrates (ADP and inorganic phosphate), and ATP formation, respectively. The membrane-embedded ring formed by subunits III of CF_{0} sector has been proposed to be a part of a rotary motor, converting electrochemical energy into chemical energy stored in ATP. X-ray structure analysis revealed
a varied number of subunits III in yeast and *E. coli* depending on the carbon source, which could reflect a regulatory mechanism of ATP synthesis (Stock et al. 1999).

**Fig. 1.** Model scheme of the thylakoid membrane. The main protein complexes are indicated. Subunits marked in green are plastid-encoded, those in yellow are nuclear-encoded (adapted from Race et al. 1999).

The NAD(P)H-dehydrogenase complex (Ndh complex), homologous to the complex I of the mitochondrial respiratory chain, is formed by eleven plastid-encoded gene products (*ndhA*-*K*). Mutation studies of NAD(P)H dehydrogenase complex indicated its role in the cyclic electron flow around photosystem I in the light and chlororespiration in the dark (Yu et al. 1993, Burrows et al. 1998, Kofer et al. 1998, Shikanai et al. 1998, Casano et al. 2000, Joët et al. 2001). Except from its function in linear photosynthetic electron transfer, PSI also functions in a cyclic process, transferring electrons through the cytochrome *b*$_{6f}$ complex so that additional protons can be translocated across the membrane for the purpose of supplying additional ATP (Bendall and Manasse 1995). The cyclic electron flow has been shown to be an important mechanism by which plants can produce extra ATP when exposed to stress.
conditions, such as high light, high temperature or drought (Bendall and Manasse 1995). In
the light, the Ndh complex may mediate an electron flow from the stromal reductant
NAD(P)H produced indirectly by PSI to the plastoquinone. In the dark, Ndh complex may
function as a component of a respiratory chain, whose purpose will be to generate ATP in
plastids using the stromal reductant NAD(P)H (Burrows et al. 1998). In the chlororespiration
process, reduction of the plastoquinone pool by NAD(P)H is followed by an oxygen-
dependent oxidation, resulting in the formation of a pH gradient across the thylakoid
membranes (Bennoun 1982). Interestingly, it has been shown that ndh mutants displayed a
reduced ability of nonphotochemical quenching. The major component of non-photochemical
quenching, qE, arises from the trans-thylakoid pH gradient. In the wild-type, oxidation of
stromal reductant by the NAD(P)H dehydrogenase complex, accompanied by pumping
protons, generates a proton gradient necessary to form qE. Thus, the Ndh complex-mediated
electron transport may also be important for the down-regulation of PSII (Burrows et al. 1998).

One of the striking features of thylakoid membranes and the photosynthetic apparatus is their
flexibility to acclimate to changing environmental conditions. The adaptation processes can
differ in time ranges from seconds or minutes (short-term), to days and even longer periods
(long-term). One of the key processes of short-term adaptation is the redox-controlled
kinase/phosphatase system operating in the energy spill-over between the two photosystems, a
process also known as state 1/state 2 transitions (Allen 1992, Allen and Nillson 1997, Gal et
al. 1997). During the operation of the state transition mechanism, reduction of plastoquinone
(an electron carrier ultimately connecting both photosystems) and the cytochrome \( b_{6f} \)
complex activates a thylakoid-bound kinase which then phosphorylates the LHCII antenna,
causing a change in protein recognition that results in redistribution of energy from
photosystem II (state 1) to photosystem I (state 2). As discussed above, the cytochrome \( b_{6f} \)
complex plays a role of a redox sensor, which upon binding of a reduced plastoquinone
molecule at the Qo site changes the position of the Rieske protein from distal to proximal
(Zito et al. 1999, Vener et al. 1997, Wollman 2001). The reverse transition is initiated by a
turnover of plastoquinol at the cytochrome \( b_{6f} \) binding site, followed by a switch of the Rieske
protein conformation from proximal to distal, subsequent inactivation of the protein kinase,
and return of dephosphorylated LHCII to PSII (Figure 2). Activation of an LHCII kinase
therefore decreases the absorption of light by photosystem II and increases that by
photosystem I. This adaptation typically occurs in the range of a few seconds. Thylakoid
membrane proteins which can be phosphorylated in a redox-regulated and light-dependent way are PSII proteins, namely D1 and D2, CP43, PsbH, antenna apoprotein and the PSI protein PsaH which is required for the docking of phospho-LHCII when it acts as a light-harvesting antenna of photosystem I (Gal et al. 1997, Lunde et al. 2000, Vener et al. 2001). The existence of two independent kinase systems, one for PSII core proteins and another for LHCII proteins has been proposed (Gal et al. 1997, Rintamäki et al. 1997, Carlberg et al. 1999, Rintamäki et al. 2000, Lunde et al. 2000), although none of them has been verified until now. Recently, Rintamäki et al. (2000) have shown that regulation of the activity of LHCII kinase is likely to be mediated by the chloroplast ferredoxin-thioredoxin system in which the LHCII kinase activity is modulated not only by the redox state of plastoquinon and the cytochrome b$_{6f}$ complex, but also by redox reactions of its disulfide bonds through thioredoxin. In spite of various attempts, no distinct LHCII kinase has been identified with certainty, however, recent analyses of Snyders and Kohorn (1999) identified three related kinases in Arabidopsis, designated TAK 1 - 3, which originate in nuclear genes and could be imported into chloroplasts. They reside in thylakoids, and were able to phosphorylate LHCII in vitro.

**Fig. 2.** Control of distribution of excitation energy between PSII (P680) and PSI (P700). The redox status of the plastoquinone pool activates or inactivates an LHCII protein kinase resulting in phosphorylation or dephosphorylation of LHCII, respectively. The LHCII phosphatase is redox-independent. The plastoquinol pool is oxidized by PSI and reduced by PSII as well as by NAD(P)H dehydrogenase (ndh). Adapted from Allen 1992.
Protection against photooxidative damages of the photosynthetic apparatus. When light energy absorbed by plants becomes excessive relative to the capacity of photosynthesis, the xanthophyll cycle represents one means to protect plants from photooxidative damage. During illumination of chloroplasts at high photon flux densities the accumulation of excitation energy in the light-harvesting antenna of the photosystems favors the production of triplet excited chlorophyll molecules ($^3$Chl) that can interact with O$_2$ to generate reactive singlet oxygen ($^1$O$_2$), and subsequently a production of damaging reactive oxygen species, such as superoxide, hydrogen peroxide, or hydroxyl radicals. In the xanthophyll cycle, the xanthophyll violaxanthin is reversibly deepoxidized to zeaxanthin via the intermediate antheraxanthin under the action of the enzyme violaxanthin deepoxidase (Figure 3). Deepoxidation of violaxanthin is involved in the conversion of PSII to a state of high thermal energy dissipation and low chlorophyll fluorescence emission. The violaxanthin cycle takes place in PSII and PSI from a free pool of violaxanthin at the interface between the LHCs and a lipid environment. It is correlated with the process of nonphotochemical thermal dissipation of the absorbed photons excess (non-photochemical quenching, NPQ). The minor PSII antenna proteins, CP24, CP26, and CP29, contain protonatable amino acid residues which might sense a low lumenal pH, and trigger NPQ as an answer to the acidification of the thylakoid lumen (Walters et al. 1994, Walters et al. 1996, Pesaresi et al. 1997, Ruban et al. 1998). However, recently reported antisense inhibition of the CP26 and CP29 proteins in Arabidopsis excluded their direct role in the formation of NPQ (Andersson et al. 2001). The involvement of minor antenna proteins in the NPQ mechanism is probably due to carotenoid binding. Carotenoids are able to function in photoprotection by both quenching $^1$O$_2$* and/or by preventing its formation from $^3$Chl*. Transition to the S$_1$ excited state of carotenoids makes an energy transfer from the $^1$O$_2$* to the xanthophyll molecule possible. In principle, any xanthophyll molecule with more than 9 conjugated double bonds should exhibit an S$_1$ state level lower than singlet oxygen. The process of $^3$Chl* quenching has been proposed to act in a similar way (Demmig-Adams 1990, Horton 1996, Ramelli et al. 1999, Croce et al. 1999, Bassi et al. 1999, Verhoeven et al. 1999, Havaux and Niyogi 1999, Ruban et al. 1999, Ruban et al. 2000). As yet, the molecular mechanism of NPQ is not fully understood. However, an increasing number of mutants with impaired violaxanthin cycle and NPQ, such as the Arabidopsis violaxanthin de-epoxidation mutant (Niyogi et al. 1998) or the barley chlorina f2 mutant lacking chlorophyll b and LHCII apoproteins (Tardy and Havaux 1997), show a clear interaction between the processes of NPQ and the xanthophyll cycle. The recently isolated Arabidopsis mutant npq4-1, defective in the qE component of NPQ, was lacking PsbS, a
nuclear-encoded, intrinsic chlorophyll-binding protein of PSII. The structure of the four-helical PsbS protein was proposed to resemble that of the LHC protein ancestor. However, its primary function would be energy dissipation rather than light harvesting (Xiao-Ping et al. 2000).

**Fig. 3.** Molecular structure of xanthophyll – cycle pigments.
The genetic system of plastids. At the beginning of this century the observation of non-
Mendelian mutants (Baur 1909, Correns 1909) suggested the existence of a genetic system in
plastids. More than thirty years ago the existence of a unique DNA species in plastids was
verified (Sager and Ishida 1963). Since then, the structure of the plastid genome and
expression of its genes is a field of intensive studies. It is now undisputable that chloroplasts,
as well as mitochondria, are of prokaryotic origin and arose from free-living, oxygen-
producing cyanobacteria (chloroplast) and oxygen-consuming proteobacteria (mitochondria),
respectively, being incorporated into a proto-eukaryotic and eukaryotic cell by an
endosymbiotic process (Margulis 1970, Gray 1992, Bhattacharya and Medlin 1998, Herrmann
1997, Martin and Müller 1998). During evolution, the incorporated genomes exported their
genes to the nucleus through endosymbiotic gene transfer, which resulted in the
compartmented, integrated eukaryotic genetic system under the regulatory dominance of the
nucleus (Brennicke et al. 1993, Herrmann 1997, Martin and Herrmann 1998, Martin et al.

Chloroplast DNAs of higher plants are double-stranded, circular molecules with sizes
between 120 - 160 kbp. They consist of two copies of inverted repeats (IR_A and IR_B),
separated by a large and a small single-copy region (LSC and SSC, respectively). The genes
found in plastid chromosomes have been divided into several categories:
1. Genes for the photosynthetic apparatus:
   a. Photosystem I, II and cytochrome b6f genes;
   b. ATP synthase genes;
   c. NAD(P)H dehydrogenase genes;
   d. RuBisCo;
2. RNA genes and genes for the genetic apparatus;
3. Other genes:
   a. conserved ORFs;
   b. potential protein genes.

During the last decade, the plastid chromosomes from several photosynthetic organisms,
including Nicotiana tabacum (Shinozaki et al. 1986), Pisum sativum (Bookjans et al. 1986),
Marchantia polymorpha (Ohyama et al. 1986), Oryza sativa (Hiratsuka et al. 1989), Pinus
thunbergii (Wakasugi et al. 1994), Zea mays (Maier et al. 1995), Arabidopsis thaliana (Sato
et al. 1999), *Oenothera elata* (Hupfer et al. 2000), and *Spinacia oleracea* (Schmitz-Linneweber et al. 2001), have been sequenced and analyzed.

![Gene map of the plastid chromosome of *Nicotiana tabacum* (Shinozaki et al. 1986). Genes inside the circle are transcribed counterclockwise, those outside the circle clockwise. Asterisks indicate genes containing introns. Pseudogenes are marked ψ. *Ycf9* and *cemA* (*ycf10*) reading frames are indicated by arrowheads.](image-url)
In addition to approximately 100 identified genes, plastid genomes encode several reading frames of unknown function (named $ycf$, hypothetical chloroplast reading frame) which most probably represent functional genes, since they are often genetically conserved. The principal aim of this thesis was the identification and functional characterization of the proteins encoded by two such reading frames of the tobacco plastid genome, namely $ycf9$ (ORF62) and $ycf10$ (ORF229).

The function of plastid-encoded genes can be effectively analyzed using reverse genetics methods (Boynton et al. 1988, Svab et al. 1990). Plastid DNA transformation techniques have been developed during the past 20 years and were optimized for transformation of algae as well as for three higher plant species: tobacco (Svab et al. 1990), potato (Sidorov et al. 1999) and tomato (Ruf et al. 2001). Until now, two methods for introduction of foreign DNA into chloroplast DNA have been established: particle bombardment, first established for *Chlamydomonas* (Boynton et al. 1988), and subsequently for *Nicotiana tabacum* (Svab et al. 1990), and a polyethylene glycol (PEG)-based approach with protoplasts (Golds et al. 1993, O’Neil et al. 1993, Koop and Kofer 1995, Koop et al. 1996). With both methods, once the foreign DNA is delivered into the chloroplasts, it is becoming incorporated into plastid genome by homologous recombination. The transformation of plastid chromosomes requires a plastid-specific selectable marker gene to promote segregation of transformed and wild-type genome copies. The most popular marker is the prokaryotic $aadA$ (aminoglycoside adenyl transferase) gene conferring resistance to spectinomycin and streptomycin (Goldschmidt-Clermont 1991).

Because the chloroplast genome is present in multiple copies, the primary plastid transformant is usually heteroplasmatic, containing a mixture of transformed and wild-type genome copies. The regeneration of transformants under selective pressure allows plastome segregation and leads to cell lines that may carry only the mutated plastome copies. This state is designated as “homoplasmomic” (Allison et al. 1996, Koop et al. 1998). However, only disruption of chloroplast genes whose functions are not necessary for cell viability leads to the homoplasmomic state after a few regeneration cycles. Inactivation of an essential gene leads to a heteroplasmolic state which remains as long as the selective pressure for the disrupted gene is maintained (Drescher et al. 2000).
Estimation of the homoplastomic status of chloroplast transformants is crucial and one of the more controversial points of molecular analysis. In contrast to unicellular algae like *Chlamydomonas reinhardtii* (Rochaix 1995 and 1997) that contain only a single plastid with approximately 80 chromosome copies per cell, the higher plant plastome with its 5.000 - 10.000 chromosomes per cell (Herrmann and Possingham 1980) is an especially difficult material for homoplastomy state analyses. Analysis is even more complicated by the finding that higher plants may contain copies of promiscuous plastid DNA fragments in both, nucleus and mitochondria (Ellis 1982, Stern and Lonsdale 1982, Ayliffe and Timmis 1992) that analytically may be interpreted as residual wild-type plastid DNA copies even if all non-transformed plastid chromosome copies have been segregated and eliminated. Thus, a method for isolation of pure, uncontaminated plastid DNA is lacking. In this thesis a protocol of pulsed-field gel electrophoresis (Schwartz and Cantor 1984) was established that can fulfil this task.

The gene *ycf9* is ubiquitous among organisms performing oxygenic photosynthesis, both in prokaryotes (cyanobacteria) and eukaryotes (Cryptophyta, Euglenoids, Glaucocystophyceae, Rhodophyta, Stramenopiles and Viridiplantae). It is a small open reading frame of 62 codons present in all chloroplast genomes of photosynthetically active organisms analyzed so far, and has not been phylogenetically transferred into the nucleus, as various other chloroplast open reading frames. In tobacco, *ycf9* is located in a small gene cluster, flanked upstream by the PSII genes *psbD* and *psbC* and downstream by the *trnG* gene. To understand the function of the *ycf9* the site-directed inactivation approach was applied to tobacco plants. *Ycf9*, which is predicted to encode a 6.5 kDa protein with two putative membrane spanning segments, was an interesting case for reverse genetic analyses, because no reliable analyses of its function in plant cells had been presented at the beginning of my work. It was previously observed that *Chlamydomonas* strains with inactivated *ycf9* displayed a wild-type phenotype when grown under heterotrophic conditions (i.e. on medium containing acetate as a carbon source) (Higgs et al. 1998). Presently, several other research groups attempted to clarify the role of *ycf9* in higher plants. However, this yielded widely different conclusions not only of localization of its protein product, but also of its function for cell viability, ranging from potential lethality to neutrality. Mäenpää et al. (2000) presented results of experiments based on the *ycf9* gene inactivation in tobacco and concluded that it was an essential gene, since no homoplastomic transformants could be recovered. These authors had suggested that the product was localized in the chloroplast stromal membranes and likely to be associated with PSI. In yet another
study, also based on chloroplast gene inactivation in tobacco (Ruf et al. 2000), homoplastomic transformants were recovered, but the ycf9 product was now presented as the first member of a new family of antenna proteins.

The gene ycf10 is conserved in algae and in higher and lower plants. It consists of 229 - 231 amino acid residues forming four membrane spanning domains. The gene was shown to be cotranscribed with the petA gene in pea (Willey and Gray 1990), and to be located upstream of the petA gene in tobacco (Shinozaki et al. 1986), rice (Hiratsuka et al. 1989), and Marchantia (Ohyama et al. 1986). The putative ycf10 gene product was first designed HBP (for heme-binding protein) because of a short region of homology to the heme-binding domain of two cytochrome b polypeptides (Willey and Gray 1990). After immunolocalization in the inner envelope membrane of the pea chloroplasts (Sasaki et al. 1993), the gene was renamed cemA (chloroplast envelope membrane protein A). The cyanobacterial cotA gene, which displays high sequence identity to ycf10 from higher plants (Katoh et al. 1996a) has been shown to play a role in the light-induced proton extrusion (Katoh et al. 1996b). In Chlamydomonas reinhardtii ycf10 was disrupted using biolistic chloroplast transformation, and its role in the inorganic carbon (Ci) uptake into chloroplasts has been proposed (Rolland et al. 1997). Results presented below are based on the application of the site-directed mutagenesis approach in Nicotiana tabacum to clarify the role of the ycf10 gene product in higher plants.
2. MATERIALS AND METHODS

2.1 Materials

2.1.1 Plant material

For plant transformation experiments as well as for biochemical and biophysical analyses tobacco plants (*Nicotiana tabacum* L. cv. Petit Havana) were grown axenically under standard light conditions (100 µE m\(^{-2}\) s\(^{-1}\)) or under low (ca. 10 µE m\(^{-2}\) s\(^{-1}\)) light regimes generally at 20°C or 25°C, and 16/8 h light/dark cycles (Osram L85W/25 Universal White fluorescent lamps). For a phenotypic comparison, plants were also grown in soil under greenhouse conditions. For the NPQ analysis plants were additionally grown at 17°C under light regimes indicated in the experimental procedures.

2.1.2 Bacterial strains

\[ E. coli \] DH5\(\alpha\) Hanahan (1983), Gibco/BRL (Eggenstein)  
\[ E. coli \] XL1-Blue Stratagene (Heidelberg)  
\[ E. coli \] Top 10 Invitrogen (San Diego, USA)

2.1.3 Vectors

pGEM-Zf(-) Promega (Madison, USA)  
pThioHis Invitrogen (San Diego, USA)

2.1.4 Recombinant plasmids (cDNA insert)

pTB20 Shinozaki et al. (1986)  
pTB22 Shinozaki et al. (1986)  
pUC16S \(aadA\) Koop et al. (1996)

2.1.5 Primer list (5’ to 3’ end)

Oligonucleotides were obtained from MWG-Biotech GmbH (Ebersberg).
**Materials and Methods**

*ycf9* cloning and sequencing

- **ycf9** for: `CATATAGAATCAATGGATTCAT`
- **ycf9** rev: `CATATTCTCTTGGAACGAT`
- **M13** for: `GTAAAACGACGGCCAGT`
- **M13** rev: `CAGGAAACAGCTATGAC`

*ycf9* overexpression

- **ycf9EcoRI1**: `CGGAATTCTGACCTTTGCTTTCCA`
- **ycf9EcoRI2**: `CGGAATTCCCCGTTGTATTTGCTTCT`
- **ycf9XbaI1**: `GCTCTAGATCAAGAGATGAGAGAATTAA`
- **ycf9XbaI2**: `GCTCTAGAACGAAAAATACAACATT`

*ycf10* cloning and sequencing

- **ycf10** for: `AATAGATTCTAGATTCGATA`
- **ycf10** rev: `TCCCCGATGGTTAGAAAT`
- **ycf10** rev2: `AGATGATATTTTTCGAGAAG`

**aad4** primers

- **aadALI 59**: `TGCTGGCCGTACATTTGTACG`
- **aadARE 60**: `CACTACATTTCGTCCATCGCC`

Primers for pulsed-field gel electrophoresis isolated DNA

**Plastid DNA:**

- **ycf9** for PF: `CATATAGAATCAATGGATTCAT`
- **ycf9** rev PF: `ACAACGGGTACGCTAATCAA`
- **ycf10** for PF: `CTCAGTTAATAATGTCTGGAA`
- **ycf10** rev PF: `TGTATCTTGATTAATTGGAT`

**Mitochondrial DNA:**

- **cox** for: `CAATGGACGAGGTAGTCGTA`
- **cox** rev: `CTATTATAGTCCGAATACTCAT`
MATERIALS AND METHODS

nuclear DNA:
Hex3P       CCCTCCTATTTGATGCGCACAC
Hex9M       GGATGCATAGGGTATGCACGTCC

Primers for construction of RNA radiolabeled probes
ycf9 for2   TTTGCATTAATTGCAACTTC
ycf9 revT7  CGCGCGTAATACGACTCACTATAGGGCATATTCTC
             TTCGGAACGAT
ycf9 rev2T7  CGCGCGTAATACGACTCACTATAGGGTCTAAATCG
             TCCACACTAAC
psbC for    GGAGCAATGAACCTATTTGA
psbC revT7  CGCGCGTAATACGACTCACTATAGGGTCTAAATCG
             TCCACACTAAC
trnG for    TCCCCGACTGCACAA
trnG revT7  CGCGCGTAATACGACTCACTATAGGGATATATTA
             ATGCTATTTATG
aadA60 revT7 CGCGCGTAATACGACTCACTATAGGGCACTACAT
             TCGCTCATCGCC
ycf10 revT7 CGCGCGTAATACGACTCACTATAGGGATGATAT
             TGTCGAGAAG
ycf4 for    TGGCGATCAGAACATATATG
ycf4 revT7  CGCGCGTAATACGACTCACTATAGGGAGGAATATG
             CGACGATT
petA for    CTGTTTCTTTTTTACCAGGAG
petA revT7  CGCGCGTAATACGACTCACTATAGGGTACTCGTT
             AATGGTTGATCA

2.1.6 Antibodies

Polyclonal goat anti-rabbit IgG peroxidase-conjugate antibodies were obtained from Sigma Chemical Company (Munich). Rabbit polyclonal anti-phosphothreonine antibodies were obtained from Zymed Laboratories (San Francisco, USA).
2.1.7 Chemicals, enzymes, molecular weight markers

Chemicals
The chemicals used in this work were obtained from Biomol (Hamburg), Difco Laboratories (Detroit, USA), Merck AG (Darmstadt), Roche (Basel, Switzerland), Roth GmbH & Co. (Karlsruhe), Serva Feinbiochemica (Heidelberg), Sigma Chemical Company (Munich), and Qiagen (Hilden).

Restriction enzymes
Restriction enzymes and nucleoside triphosphates were obtained from New England Biolabs (Bad Schwalbach), MBI Fermentas (Vilnius, Lithuania), and Roche (Basel, Switzerland).

Radiolabeled nucleotides
Radiolabeled nucleotides were obtained from Amersham Pharmacia biotech (Freiburg i. Br.).

Molecular weight standards
DNA marker GIBCO/BRL (Karlsruhe)
RNA marker GIBCO/BRL (Karlsruhe)
λDNA-PFGE marker Amersham Pharmacia biotech (Freiburg i. Br.)

Protein molecular weight standards for SDS page
SDS 17S Sigma Chemical Comp. (Munich)
SDS 7 Sigma Chemical Comp. (Munich)
See Blue prestained marker Invitrogen (San Diego, USA)

2.1.8 Media for growth of plants and bacteria

Plant growth media
The B5 medium was prepared according to Gamborg et al. (1968), protoplast culture media (PCN and PIN) according to Koop and Kofer (1995). RMOP medium for shoot regeneration was prepared according to Svab and Maliga (1993).
Table 1. Composition of plant growth media (adapted from Brunner 1997).

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<th>F-PIN</th>
<th>RMOP</th>
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</tr>
<tr>
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</tr>
<tr>
<td>(NH₄)₂SO₄</td>
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<tr>
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<tr>
<td>Macro salts</td>
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<tr>
<td>EDTA-Fe(III)Na</td>
<td>40</td>
<td>40</td>
<td>40</td>
<td>40</td>
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</tr>
<tr>
<td>KJ</td>
<td>0.75</td>
<td>0.75</td>
<td>0.75</td>
<td>0.83</td>
<td></td>
</tr>
<tr>
<td>H₃BO₃</td>
<td>3</td>
<td>3</td>
<td>3</td>
<td>6.2</td>
<td></td>
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<tr>
<td>MnSO₄*H₂O</td>
<td>10</td>
<td>10</td>
<td>10</td>
<td>22.3</td>
<td></td>
</tr>
<tr>
<td>ZnSO₄*7H₂O</td>
<td>2</td>
<td>2</td>
<td>2</td>
<td>8.6</td>
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<tr>
<td>Na₂MoO*2H₂O</td>
<td>0.25</td>
<td>0.25</td>
<td>0.25</td>
<td>0.25</td>
<td></td>
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<tr>
<td>CuSO₄*5H₂O</td>
<td>0.025</td>
<td>0.025</td>
<td>0.025</td>
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<td></td>
</tr>
<tr>
<td>CoCl₂*6H₂O</td>
<td>0.025</td>
<td>0.025</td>
<td>0.025</td>
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<td></td>
</tr>
<tr>
<td>Micro salts</td>
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<tr>
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<td></td>
</tr>
<tr>
<td>Vitamins</td>
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<td></td>
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</tr>
<tr>
<td>BAP</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>NAA</td>
<td>0.1</td>
<td>0.1</td>
<td>0.1</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Polybuffer 74</td>
<td>10 ml</td>
<td>10 ml</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Others</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sucrose</td>
<td>20 000</td>
<td>130 000</td>
<td>30 000</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Glucose</td>
<td>85 000</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>pH</td>
<td>5.7 (KOH)</td>
<td>5.8 (KOH)</td>
<td>5.8 (KOH)</td>
<td>5.8 (KOH)</td>
<td></td>
</tr>
<tr>
<td>Osmolarity</td>
<td>550 (mosm/l)</td>
<td>550 (mosm/l)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Agar</td>
<td>7000</td>
<td></td>
<td></td>
<td></td>
<td>8000</td>
</tr>
</tbody>
</table>
Protoplast isolation (Dovzhenko et al. 1998)

| Enzymes               | 100 mg/ml | Macerozym R10 and Cellulase R10 (Serva, Heidelberg) |

Bacteria growth medium

| LB medium (1 l)       | 10.0 g    | peptone                        |
|                       | 5.0 g     | yeast extract                  |
|                       | 10.0 g    | NaCl                           |

2.1.9 Buffers and solutions

| TE buffer             | 10.0 mM   | Tris-HCl, pH 8.0               |
|                       | 1.0 mM    | EDTA                           |
| 10xTBE                | 1.0 M     | Tris-HCl, pH 8.0               |
|                       | 20.0 mM   | EDTA                           |
|                       | 0.5 M     | boric acid                     |
| 10xMOPS               | 0.05 M    | Na-acetate, pH 7.0             |
|                       | 0.1 M     | MOPS                           |
|                       | 0.1 M     | EDTA                           |
| 20xSSC                | 3.0 M     | NaCl                           |
|                       | 0.2 M     | trisodium citrate x 2H₂O       |
|                       | pH 7.4    |                                |
| 20xSSPE               | 0.2 M     | NaH₂PO₄                         |
|                       | 0.3 M     | NaCl                           |
|                       | 20.0 mM   | EDTA                           |
|                       | pH 7.4    |                                |
| 10xPBS                | 0.75 M    | NaCl                           |
|                       | 30.0 mM   | KCl                            |
|                       | 45.0 mM   | Na₂HPO₄                        |
|                       | 15.0 mM   | KH₂PO₄                         |
50 x Denhardt solution 10% (w/v) bovine serum albumin (BSA)
10% (w/v) polyvinylpyrrolidone 10 000
10% (w/v) Ficoll 400

2.1.10 Transfer membranes

Hybond nitrocellulose membranes for nucleic acid transfer were purchased from Amersham (Braunschweig). PVDF and nitrocellulose PROTRAN membranes for protein transfer were obtained from Schleicher & Schuell (Dassel) or PALL (Portsmouth, England).

2.2 Methods

2.2.1 Construction of plant transformation vectors

2.2.1.1 Construction of the recombinant plasmid for ycf9 inactivation

A 2425 bp BamHI-EcoRI fragment of tobacco plastid DNA containing ycf9 was excised from the plasmid pTB20 with BamHI (nucleotide position 36463) and EcoRI (position 38889), ligated into the vector pGEM-Zf(-), and cloned into E. coli DH5α. The ycf9 gene was inactivated by digestion with MunI at nucleotide position 37609, followed by filling in the recessed ends with the Klenow fragment of DNA polymerase I, and subsequent ligation with the blunt-ended, terminator-less aadA cassette. An individual clone carrying the aadA cassette in the same polarity as ycf9 was used for transformation.

2.2.1.2 Construction of the recombinant plasmid for ycf10 inactivation

A 3235 bp fragment of tobacco plastid DNA containing ycf10 was excised from the plasmid pTB22 with KpnI (nucleotide position 62075) and SalI (nucleotide position 65310), ligated into the vector pGEM-Zf(-), and cloned into E. coli DH5α. The ycf10 gene was inactivated by digestion with BbsI at nucleotide position 63571, and ligation with the blunt-ended, terminator-less aadA cassette. An individual clone carrying the aadA cassette in the same polarity as ycf10 was used for transformation.
2.2.2 Plant transformation

2.2.2.1 Seed sterilization

Tobacco seeds were sterilized by 1 min incubation in 70% (v/v) ethanol followed by 5 min incubation in 5% (w/v) Dimanine C (Bayer, Leverkusen) solution. The sterilized seeds were washed 3 x in sterile water.

2.2.2.2 Plastid transformation

The biolistic transformation technique (Boynton et al. 1988, Svab et al. 1990, Svab and Maliga 1993) was applied. 60 mg of gold particles (0.6 μm; BioRad, USA) were suspended in 220 μl sterile water and mixed with 25 μl of purified plasmid DNA (1 μg/μl), 250 μl of 2.5 M CaCl₂, 50 μl spermidin (0.1 M), followed by 2x washing of DNA-gold particles in 100% ethanol (p.a.), and final suspension of DNA-gold particles in 72 μl of 100% ethanol. The gold particles were delivered into plant cells using the Particle Gun - PDS 1000/He, Bio-Rad (USA).

Parameters used with the Particle Gun

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Helium pressure</td>
<td>1100 psi</td>
</tr>
<tr>
<td>Rupture discs</td>
<td>900 psi</td>
</tr>
<tr>
<td>Distance rupture disc/macrocarrier</td>
<td>8 - 10 mm</td>
</tr>
<tr>
<td>Distance macrocarrier/stopping screen</td>
<td>10 mm</td>
</tr>
<tr>
<td>Distance stopping plate/table</td>
<td>7 cm</td>
</tr>
<tr>
<td>Vacuum</td>
<td>0.85 bar</td>
</tr>
<tr>
<td></td>
<td>(26 – 27 inches Hg)</td>
</tr>
</tbody>
</table>

For each construct 20 leaves were shot and dissected 48 h after particle bombardment into 3 x 3 mm segments that were kept under non-selective conditions for 3 days, and then under selective conditions.
2.2.2.3 Selection and regeneration of mutants

Selection with 500 mg/liter of spectinomycin started three days after transformation. Green colonies began to appear and grow vigorously after approximately 3 weeks, while untransformed cell lines were bleaching and showed impaired growth. Resistant calluses were transferred to RMOP medium and cultivated in Petri dishes until shoot formation. For further segregation transplastomic lines were transferred into Petri dishes containing fresh medium at 3 - 4 week intervals. After several rounds of segregation, small green shoots of homoplastomic lines were transferred to 750 ml glass jars containing B5 agar with antibiotic, and grown for 4 - 6 weeks until use.

2.2.3 Isolation and fractionation of chloroplasts

2.2.3.1 Isolation of intact chloroplasts

Isolation of intact plastids was performed according to Müller and Eichacker (1999).

Homogenization Medium

<table>
<thead>
<tr>
<th>Component</th>
<th>Concentration</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.4 M sorbitol</td>
<td></td>
</tr>
<tr>
<td>50 mM HEPES-KOH</td>
<td>pH 8.0</td>
</tr>
<tr>
<td>2 mM EDTA</td>
<td>pH 7.5</td>
</tr>
</tbody>
</table>

Percoll Gradient Solution

<table>
<thead>
<tr>
<th>Component</th>
<th>Concentration</th>
</tr>
</thead>
<tbody>
<tr>
<td>45% / 85% Percoll</td>
<td>(w/v)</td>
</tr>
<tr>
<td>0.4 M sorbitol</td>
<td></td>
</tr>
<tr>
<td>50 mM HEPES-KOH</td>
<td>pH 8.0</td>
</tr>
<tr>
<td>2 mM EDTA</td>
<td>pH 7.5</td>
</tr>
</tbody>
</table>

Washing Buffer

<table>
<thead>
<tr>
<th>Component</th>
<th>Concentration</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.4 M sorbitol</td>
<td></td>
</tr>
<tr>
<td>50 mM HEPES-KOH</td>
<td>pH 8.0</td>
</tr>
</tbody>
</table>
10 - 20 g of freshly harvested leaf material was homogenized in approximately 100 ml of homogenization medium. The homogenate was filtrated through two layers of Miracloth (100 µm, Calbiochem, La Jolla, USA), and centrifuged at 4000 x g for 1 min. The sediment was carefully suspended in 1 ml of homogenization medium and loaded onto 45 - 85% Percoll gradients. The gradients were prepared by loading 10 ml of 45% and 5 ml of 85% Percoll in Corex tubes. After centrifugation of 8 min at 4000 x g in a “swing out” rotor the lower band containing intact plastids was collected using a Pasteur pipette and washed once in washing buffer (centrifugation 3 min, 1000 x g). All preparation steps were performed at 4°C, and isolated plastids were stored in darkness on ice.

### 2.2.3.2 Fractionation of chloroplasts into stroma and thylakoid membranes

**TMK Buffer**

<table>
<thead>
<tr>
<th>Component</th>
<th>Concentration</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tris-HCl</td>
<td>10 mM</td>
</tr>
<tr>
<td>MgCl₂</td>
<td>10 mM</td>
</tr>
<tr>
<td>KCl</td>
<td>20 mM</td>
</tr>
</tbody>
</table>

Freshly prepared chloroplasts were osmotically ruptured with TMK buffer in a ratio of 100 µg chlorophyll to 0.5 ml buffer. The chloroplasts were lysed for 10 min in darkness on ice and centrifuged for 3 - 5 min at 2000 x g. The supernatant (stroma fraction) was collected, the pellet (thylakoids) was washed twice by centrifugation in TMK buffer. The final pellet was resuspended in TMK buffer. Aliquots of the thylakoids were stored at −70°C.

### 2.2.3.3 Isolation of the major thylakoid protein complexes

The major thylakoid protein complexes, equivalent to 3 x 10⁸ chloroplasts were solubilized 10 min on ice by 1.5% (w/v) β-dodecylmaltoside (final concentration) as described by Müller and Eichacker (1999). The thylakoid lysate was loaded onto linear 0.1-1.0 M sucrose gradients and centrifuged for 16.5 h at 36 000 x g at 4°C (Beckman “swing out” rotor SP 40 Ti). The sucrose gradients were top-to-bottom fractionated into 35 fractions of 300 µl each using an ISCO 640 gradient fractionator (Instrumentation Specialties Company, USA). For further analyses individual gradient fractions were precipitated with 10% (w/v) trichloroacetic acid and washed with 100% acetone.
2.2.3.4 Washing of thylakoid membranes with Na$_2$CO$_3$

HS Buffer

0.1 M sucrose
10 mM HEPES-KOH, pH 8.0

Carbonate Buffer

0.1 M Na$_2$CO$_3$
0.1 M sucrose
10 mM HEPES-KOH, pH 8.0

To dissociate peripherally associated proteins from thylakoid membranes a Na$_2$CO$_3$ washing step was applied. Thylakoid samples at a concentration of 0.5 mg chlorophyll/ml were washed with HS buffer and incubated with 100 µl Na$_2$CO$_3$ for 10 min on ice. Washed thylakoids were pelleted by centrifugation for 30 min at 18 000 x g and suspended in an appropriate volume of HS buffer.

2.2.3.5 Purification and fractionation of chloroplast envelope membranes

Tricine Solution (1 l)

20 ml 0.5 M Tricine, pH 7.9
2 ml 0.5 M EDTA, pH 8.0
72 µl β-mercaptoethanol
1 ml PMSF

0.65 M sucrose (200 ml)

44.48 g sucrose
4.0 ml 0.5 M Tricine, pH 7.9
14.4 µl β-mercaptoethanol
0.2 ml PMSF
### Sucrose Gradients (each 200 ml)

<table>
<thead>
<tr>
<th></th>
<th>0.996 M</th>
<th>0.800 M</th>
<th>0.465 M</th>
</tr>
</thead>
<tbody>
<tr>
<td>sucrose</td>
<td>68.57 g</td>
<td>54.76 g</td>
<td>31.82 g</td>
</tr>
<tr>
<td>0.5 M Na₃PO₄, pH 7.9</td>
<td>4 ml</td>
<td>4 ml</td>
<td>4 ml</td>
</tr>
<tr>
<td>0.5 M EDTA, pH 8.0</td>
<td>0.4 ml</td>
<td>0.4 ml</td>
<td>0.4 ml</td>
</tr>
</tbody>
</table>

### Na₃PO₄ Buffer Stock

- 93.2 ml 0.5 M Na₂HPO₄
- 6.5 ml 0.5 M NaH₂PO₄

pH 7.9

### Na₃PO₄ Buffer (1l)

- 20 ml 0.5 M Na₃PO₄ Buffer Stock, pH 7.9
- 72 µl β-mercaptoethanol
- 1 ml PMSF

Purified intact plastids (see Section 2.2.3.1) were suspended in 10 ml of 0.65 M sucrose and ruptured by homogenization in a Dounce homogenizer on ice. The suspension of broken chloroplasts was filled up to 40 ml with Tricine solution and centrifuged for 1 h at 186 000 x g (Beckman TT45 rotor) at 4°C. The thylakoid pellet containing the envelope fraction was then suspended in Tricine solution and loaded onto sucrose gradients consisting of 7 ml 0.996 M / 10 ml 0.800 M / 8 ml 0.465 M sucrose solutions. Envelope membranes were isolated by centrifugation at 113 000 x g (Beckman SW27 rotor) for 3 h at 4°C. During this centrifugation step envelope membranes separate into three bands: outer envelope + RuBisCo (in 0.465 M sucrose), outer envelope (0.465 M / 0.800 M sucrose), inner envelope (0.800 M / 0.966 M sucrose), and thylakoid membrane pellet. Inner and outer envelope membranes were collected and washed 1:3 in Na₃PO₄ buffer by 1 h centrifugation at 113 000 x g at 4°C.

### 2.2.4 Overexpression of proteins in vitro

#### 2.2.4.1 Cloning of ycf9 for overexpression

A DNA fragment containing the ycf9 gene was amplified using the relevant primer pair with introduced EcoRI and XbaI restriction sites on their 5′ ends (listed in Section 2.1.5). For
overexpression of truncated proteins three additional PCR products were amplified using the primer pairs ycf9EcoRI1 and ycf9XbaI2, ycf9EcoRI2 and ycf9XbaI1, ycf9EcoRI1 and ycf9XbaI2. Amplified ycf9 fragments were digested with EcoRI and XbaI enzymes and cloned into the pThioHis expression vector (His-Path ThioFusion Expression system, Invitrogen, USA).

2.2.4.2 Overexpression of proteins in E. coli

An overnight culture of E. coli Top 10 cells was transformed with the overexpression plasmid and induced by 100 mM IPTG. The samples from 0, 1, 2, 3 and 4 hrs of induction were fractionated to soluble and insoluble fraction. Bacterial cells were suspended in buffer containing 20 mM Tris-HCl (pH 8.0), 2.5 mM EDTA and 5 mM imidazol and disrupted by three sonication-freeze-thaw cycles (Branson Sonifier W-250). Soluble and insoluble protein fractions were separated by a 10 min centrifugation at 10 000 x g. Additionally, remaining homogenat samples were precipitated with 10% (v/v) TCA followed by washing of proteins with 100% acetone. The samples were analyzed by SDS-PAGE and Western analyses.

2.2.5 Protein analysis

2.2.5.1 Determination of the protein concentration

The Bradford assay (Bradford, 1976) was used for the determination of protein concentrations.

**Bradford Stock Solution**

- 100 ml 100% ethanol
- 200 ml 80% phosphoric acid
- 359 mg Serva Blue G

**Bradford Working Buffer**

- 425 ml H₂O
- 15 ml 100% ethanol
- 30 ml 80% phosphoric acid
- 30 ml Bradford Stock Solution
A protein solution (maximal 100 µl) was mixed with 1 ml of Bradford Working buffer. After 10 min the absorbance of the sample was measured at 595 nm. The protein concentration in the sample was determined using a standard curve with samples of known protein concentrations (BSA from 1 to 20 µg)

### 2.2.5.2 Phosphorylation of proteins in vitro

**Phosphorylation Mixture**

<table>
<thead>
<tr>
<th>Component</th>
<th>Concentration</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tricine-NaOH</td>
<td>50 mM</td>
</tr>
<tr>
<td>sorbitol</td>
<td>100 mM</td>
</tr>
<tr>
<td>MgCl₂</td>
<td>5.0 mM</td>
</tr>
<tr>
<td>NaF</td>
<td>10 mM</td>
</tr>
<tr>
<td>[γ³²P]-ATP</td>
<td>0.5 µl/100 µl reaction volume</td>
</tr>
</tbody>
</table>

**4x LSB (Laemmli Sample Buffer)**

<table>
<thead>
<tr>
<th>Component</th>
<th>Concentration</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tris-HCl</td>
<td>0.25 M</td>
</tr>
<tr>
<td>SDS</td>
<td>8% (w/v)</td>
</tr>
<tr>
<td>glycerol</td>
<td>40% (w/v)</td>
</tr>
<tr>
<td>β-mercaptoethanol</td>
<td>20% (v/v)</td>
</tr>
<tr>
<td>Bromophenol Blue</td>
<td>0.016% (v/v)</td>
</tr>
</tbody>
</table>

Thylakoid membranes with a concentration of 1.0 - 1.5 mg chlorophyll/ml were pelleted and resuspended in 100 µl of phosphorylation mix. The phosphorylation reaction was carried out for 30 min in light at room temperature and stopped directly by addition of 30 µl of 4 x LBS. Autoradiographs for the phosphorylation analyses were evaluated using Fuji Bio Imaging plates type BASIII, a Fuji Bio Imaging analyzer, the BAS200025 software package (Fuji, Japan) and the TINA software package v2.08 beta (Raytest, Spröckhovel).
2.2.5.3 Protein gel electrophoresis

2.2.5.3.1 SDS-denaturing polyacrylamide (PAA) gels

2.2.5.3.1.1 Laemmli gel system (Laemmli 1970)

10 x Laemmli Buffer

<table>
<thead>
<tr>
<th></th>
<th>15% separating gel</th>
<th>12% separating gel</th>
<th>stacking gel</th>
</tr>
</thead>
<tbody>
<tr>
<td>30% acrylamide / 0.8% bisacrylamide</td>
<td>30 ml</td>
<td>24 ml</td>
<td>2.7 ml</td>
</tr>
<tr>
<td>3M Tris-HCl, pH 8.8</td>
<td>7.5 ml</td>
<td>7.5 ml</td>
<td>-</td>
</tr>
<tr>
<td>1M Tris-HCl, pH 6.8</td>
<td>-</td>
<td>-</td>
<td>1.25 ml</td>
</tr>
<tr>
<td>10% (w/v) SDS</td>
<td>0.6 ml</td>
<td>0.6 ml</td>
<td>0.1 ml</td>
</tr>
<tr>
<td>10% (w/v) APS</td>
<td>180 µl</td>
<td>180 µl</td>
<td>50 µl</td>
</tr>
<tr>
<td>TEMED</td>
<td>60 µl</td>
<td>60 µl</td>
<td>17 µl</td>
</tr>
<tr>
<td>H₂O</td>
<td>ad 60 ml</td>
<td>ad 60 ml</td>
<td>ad 10 ml</td>
</tr>
</tbody>
</table>

2.2.5.3.1.2 Schägger/von Jagow gel system (Schägger and von Jagow 1987)

Gel Buffer

<table>
<thead>
<tr>
<th></th>
<th>Tris-HCl, pH 8.45</th>
</tr>
</thead>
<tbody>
<tr>
<td>3 M</td>
<td>SDS</td>
</tr>
<tr>
<td>0.3% (w/v)</td>
<td></td>
</tr>
</tbody>
</table>
**MATERIALS AND METHODS**

**5 x Cathode Buffer**

<table>
<thead>
<tr>
<th>Concentration</th>
<th>Component</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.5 M</td>
<td>Tris</td>
<td></td>
</tr>
<tr>
<td>0.5 M</td>
<td>Tricine</td>
<td></td>
</tr>
<tr>
<td>0.5% (w/v)</td>
<td>SDS</td>
<td>pH 8.25</td>
</tr>
</tbody>
</table>

**5 x Anode Buffer**

<table>
<thead>
<tr>
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<th>Component</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 M</td>
<td>Tris-HCl</td>
<td>pH 8.9</td>
</tr>
</tbody>
</table>

**Sample Buffer (carbonate buffer)**

<table>
<thead>
<tr>
<th>Concentration</th>
<th>Component</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>100 mM</td>
<td>Na₂CO₃</td>
<td></td>
</tr>
<tr>
<td>10% (w/v)</td>
<td>sucrose</td>
<td></td>
</tr>
<tr>
<td>50 mM</td>
<td>DTT</td>
<td></td>
</tr>
</tbody>
</table>

**Pipetting scheme of gels**

<table>
<thead>
<tr>
<th>Component</th>
<th>10% separating gel</th>
<th>13% separating gel</th>
<th>20% separating gel</th>
<th>stacking gel</th>
</tr>
</thead>
<tbody>
<tr>
<td>49.5% acrylamide / 3% bisacrylamide</td>
<td>12.2 ml</td>
<td>15.75 ml</td>
<td>20 ml</td>
<td>2 ml</td>
</tr>
<tr>
<td>Gel Buffer</td>
<td>20 ml</td>
<td>20 ml</td>
<td>20 ml</td>
<td>6.2 ml</td>
</tr>
<tr>
<td>glycerol</td>
<td>8 g</td>
<td>8 g</td>
<td>8 g</td>
<td>-</td>
</tr>
<tr>
<td>10% (w/v) APS</td>
<td>200 µl</td>
<td>200 µl</td>
<td>200 µl</td>
<td>200 µl</td>
</tr>
<tr>
<td>TEMED</td>
<td>20 µl</td>
<td>20 µl</td>
<td>20 µl</td>
<td>20 µl</td>
</tr>
<tr>
<td>H₂O</td>
<td>ad 60 ml</td>
<td>ad 60 ml</td>
<td>ad 60 ml</td>
<td>ad 25 ml</td>
</tr>
</tbody>
</table>
2.2.5.3.2 Non-denaturing PAGE

2.2.5.3.2.1 Non-denaturing Deriphat-PAGE

Pipetting scheme of Deriphat-PAGE gels

<table>
<thead>
<tr>
<th></th>
<th>4% separating gel</th>
<th>8% separating gel</th>
<th>4% stacking gel</th>
</tr>
</thead>
<tbody>
<tr>
<td>48% acrylamide/1.5% bisacrylamide</td>
<td>1.3 ml</td>
<td>2.6 ml</td>
<td>0.6 ml</td>
</tr>
<tr>
<td>120 mM Tris, 480 mM glycine</td>
<td>1.6 ml</td>
<td>1.6 ml</td>
<td>1 ml</td>
</tr>
<tr>
<td>glycerol (50% w/v)</td>
<td>3.2 ml</td>
<td>3.2 ml</td>
<td>2 ml</td>
</tr>
<tr>
<td>APS 10% (w/v)</td>
<td>32 µl</td>
<td>32 µl</td>
<td>40 µl</td>
</tr>
<tr>
<td>TEMED</td>
<td>12 µl</td>
<td>12 µl</td>
<td>10 µl</td>
</tr>
<tr>
<td>H₂O</td>
<td>ad 16 ml</td>
<td>ad 16 ml</td>
<td>ad 10 ml</td>
</tr>
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</table>

Cathode Buffer

<table>
<thead>
<tr>
<th>Concentration</th>
<th>Buffer Component</th>
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</thead>
<tbody>
<tr>
<td>12 mM</td>
<td>Tris</td>
</tr>
<tr>
<td>96 mM</td>
<td>glycine</td>
</tr>
<tr>
<td>0.1% (w/v)</td>
<td>Deriphat-160</td>
</tr>
<tr>
<td></td>
<td>pH 8.3</td>
</tr>
</tbody>
</table>

Anode Buffer

<table>
<thead>
<tr>
<th>Concentration</th>
<th>Buffer Component</th>
</tr>
</thead>
<tbody>
<tr>
<td>12 mM</td>
<td>Tris</td>
</tr>
<tr>
<td>96 mM</td>
<td>glycine</td>
</tr>
<tr>
<td></td>
<td>pH 8.3</td>
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### Solubilization Mixture

<table>
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<th>Component</th>
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</thead>
<tbody>
<tr>
<td>1.6% (w/v)</td>
<td>α-dodecylmaltoside</td>
</tr>
<tr>
<td>12 mM</td>
<td>Tris-HCl, pH 8.6</td>
</tr>
<tr>
<td>96 mM</td>
<td>glycine</td>
</tr>
<tr>
<td>10% (w/v)</td>
<td>glycerol</td>
</tr>
</tbody>
</table>

From a preparation of thylakoid membranes with a chlorophyll concentration of 2 mg/ml, a sample corresponding to 12.5 µg chlorophyll was taken and solubilized by vortexing for 1 min with an equal volume of solubilization buffer. The sample was then centrifuged for 10 min at 10 000 x g and the supernatant was applied to the gel. The Deriphat-PAGE experiments were performed by Dr. A. Sokolenko.

#### 2.2.5.3.2.2 Non-denaturing Blue-Native PAGE (Schägger and von Jagow 1991)

##### Stacking Gel (4%)

<table>
<thead>
<tr>
<th>Concentration</th>
<th>Component</th>
</tr>
</thead>
<tbody>
<tr>
<td>4% PAA</td>
<td>[acrylamid/bisacrylamid = 30/0.8]</td>
</tr>
<tr>
<td>500 mM</td>
<td>ε-aminocapronic acid</td>
</tr>
<tr>
<td>50 mM</td>
<td>bis-Tris/HCl, pH 7.0</td>
</tr>
</tbody>
</table>

##### Separating Gel

<table>
<thead>
<tr>
<th>Concentration</th>
<th>Component</th>
</tr>
</thead>
<tbody>
<tr>
<td>6-12% PAA</td>
<td>[acrylamid/bisacrylamid = 30/0.8]</td>
</tr>
<tr>
<td>500 mM</td>
<td>ε-aminocapronic acid</td>
</tr>
<tr>
<td>50 mM</td>
<td>bis-Tris/HCl, pH 7.0</td>
</tr>
</tbody>
</table>

##### Cathode Buffer

<table>
<thead>
<tr>
<th>Concentration</th>
<th>Component</th>
</tr>
</thead>
<tbody>
<tr>
<td>50 mM</td>
<td>Tricine</td>
</tr>
<tr>
<td>15 mM</td>
<td>bis-Tris/HCl, pH 7.0</td>
</tr>
<tr>
<td>0.02% (w/v)</td>
<td>Serva Blue G</td>
</tr>
</tbody>
</table>

##### Anode Buffer

<table>
<thead>
<tr>
<th>Concentration</th>
<th>Component</th>
</tr>
</thead>
<tbody>
<tr>
<td>50 mM</td>
<td>bis-Tris/HCl, pH 7.0</td>
</tr>
</tbody>
</table>
ACA Buffer
750 mM $\epsilon$-aminocapronic acid
50 mM bis-Tris/HCl, pH 7.0
0.5 mM EDTA

Solubilization Buffer
2% (w/v) SDS
66 mM DTT
66 mM $\text{Na}_2\text{CO}_3$

The method is based on the Schägger and von Jagow (1991) system for separation of “native” protein complexes. Thylakoid membranes corresponding to $1 \times 10^8$ plastids were suspended in 60 µl of ACA buffer and solubilized in 5 µl DM (10% (w/v) in H$_2$O). Solubilized thylakoids were mixed with a buffer containing 5 µl of 5% (w/v) Serva Blue G, 750 mM $\epsilon$-aminocapronic acid. The samples were then loaded onto the gel and electrophoresed at 150 V. After entering the separation gel, electrophoresis was continued at 500 - 1000 V. When the gel front reached half the distance, cathode buffer containing Serva Blue G was replaced by the same buffer without staining reagent. All solutions were pre-cooled to 4°C; electrophoresis was performed at 10°C. The specifications refer to $20 \times 20 \times 0.075$ cm gels.

For the 2$^{nd}$ dimension electrophoresis, gel slices were cut out and incubated at room temperature in solubilization buffer for 20 min. The incubated gel slices were loaded onto denaturing PAA gels.

2.2.5.4 Staining of PAA gels

2.2.5.4.1 Coomassie Brilliant Blue staining

Staining Solution
40% ethanol
5% acetic acid
0.3% (w/v) Coomassie Brilliant Blue R 250 (Serva, Heidelberg)
Destaining Solution

20% ethanol
7% acetic acid

The gels were incubated in staining solution for 30 min under constant shaking, and destained until protein bands appeared well.

2.2.5.4.2 Silver staining

Fixation Solution

50% methanol
12 % acetic acid
0.05% 37% formaldehyde

Thiosulfate Solution

0.02% (w/v) Na₂S₂O₃

Silver Solution

0.2% (w/v) AgNO₃
5% 37% formaldehyde

Developing Solution

6% (w/v) Na₂CO₃
0.05% 37% formaldehyde
4 mg/ml Na₂S₂O₃

Stopping Solution

50% methanol
12% acetic acid

The 1.5 mm gel was incubated for at least 1 h in fixation solution, washed three times for 30 min in 50% ethanol and soaked for 1.5 min in thiosulfate solution. Then, the gel was washed three times for 30 sec with water and incubated in silver solution for 30 min in darkness with constant agitation. The gel was then washed again with water and incubated in developing
solution until the bands reached the desired intensity. The reaction was ended in stopping solution.

2.2.6 Immunological detection of proteins on membranes

2.2.6.1 Transfer of proteins onto nitrocellulose and PVDF membranes

Anode Buffer I
0.025 M Tris

Anode Buffer II
0.3 M Tris

Cathode Buffer
40 mM $\varepsilon$-aminocapronic acid
0.01% (w/v) SDS

Transfer membranes and PAA gels were incubated for 10 min in anode buffer II prior to transfer. PVDF membranes were prewetted in 100% methanol. The proteins were transferred onto membranes using a semi-dry blotting system. Three sheets of Whatman paper were soaked in cathode buffer and placed onto the cathode. The gel was placed on the cathode buffer-soaked Whatman paper and covered with the transfer membrane. Gel and transfer membrane assembly was covered by two layers of Whatman paper soaked in anode buffer II and three layers of Whatman paper soaked in anode buffer I. The transfer was performed for 1.5 - 2.5 h with constant current according to the formula: $mA = \text{gel size in cm}^2 \times 0.8$

2.2.6.2 Staining of blots with Ponceau S

Ponceau S Solution
0.2% (w/v) Ponceau S
1.0% acetic acid
After blotting the membrane was incubated in Ponceau S (3-hydroxy-4-[2-sulfo-4-(sulfo-phenylazo)phenylazo]-2,7-naphtalene disulfonic acid) solution for 10 min at room temperature with agitation. The membrane was rinsed with water and the positions of the molecular marker bands were marked. The membrane was finally destained in water.

2.2.6.3 Immunological detection of proteins

2.2.6.3.1 Western analysis using horseradish peroxidase-conjugated antibodies

Blocking Buffer
1x PBS  
5% (w/v)  dry milk powder  
1% (v/v)  Tween 20

Washing Buffer
1x PBS  
1% (v/v)  Tween 20

Solution I Stock
0.25 M  luminol (in DMSO)  
0.09 M  p-coumaric acid (in DMSO)

Solution I
2.5 mM  luminol  
0.4 mM  p-coumaric acid  
0.1 M  Tris-HCl pH 8.5

Solution 2
5.4 mM  H₂O₂  
0.1 M  Tris-HCl, pH 8.5

After electrophoretic transfer of the proteins from a PAA gel the membrane was incubated in blocking buffer for 1 h at room temperature. The antisera diluted to the desired concentration in blocking buffer were incubated with the membrane for 2 h at room temperature or
overnight at 4°C. The first antibodies were removed by washing the membrane four times for 10 min in blocking buffer. Anti-rabbit antibodies were diluted in blocking buffer and incubated with the membrane for 1 h at room temperature. The membranes were then washed four times for 10 min in washing buffer and developed in a mixture of solution 1 and 2 (1:1) by incubation for 1.5 min. They were exposed to X-ray films (Hyperfilm; Amersham Life Science, England) for 1 - 20 min.

2.2.7 Antibody generation

2.2.7.1 Preparation of probes for the rabbit immunization

Overexpressed fusion proteins pThioHis-ycf9 were prepared as described in Sections 2.2.4.1 and 2.2.4.2 and loaded onto 12% PAA gels. After separation, proteins were transferred onto nitrocellulose membrane and, after Ponceau S staining, a band corresponding to overexpressed fusion protein pThioHis-ycf9 was excised from the membrane. This membrane sector was then dissolved in DMSO and mixed with adjuvant TiterMax™ (CytRX Corporation, Atlanta, USA) at a ratio 1:1. Alternatively, a synthetized Ycf9 oligopeptide, PVVFASPDGSSNKNVVF, coupled with BSA was used for immunization of rabbits by mixing 350 µg of BSA-Ycf9 protein with 200 µl TiterMax™.

2.2.7.2 Subcutaneous injection of rabbits and antibody preparation (Harlow and Lane 1988)

The injections were started near the back of the neck. The antigen-TiterMax™ suspension was injected generally at three sites. After four weeks, the injection was repeated. Serum samples were taken in two-week intervals after the second boost. Bleeds were done from the ear vein of the rabbits and about 15 ml of blood was taken from each animal. After collection, the blood was allowed to clot for 30 - 60 min at 37°C. The clot was placed at 4°C overnight. The serum was then separated from the clot by centrifugation at 10 000 x g for 10 min at 4°C. Na-azide (0.01%) was added to the supernatant, which was aliquoted and stored at –20°C.
2.2.8 Nucleic acid analysis

2.2.8.1 Standard methods

The presented methods were described (if not indicated otherwise) by Sambrook et al. (1989):

- isolation of plasmid DNA (Birnboim and Doley 1979);
- determination of DNA and RNA concentrations;
- electrophoresis of DNA in agarose gels;
- electrophoresis of RNA in agarose/MOPS/formaldehyd gels;
- restriction of DNA;
- sequencing by the chain termination method (Sanger et al. 1977);
- dephosphorylation with alkaline phosphatase;
- “fill in”-reaction with Klenow DNA polymerase;
- ligation with T4 DNA ligase;
- preparation of competent *E. coli* cells using the CaCl₂ method;
- transformation of *E. coli*;
- polymerase chain reaction (“PCR”, White et al. 1989);
- Southern analysis of DNA;
- Northern analysis of RNA (Ausubel et al. 1987).

2.2.8.2 DNA analysis

2.2.8.2.1 Sequence searches and alignments

Amino acid sequences for predicted *ycf9* and *ycf10* ORFs were retrieved from the fourth iteration of a ψ-blast search (Altschul et al. 1997) run on the server of the National Centre for Biotechnology Information (http://www.ncbi.nlm.nih.gov/blast/psiblast.cgi) with the default parameters. Sequence alignments were performed using the CLUSTAL X multiple sequence alignment program (Jeanmougin et al. 1998). Amino acid sequence and protein bank accession numbers are obtained from the following references: *Arabidopsis thaliana* (GI:7525030) (Sato et al. 1999), *Chlamydomonas reinhardtii* (GI:3123110) (Higgs et al. 1998), *Marchantia polymorpha* (GI:140316) (Ohyama et al. 1986), *Nicotiana tabacum* (GI:140318) (Shinozaki et al. 1986), *Oenothera elata* (GI:6723731) (Hupfer et al. 2000),

2.2.8.2.2 Isolation of total DNA and plastid DNA

Isolation of DNA from plants was performed according to Doyle and Doyle (1990).

**Extraction Buffer**

<p>| | |</p>
<table>
<thead>
<tr>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>2% (w/v)</td>
<td>CTAB</td>
</tr>
<tr>
<td>1.4 M</td>
<td>NaCl</td>
</tr>
<tr>
<td>20 mM</td>
<td>EDTA, pH 8.0</td>
</tr>
<tr>
<td>100 mM</td>
<td>Tris-HCl, pH 8.0</td>
</tr>
<tr>
<td>100 mM</td>
<td>β-mercaptoethanol</td>
</tr>
</tbody>
</table>

For total cell DNA preparations fresh leaf tissue was used. Plastid DNA preparations started from intact plastids (see Section 2.2.3.1). Small amounts of plant material were homogenized with extraction buffer (100 µl for a leaf piece, up to 500 µl for plastid suspensions) and heated for 30 min at 60°C. Proteins were removed from samples by two chloroform/isoamylalcohol (24:1) washing steps, and DNA was precipitated with 2/3 assay volume of 2-propanol. The DNA was pelleted by centrifugation for 20 min at 10 000 x g, washed with 70% ethanol, and dissolved in TE buffer or sterile H$_2$O.

2.2.8.2.3 Southern analysis of DNA

For a Southern analysis 5 µg of plastid DNA was restricted with the desired enzyme and electrophoretically separated in an agarose gel. DNA transfer from the gel to nitrocellulose membranes was performed by capillary blotting as described in Sambrook et al. (1989). DNA was fixed to membranes by UV crosslinking (2 x Autocrosslink on “UV-Stratalinker™ 2400”, Stratagene).
2.2.8.2.4 Radioactive probe labeling

2.2.8.2.4.1 Radioactive labeling of PCR products

PCR products (100 ng DNA) were radio-labeled by the random priming method with Klenow enzyme and \(^{32}\)P-dATP or \(^{32}\)P-dCTP (according to Sambrook et al. 1989). For this assay, the Random Primed DNA Labeling Kit from Roche (Basel, Switzerland) was used according to the manufacturer’s protocol.

2.2.8.2.4.2 Radioactive labeling of oligodesoxynucleotides

Reaction Mixture (25 µl)

- 0.5 µg oligodesoxynucleotides
- 2.5 µl 10 x PNK buffer
- 7 µl \([\gamma-^{32}\text{P}]-\text{ATP}\)
- 1 µl T4 polynucleotidkinase

The 5’ hydroxy group of synthetic oligonucleotides (primers) was radioactively labeled by 30 minutes incubation at 37°C with T4 polynucleotidkinase (New England Biolabs, Bad Schwalbach). Residual radioactive mononucleotides were removed with 100% ethanol.

2.2.8.2.5 Hybridization procedure

Hybridization Buffer (0.2 l)

- 10 ml 100 x Denhardt solution
- 50 ml 20 x SSPE
- 2 ml 20% (w/v) SDS
- 4 ml herring sperm DNA (10 mg/ml)

The membrane was prehybridized for at least 2 h at 65°C in the hybridization buffer. Hybridization was performed by incubation of the membrane with the denatured radiolabeled probe in 10 ml of hybridization buffer overnight at 65°C.
2.2.8.2.6 Isolation of DNA from agarose gels

The GenElute™ Agarose Spin Columns (Supelco, Bellefonte, USA) were used to isolate DNA from agarose gels. The agarose gel slice containing the DNA was placed in the TE buffer washed GenElute Agarose spin column and centrifuged for 10 min at 10,000 x g at room temperature. DNA was precipitated by 3 M sodium acetate : 100% ethanol (0.1 : 2, v/v), and washed with 70% ethanol.

2.2.8.3 RNA analysis

2.2.8.3.1 Isolation of total RNA from plants

Total RNA from plants was isolated using TRIzol Reagent (GIBCO BRL, Karlsruhe) according to the manufacturer’s protocol. A small amount of fresh tissue (100 - 200 mg) was frozen in liquid nitrogen and homogenized with TRIzol (1 ml reagent per 100 mg material). The homogenized sample was incubated for 5 min at room temperature, mixed with chloroform (0.2 ml chloroform per 1 ml of TRIzol reagent) and centrifuged for 15 min at 12,000 x g at 4°C. RNA was precipitated from the aqueous phase with 100% 2-propanol. The RNA pellet was washed once with 70% ethanol, dried and dissolved in RNAse-free water. For Northern analyses 5 µg of total plant RNA was used.

2.2.8.3.2 Construction of radiolabeled RNA probes

To obtain a probe for radioactive in vitro transcription, PCR with a single primer containing the T7 promoter sequence was performed. This resulted in an antisense transcript for the desired RNA. Approximately 100 ng of this PCR product was used for radiolabeling.

Radioactive in vitro Transcription Mixture (20 µl end volume)

2.0 µl 10 x transcription buffer (NEB)
2.0 µl 100 mM DTT
0.5 µl RNasin (RNase inhibitor, Promega)
4.0 µl rNTP mix-UTP (2.5 mM per base)
2.4 µl 100 µM UTP
x µl PCR product
5.0 µl  $[\alpha^{32}\text{P}]-\text{UTP}$
1.0 µl  T7 RNA polymerase (20U/µl)

The transcription reaction was carried for 2 h at 37°C. Subsequently, 1 µl of DNase I was added to the mix and the sample was incubated for 15 min at 37°C. After incubation, 80 µl of sterile water was added and two steps of chloroform purification were performed. RNA was precipitated from the aqueous phase by 2.5 volumes of 100% ethanol and 0.1 volume of 3M Na-acetate and then incubated for at least 30 min at –20°C. The RNA pellet was washed once with 70% ethanol, dried and dissolved in RNase-free water. Before hybridization the radioactive RNA sample was denatured for 5 min at 65°C. For the hybridization procedure see Section 2.2.8.2.5. Autoradiographs both for RNA and DNA analyses were evaluated using Fuji Bio Imaging plates type BASIII, a Fuji Bio Imaging analyzer, the BAS200025 software package (Fuji) and the TINA software package v2.08 beta (Raytest, Spröckhovel).

2.2.8.4 Pulsed-field gel electrophoresis of DNA

2.2.8.4.1 Sample preparation

**LMP Buffer**
- 125 mM EDTA
- 330 mM sorbitol
- 25 mM Na-Citrate/HCl, pH 7.0
- 90 mM β-mercaptoethanol

**Buffer A**
- 450 mM EDTA, pH 8.0
- 10 mM Tris-HCl, pH 8.0

**Buffer A + detergent**
- 450 mM EDTA, pH 8.0
- 10 mM Tris-HCl, pH 8.0
- 1% (w/v) SLS
Buffer B
500 mM EDTA, pH 9.0

For a pulsed-field gel, samples of intact plastids were prepared as described in Section 2.2.3.1. The chloroplast suspension was adjusted to a chlorophyll concentration of 5 mg/ml with isolation medium and incubated for 10 min at 42°C. 0.9% Incert Agarose in LMP buffer was melted at 80°C and, after cooling down to 42°C, carefully mixed with the chloroplast suspension (3 volumes of agarose for 1 volume of chloroplasts). The chloroplasts/agarose mixture was loaded into block forms and left for solidification at 4°C in the dark. Solidified blocks were washed for 15 min at room temperature in 1.5 ml/block buffer A with SLS, and incubated overnight in 1.5 ml buffer A + SLS + 20 µg/ml RNase at 37°C with gentle agitation. Subsequently blocks were incubated for 24 h in 1.5 ml buffer A + 0.2 mg/ml Proteinase K at 50°C with gentle agitation. After 24 h the Proteinase K solution was removed and the blocks were washed 2 x for 4 h, and then overnight in 1.5 ml buffer B (50°C). The blocks were then transferred to 1.5 ml of buffer B and stored at 4°C.

2.2.8.4.2 Pulsed-field gel electrophoresis

The blocks with immobilized, treated chloroplasts were loaded onto 1% SeaKem LE agarose gels in 1 x TBE buffer, with λ-DNA PFGE as a molecular weight marker. The gel was run for 23 h at 170 mV at 12°C. Pulse periods started with 60 s and decreased to 0.1 s. After the run, gels were stained for 1 h in 150 ml TBE buffer with 7.5 µl ethidium bromide (10 mg/ml), destained in water and photographed under UV light.

2.2.8.4.3 Isolation of DNA from pulsed-field gels

The band of monomeric plastid DNA (around 150 000 bp) was excised from the gel with a scalpel blade. DNA was eluted from the agarose slice with the QIAEX®II (Qiagen, Hilden) kit for DNA extraction from agarose gels according to the manufacturer’s protocol.
2.2.9 Electron microscopy

Electron microscopy was performed by Prof. G. Wanner (Botanisches Institut der LMU, München). Small leaf pieces were fixed for 2 h in 2.5% glutardialdehyde (in 75 mM sodium cacodylate, 2 mM MgCl2, pH 7.0) at room temperature. Subsequently, the material was rinsed several times in fixative buffer and post-fixed for 1 h with 1% osmium tetroxide in fixative buffer at room temperature. After two washing steps in distilled water, the tissue pieces were stained with 1% uranyl acetate in 20% acetone for 1h. Dehydration was performed with a graded acetone series. Tissue samples were then infiltrated and embedded in Spurr's low-viscosity resin (Spurr 1969). After polymerization, ultra thin sections with thicknesses between 50 and 70 nm were cut with a diamond knife and mounted onto collodion-coated copper grids. The sections were post-stained with aqueous lead citrate (100 mM, pH 13.0). All micrographs were taken with EM 912 or EM 109 electron microscopes (LEO, Oberkochen).

2.2.10 Pigment analysis

Chlorophyll concentrations were determined according to Arnon (1949) with the formula:

\[ C \, (\mu g/ml) = 20.2 \times A_{645} + 8.02 \times A_{663}. \]

Pigment analyses were performed by Dr. R. Bassi (University of Verona, Italy). HPLC analysis was done according to Gilmore and Yamamoto (1991).

2.2.11 Non-photochemical quenching

Non-photochemical quenching (NPQ) measurements were done by Dr. R. Bassi (University of Verona, Italy). Fluorescence images were harvested (25 ms exposure) by a solid state CCD video camera (mod.4710 COHU, San Diego, USA) through a red filter. For determination of NPQ, plants were dark-adapted for 15 min. Measurements were performed as follows: plants were illuminated with relatively strong actinic light (220 µE m\(^{-2}\) s\(^{-1}\)) for 10 min followed by 12 min dark recovery. Actinic light was interrupted for periods of 2 sec during which the maximum fluorescence yield (F’max) was recorded with an orange LED light of 800 µE m\(^{-2}\) s\(^{-1}\). Images for fluorescence determination were taken 0.9 sec after the onset of the detection
light. NPQ was calculated as: \((F_{\text{max}} - F'_{\text{max}})/F'_{\text{max}}\) (Li 2000) where \(F_{\text{max}}\) is the maximum fluorescence in the dark adapted state and \(F'_{\text{max}}\) is the maximum fluorescence in any light-adapted state. Duration of illumination by the detecting light was 1.8 sec during the treatment with actinic light and 0.9 sec during dark recovery.

2.2.12 Measurement of photosynthetic O\(_2\) evolution with the Clark Electrode

Oxygen evolution performed by intact tobacco chloroplasts or protoplasts was measured using a Clark-type electrode (Hansatech Instruments, Bachofen GmbH, Reutlingen). Calibration of the electrode was performed at 25°C. The zero point for O\(_2\) synthesis was determined by calculation of the difference between measurements with oxygen-free water (1 ml H\(_2\)O + a few crystals of Na\(_2\)S\(_2\)O\(_4\)), and oxygen-saturated water (1 ml). Using the constant values of the oxygen content of air-saturated water (Seidell and Linke 1965), the \(\mu\)mol amount of produced O\(_2\) was calculated for 1 ml of solution and 1 cm of recorder print. For measurements, 1 ml of suspension of intact plastids or protoplasts, equivalent to a chlorophyll concentration of approximately 0.1 mg/ml was used. Plastid (protoplast) suspensions were incubated in darkness for 5 min and then exposed to light (light source of 15 cm distance from the reaction chamber). Oxygen evolution was measured for 10 min and the desired concentration of NaHCO\(_3\) was added into the reaction chamber. Measurements were performed until the maximum of O\(_2\) production was reached.

2.2.13 Chlorophyll fluorescence analysis

2.2.13.1 77K fluorescence analysis

77K fluorescence analyses were performed by Dr. A. Sokolenko. 77K emission and excitation spectra were made in a JASCO FP-777 (Japan) luminescence spectrometer. For 77K measurements thylakoid membranes from tobacco plants were injected into silica tubes and frozen in liquid nitrogen. The excitation and emission spectra were measured with a slit width of 5 nm and excitation wavelengths at 475, 690 and 735 nm.
2.2.13.2 Chlorophyll fluorescence measurements using the PAM fluorometer

*In vivo* fluorescence of single leaves was excited and detected with a pulse amplitude modulated fluorometer (PAM 2000; Walz, Germany) as described by Meurer et al. (1996). 800 ms white light pulses of 4000 µE m\(^{-2}\) s\(^{-1}\) were used to determine the maximum fluorescence (F\(_M\)) and the ratio (F\(_{M}'\)-F\(_0\))/F\(_M\) = F\(_V\)/F\(_M\). Actinic light of 40 µE m\(^{-2}\) s\(^{-1}\) intensity was used to drive photosynthesis. In addition, fluorescence quenching parameters q\(_P\) (photochemical quenching) = (F\(_M\)-F\(_S\))/(F\(_M\)-F\(_0\)'), and q\(_N\) (non-photochemical quenching) = 1-(F\(_{M}'\)-F\(_0\'))/(F\(_M\)-F\(_0\)) (Schreiber 1986) were recorded.

2.2.13.3 Fluorescence detection using the FluorCam-Video Imaging System

*In vivo* fluorescence of single leaves was excited and detected with an imaging fluorometer FluorCam-Video Imaging System (P.S.Instruments; Brno, Czech Republic) according to the manufacturer’s protocol. The continuous actinic light of 100 µE (30% of maximal 350 µmol photons m\(^{-2}\) s\(^{-1}\)) was given for 5 s. Images were captured with a sensitivity of 30% of its maximum and an electronic shutter intensity of 7 (2 ms).
3. RESULTS

3.1 Functional analysis of the \textit{ycf9} gene product

3.1.1 Comparison of the Ycf9 sequences

\textit{Ycf9} genes from a variety of photosynthetic organisms encode a well-conserved protein (Figure 6). Sequence similarity exists through all photosynthetically active organisms, with the dominant feature being two hydrophobic stretches long enough to give rise to trans-membrane (TM) \(\alpha\)-helices (Figure 5). These putative TM helices are separated by a loop of about 15 amino acids that can be predicted to be lumenal, provided the N-terminus of the protein is retained at the stromal side of the membranes. Sequence evaluation predicts a processing site for the peptidase, VFA or ALA (bold in Figure 6), located at the end of the first putative TM helix. Processing of the 6.5 kDa polypeptide at this site would yield proteins of about 3.5 and 3.0 kDa. The \textit{Chlamydomonas} sequence shows about 60\% identity with those of vascular plants, including tobacco (Figure 6).

\textbf{Fig. 5.} Kyte and Doolittle (1982) hydrophathy profile of Ycf9. Amino acid positions are indicated at the bottom of the graph. The scan window size was 13. Hydrophobic amino acid residues, placed above the zero line, form two hydrophobic stretches, I and II, long enough to give rise to trans-membrane \(\alpha\)-helices (see also Figure 6).
RESULTS

**YCF9OE1**
MTIAFQLAVF ALIATSSLNL ISVPVVFASP EGWSSNKNVV FSGTSLWGL VFLVGILNSL IS

**YCF9OE4**
MTIAFQLAVF ALIATSSLNL ISVPVVFASP EGWSSNKNVV FSGTSLWGL VFLVGILNSL IS

**YCF9NT**
MTIAFQLAVF ALIATSSLNL ISVPVVFASP DGWSSNKNVV FSGTSLWGL VFLVGILNSL IS

**YCF9AT**
MTIAFQLAVF ALIATSSLNL ISVPVVFASP DGWSSNKNVV FSGTSLWGL VFLVGILNSL IS

**YCF9SO**
MTIAFQLAVF ALIATSSLNL ISVPVVFASP DGWSSNKNIV FSGTSLWGL VFLVGILNSL IS

**YCF9ZM**
MNIAFQLAVF ALIATSSLNL ISVPVVFASP DGWSSNKNVV FSGTSLWGL VFLVAILNSL IS

**YCF9OS**
MTIAFQLAVF ALIATSSLNL ISVPVVFASP DGWSSNKNVV FSGTSLWGL VFLVAILNSL IS

**YCF9MP**
MTIAFQLAVF ALIATSSLNL ISVPVVFASP DGWSSNKNVV FSGTSLWGL VFLVAILNSL IS

**YCF9ChR**
MTSILQVALL ALIVTFALV VGVVPVFATP NGWTDNKGAV FSGLSLWLLL VFLVGILNSF VV

Consensus *....*.*.. ***..*.... .......*.* .**...*..* ***.***... .*...****. ..

Fig. 6. Multiple sequence alignment of predicted *ycf9* gene products with accession numbers. The consensus is indicated at the bottom: 100% of identity among all sequences is marked by asterisks, identity from 60% - 80% by dots. A potential cleavage site for the luminal peptidase is marked bold, two hydrophobic regions are underlined. Abbreviations: OE1,4: *Oenothera elata* plastome 1 and 4, NT: *Nicotiana tabacum*, AT: *Arabidopsis thaliana*, SO: *Spinacia oleracea*, ZM: *Zea mays*, OS: *Oryza sativa*, PT: *Pinus thunbergii*, MP: *Marchantia polymorpha*, ChR: *Chlamydomonas reinhardtii*.

3.1.2 Construction of a recombinant plasmid for *ycf9* inactivation

A reverse genetic approach was employed to understand the function of *ycf9*. An inactivation vector for the *ycf9* gene was constructed by introducing the promoterless *aadA* cassette into a MunI site in the *ycf9* gene. Correct integration of the *aadA* gene into the pGEMycf9 vector was monitored by PCR, restriction and sequence analyses.
Fig. 7. Construction of the plastid transformation vector for disruption of the ycf9 gene in the tobacco plastid chromosome. A BamHI/EcoRI fragment of the plasmid TB20 (Shinozaki et al. 1986) containing the ycf9 gene and app. 1 kb flanking regions (necessary for homologous recombination) was cloned in the pGEM vector. Inactivation of ycf9 was performed by insertion of the aadA cassette into the gene internal MunI site. The plasmid clone including the aadA cassette inserted in the same orientation as ycf9 was used for further plant transformation (see Material and Methods for details). Transcription direction of the aadA gene is indicated by an arrowhead.

3.1.3 Plant transformation, selection and regeneration of mutants

The method of biolistic chloroplast transformation is based on the process of incorporation of foreign DNA into the plastome by homologous recombination. This method was developed for Chlamydomonas reinhardtii by Boynton et al. (1988), and modified for Nicotiana tabacum by Svab et al. (1990). Twenty leaves of Nicotiana tabacum were shot with gold particles carrying the recombinant plasmid pGEMycf9. After three days of cultivation on non-selective RMOP medium leaves were cut and transferred to selective RMOP medium (for details see Section 2.2.2.2). Resistant green colonies (calli) began to grow vigorously after approximately three weeks, while non-transformed material displayed retarded growth and turned brownish. The transformation with ycf9 yielded 4 independent mutant lines. Segregation of the mutant plastomes was performed by transferring early appearing shoot tops or green calluses onto fresh selection medium in three week intervals. For molecular analyses, the mutant lines had undergone at least five cycles of shoot regeneration from leaf explants.
3.1.4 Growth phenotypes of the Δycf9 plants

Δycf9 plants were regenerated from the green calluses or plant shoot (described in Section 2.2.2.3) using antibiotic-supplied medium, and further cultivated parallel in vitro and in the greenhouse. In the latter case, the mutant tobacco lines were able to survive in soil, and produced fertile seeds. This confirmed that their phenotype was photoautotrophic. However, the lines showed two clear non-wild-type phenotypes, depending on the growth conditions used. Under standard heterotrophic conditions (100 µE m⁻² s⁻¹, 25°C, growth medium supplemented with sucrose), Δycf9 plants developed pale green leaves of the reticulate type when compared to wild-type material (Figure 8A). This corresponded to a 2.5-fold decrease in the chlorophyll content compared with the wild-type plant. This effect was less apparent when plants where grown in soil in the greenhouse. No differences in growth rates between wild-type and mutant plants grown under heterotrophic and photoautotrophic conditions were observed at 25°C (at both standard and low light intensities). When transferred to the lower temperatures (e. g. 17°C or 20°C), Δycf9 plants displayed a drastic decrease in growth rate at low light intensities (10 µE m⁻² s⁻¹). The dwarf phenotype was progressively lost with increasing light intensities (100 µE m⁻² s⁻¹; Figure 8A). The same effect was observed when plants were grown in soil under the same conditions (10 µE m⁻² s⁻¹, 20°C; Figure 8B) which excludes that the growth medium notably influenced the phenotype.
Fig. 8. Phenotypes of Δycf9 plants. Δycf9 plants grown under different conditions are compared with tobacco wild-type. Panel A: plants from in vitro cultures, Panel B: plants grown autotrophically on soil. Growth conditions used (temperature and light intensity) are indicated on the picture. Standard light intensity corresponds to 100 µE m\(^{-2}\) s\(^{-1}\), low light to 10 µE m\(^{-2}\) s\(^{-1}\).
3.1.5 Ultrastructure of Δycf9 chloroplasts

Wild-type tobacco chloroplasts of shoot cultures grown in vitro are lens-shaped, relatively large, up to 12 µm in diameter and up to 4 µm thick. Their matrix is densely packed with ribosomes, and numerous stroma and grana thylakoids are prominent (Figure 9A). Starch grains are common, while only few, evenly distributed plastoglobules were present. The plastids of Δycf9 mutants did not vary in size or shape from those of the wild-type. Nevertheless, a remarkable difference in the proportions of the stroma and grana lamellae can be observed. In wild-type plastids stroma and grana thylakoids are numerous and uniformly distributed, while in the Δycf9 mutants no stacked grana lamellae can be observed, or only in reduced amounts. Grana and stroma thylakoids possess a distinct structure and function, the protein complexes of ATP-synthase and photosystem I are located in stroma thylakoids, while grana lamellae contain most of photosystem II. The abnormal structure of the grana thylakoids in the Δycf9 plants suggests a defect in the PSII structure (Figure 9B).

Fig. 9. Electron micrographs of wild-type and Δycf9 mutant of Nicotiana tabacum. (A) Typical lens-shaped wild-type chloroplast with numerous stroma (ST) and grana thylakoids (GT in wild-type and rGT in mutant); (B) Typical Δycf9 mutant. Grana thylakoids are strongly reduced (rGT). Mutant plastids are able to form starch grains. Large plastoglobules (marked Pg; possibly an effect of cultivation in vitro) show a uniform distribution.
3.1.6 Investigation of the homoplastomic state of mutant plastome

Precise approaches, including customary sequence, Southern and PCR analyses as well as pulsed-field electrophoresis with appropriate subcellular fractions were employed to assess the homoplastomic status of the mutants. Southern analysis allows to check for the correct integration of the transgene, but does not allow the demonstration of minimal numbers of wild-type DNA copies. The PCR approach is clearly superior in sensitivity of detection but it bears the risk of (co)amplifying promiscuous plastid sequences (Ellis 1982, Stern and Lonsdale 1982, Ayliffe and Timmis 1992) residing in nuclear and/or mitochondrial DNA. Pulsed-field electrophoresis including the separation of intact, virtually contamination-free circular plastid chromosomes, is an obvious approach to bypass that problem.

3.1.6.1 Sequence analysis

To confirm the correct integration of the construct carrying the aadA gene into the tobacco plastid chromosome, the fragment of plastid DNA coding for the ycf9 gene was sequenced in the Δycf9 transformants. For this purpose an amplification product of ycf9for and ycf9rev primers was purified by agarose gel electrophoresis and served as template for sequencing (Hupfer et al. 2000). The analysis of the nucleotide sequence proved the disruption of the ycf9 gene (Figure 10). Moreover, the aadA insertion causes a frame-shift in the ycf9 reading frame, preventing translational re-initiation of a transcript containing the C-terminal gene segment.

37525  GGATTCATGA TAAAGTAAAA TCCCTCGATG ACATATTTTA TCACAATTAA
37575  TATTTTTTTG GTGATAGAGG GATCAAATGG TATATAGTTC ATTTGTTGTT
37625  AGCTTGGGAGG ATTTAAAGCA TGACTCTTGTG TTTCCAATTA ATTCGGCGTCGC
                        GTTCAATGAG AATGGATAAG AAGCTCGTGG GATGGACGTG AGGGGCCAGG
                        GATGGCTATA TTTTGGGAGG CGAAGCTCGG GCCATATCAC TAGTTGTAGG
                        GAGGAGTCTA TGGCTCGTGA AGCCGGTTTC GCCGAAGTAT CGACTCAACT
                        ATCGAGAGGT ACGCTCGGCA TCTGAGGCTA TCTCGGACCAG CCGTTGCGGG
                        CCGTACATTG GTACGGCTCC GCAGTGAGAT GGGCCCTGAA GCCACACAGT
                        GATATTGATT TGCTGGTTAC GGTGACCGTA AGGCTTGATG AAACAACGCG
                        GCGAGCTTTG ATCAACGACC TTTGGGAAC TCTGCGTCTC CCTGGAGAGA
RESULTS

GCGAGATTCT CCGCGCTGTA GAAGTCACCA TTGTTGTGCA CGACGACATC
ATTCCGTGGC GTTATCCAGC TAAGCGCGAA CTGCAATTTG GAGAATGGCA
GCGCAATGAC ATTCTTGCAG GTATCTTCGA GCCAGCCACG ATCGACATTG
ATCTGGCTAT CTTGCTGACA AAAGCAAGAG AACATAGCGT TGCCCTTGTA
GGTCCAGCGC CCGAGGAACG TTGGATCCG GCACTGAAAAG AGGATCTATT
TGAGGCCGCTA AATGAAACCT TAACGCTATG GAACCGGCGC CCCGACTGGG
CTGGCGATGA GCGAAATGTA GTGCTTACGT TGTCCCAGAT TGCTGACAGC
GCAGTAAACC GCAAATCGC GCCGAGGAT GAAGCTGCAGC ACTGGCAAT
GGAGGCGCTG CCGCCCAGT ATCAGCCCAGT CATACTTGAA ACTAGACAGG
CTTATCTTGG ACAAGAAGAA GATCGCTTGG CCGGCGCCG AGATCAGTTG
GAAGAATTGG TCCACTACGT GAAAGGCGG AATCACTAAGG TAGTTGCAA

37604 ATAACTGCAAG CTTGGCTTTAC AATTTGCTTG TTTTGCAATAG TTTGCTACTT
37640 CATTAATCTTT ATTTGATGAGC GTACCCGGAT TCCTGATGCC
37690 TGTCATACGT AAAAAATGT TGTATTTTTCT GGTACATCCT TATGGATTGG
37740 ATTAATCTTT CTGGTGCTGGTA TTCTTAATTC TCTCATCTCT
37790 CGTCGAGAC CCAAAACCAA AATGACCCCC CTAATTTTTC TCGGTTGTGA
37840 GACACATAAA ATTGGAAATCT AAGTCCCCAG AGAAACGCAA AATCAAAAT
37890 AGAAAAACAA AAAATTAGAG GGGGGTCAAG CTTCTTGAAT AAAAAGAATA
37940 CAATATAAAA

Fig. 1. Nucleotide sequence of the ycf9 gene from transformed tobacco lines. Start and stop codons of ycf9 are indicated in bold, the inserted aadA sequence is underlined.

3.1.6.2 Southern analysis

The plant material obtained after 6 – 8 regeneration cycles on selective medium was used to check the ratio of transformed/wild-type plastome copies by Southern analysis. Isolated plastid DNA was restricted and used for RFLP analyses and subsequent hybridization with a radiolabeled ycf9 probe. Different amounts of wild-type ptDNA concentrations were loaded to
estimate wild-type plastome copies in $\Delta ycf9$ plants. In all transgenic lines no wild-type hybridization signal became visible (Figure 11).

![Fig. 11. Southern analysis of ycf9 knockout plants. 5 µg of purified tobacco plastid DNA was digested with BanII and hybridized with a radiolabeled ycf9 probe. Hybridization yields a 2.3 kb fragment for wild-type, and a 3.2 kb fragment for transformed plants. Lane 1: 100 % (5 µg) of wild-type ptDNA, lane 2: 50%, lane 3: 20%, lane 4: 10%, lane 5: 5% of wild-type ptDNA, lane 6, 7, 8: ptDNA of three independently generated transformed lines.](image)

3.1.6.3 PCR analysis

The PCR reaction was performed using a primer pair designed for the regions flanking the ycf9 gene (ycf9for and ycf9rev). This primer combination allowed to distinguish signals originating from the transformed and wild-type DNA sequences. Clear signals expected from the wild-type plastome appeared when total DNA was used as a template.

To exclude the possibility of contamination of the sample by nuclear and/or mitochondrial DNA sequences DNA isolated from plastids of isopycnic Percoll gradients was subsequently used as a template for PCR. A strong reduction of the wild-type plastome product was noted, although the signal arising from the hybridization between the transformed and wild-type amplification products was still present.
3.1.6.4 PCR analysis of DNA purified by pulsed-field gel electrophoresis

To investigate if wild-type PCR signal, detectable when even plastid $\Delta ycf9$ DNA served as a template, arises from the promiscuous DNA or residual wild-type plastid DNA molecules, the purification step including pulsed-field gel electrophoresis (PFGE) was employed. Three preparations of $\Delta ycf9$ DNA, total, Percoll gradient purified, and PFGE purified plastid DNA, were investigated for presence of wild-type DNA copies using a $ycf9$forPF and $ycf9$revPF primer pair. In contrast to total DNA and Percoll gradient-purified DNA preparation, no wild-type originating signal was detected when PFGE plastid DNA served as a template (Figure 13). The sensitivity of this approach can be further increased if the gel shown in Figure 13 is subjected to a Southern analysis with radiolabeled probe.
3.1.6.4.1 Southern analysis of $\Delta\text{ycf9}$ PFGE purified plastid DNA

Electrophoretically separated PCR products were visualized by ethidium bromide staining, subsequently blotted onto nylon membrane and hybridized with a radiolabeled, $\text{ycf9}$-specific probe (Figure 14). In contrast to the total- and Percoll gradient purified plastid DNA, no 200 bp, wild-type specific signal has been found in the PFGE purified plastid DNA sample. The detection level of this method was estimated at one wild-type gene copy to $10^4$ transformed gene copies, what corresponds to one wild-type gene copy detectable in one plant cell.

![Fig. 14. PCR analysis of $\Delta\text{ycf9}$ plants. $\text{Ycf9}$ was amplified from transformed plants using $\text{ycf9forPF}$ and $\text{ycf9revPF}$ primers. PCR products were electrophoretically separated, blotted onto nylon membrane and hybridized with a radiolabeled $\text{ycf9}$-specific oligonucleotide. The wild-type signal appears as a 200 bp band, while the transplastomic fragment is 900 bp larger (1100 bp). The intermediate band is an artefact. Different preparations of plant DNA were used to investigate the homoplastic status of the mutant. Lane 1: primer control, 2: $\Delta\text{ycf9}$ total DNA, 3: $\Delta\text{ycf9}$ Percoll gradient purified plastid DNA, 4: $\Delta\text{ycf9}$ PFGE purified plastid DNA.](image)

3.1.7 Northern analysis

In tobacco, $\text{ycf9}$ is flanked upstream by the PSII genes $\text{psbD}$ and $\text{psbC}$, and downstream by $\text{trnG}$. Insertion of the $\text{aadA}$ cassette was designed to interrupt $\text{ycf9}$ at the 6th codon without notably disturbing expression of the preceding genes ($\text{psbC}$ and $\text{psbD}$), nor of $\text{trnG}$ (Figure 15, upper panel). The RNA filter hybridization analyses performed with all relevant probes ($\text{ycf9}$, $\text{psbC}$, $\text{trnG}$, and $\text{aadA}$) was consistent with the desired genome structure. Six different transcripts were observed originating from the $\text{psbD-psbC-ycf9}$ cluster, whereas $\text{trnG}$ appeared to be transcribed independently, consistent with a previous report (Yao et al. 1989). Most of the transcripts identified with the $\text{psbC}$, $\text{ycf9}$ and $\text{aadA}$ probes in $\Delta\text{ycf9}$ plants were larger by about 0.9 kb, the size of the inserted $\text{aadA}$ cassette. The expression of the downstream $\text{trnG}$ gene was not detectably modified by this insertion. The complex pattern caused by post-transcriptional RNA processing was modified only in those RNA species that included the $\text{aadA}$ insertion. No wild-type signals were detected, even in overloaded gels.
Fig. 15. Maps of the *psbD/C-ycf9* gene cluster in the wild-type and in Δycf9 tobacco. The gene on the opposite strand, between *psbC* and *ycf9*, is *trnS*. Known promoters are indicated by bent arrows. The MunI site used for insertion of the *aadA* cassette is indicated in the wild-type restriction map (M). Below each map, horizontal lines indicate the positions of probes used for RNA filter hybridization. The deduced or known positions of transcripts are shown below the maps. The pattern in the wild-type chloroplasts is according to Yao et al. (1989), the transcripts in the Δycf9 transformant were deduced by extrapolation from the wild-type RNAs. Transcript sizes in the wild-type are: T1: 4.4 kb, T2: 3.6 kb, T3: 3.6 kb, T4: 2.6 kb, T5: 2.6 kb, T6: 1.7 kb, T8: 0.71 kb, T9 (*trnS*): 0.7 kb. In the transformant the sizes of transcripts modified by the *aadA* insertion (indicated by asterisks) are: T1*: 5.3 kb, T3*: 4.5 kb, T5*: 3.5 kb, and 1.3 kb T7 transcript of *aadA* gene.
3.1.8 Overexpression of the Ycf9 protein in \textit{E. coli}

To obtain an antiserum against Ycf9, the full-length \textit{ycf9} gene was amplified by PCR using a primer pair with introduced restriction sites for EcoRI and XbaI enzymes. The \textit{ycf9} gene was then cloned in the EcoRI/XbaI restricted pThioHis vector (carrying thioredoxin as a fusion protein; Invitrogen, USA). The construct was checked by sequencing analysis, and found to be correct, but no overexpression of the protein was detected after IPTG induction of \textit{E. coli}. Therefore, another strategy for overexpression of a truncated form of the protein was employed. Three polypeptide parts were cloned and individually prepared for overexpression in \textit{E. coli}:

Polypeptide 1: MTLAFQLAVFALIATSLILLISVPVVFASPDGWSSNKNVVFSG (N-terminal hydrophobic region + hydrophilic loop; Figure 16B),

Polypeptide 2: PVVFASPDGWSSNKNVVFSG (hydrophilic loop; Figure 16C),

Polypeptide 3: PVVFASPDGWSSNKNVVFSGTSLWIGLVFLVGILNSLS (hydrophilic loop and hydrophobic C-terminus).

Using an additional primer pair with introduced restriction sites for EcoRI and XbaI (primers \textit{ycf9EcoRI2}, \textit{ycf9XbaI2}) three constructs encoding the C-terminal hydrophobic region, the central hydrophilic loop and the N-terminal hydrophobic region, respectively, were cloned in the EcoRI and XbaI restriction sites of the pThioHis vector. After transformation of \textit{E. coli} and induction with 1 mM IPTG for 3 h the overexpressed polypeptides 1 and 2 with the molecular masses of 3.5 and 2 kDa (increased by the size of the fusion protein thioredoxin to approximately 16 and 15 kDa, respectively; Figure 16A), were detected (Figure 16 B, C). After fractionation of the \textit{E. coli} proteins into soluble and insoluble protein fractions, polypeptides 1 and 2 were found in the soluble protein fraction.
Fig. 16. Overexpression of truncated Ycf9 proteins. Three ycf9 subclones were inserted into pThioHis vector, transformed into E. coli, and the proteins expressed by induction with 1 mM IPTG. Polypeptide 1, representing the N-terminal hydrophobic region with the hydrophilic loop (Panel B), and polypeptide 2, representing only the central hydrophilic loop (Panel C) were successfully overexpressed, resulting in two proteins with molecular masses of 3.5 and 2 kDa, increased by approximately 15 and 13 kDa by thioredoxin (Panel A). Only the soluble protein fractions are presented. Molecular weight markers in kDa are indicated at the left. Fractions 0, 1, 2, 3, and 4 represent cultures without (0), and after 1, 2, 3 and 4 h of IPTG induction. Arrowheads: overexpressed proteins.

3.1.9 Generation of antisera against the Ycf9 protein

To identify the localization of Ycf9 in the chloroplast antisera were raised against two fragments of truncated Ycf9 protein overexpressed in the pThioHis system (Figure 17, Panel A and B), and additionally, against a BSA-coupled, synthetized oligopeptide representing the hydrophilic loop of Ycf9 (PVVFASPDGWSSNKNVVFS; Figure 17 C). Western analysis of the overexpressed proteins demonstrated a strong reaction of all three antisera with overexpressed Ycf9. A strong reaction with overexpressed fusion protein, thioredoxin, was
observed in case of two antisera directed against overexpressed Ycf9. Thus, for further analyses of plant material, antisera directed against the BSA-coupled oligopeptide were used.

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**Fig. 17.** Generation of antisera against Ycf9 protein. Two antisera directed against overexpressed fragments of truncated Ycf9 and one directed against the BSA-coupled, synthetized Ycf9 oligopeptide were used to detect *E. coli*-overexpressed Ycf9. Panels A, B and C represent antisera directed against overexpressed N-terminal hydrophobic and hydrophilic region of Ycf9, hydrophilic loop of Ycf9, and antiserum directed against synthesized Ycf9, respectively. Lane 1: protein extract from non-IPTG induced *E. coli* culture, 2: overexpressed thioredoxin, 3: overexpressed fusion protein thioredoxin-Ycf9 (3 h IPTG induction).

### 3.1.10 Immunological detection of the Ycf9 protein

To determine the localization of the Ycf9 protein, the antiserum raised against Ycf9 was employed to chloroplast subfractions (intact chloroplasts, stroma and thylakoids; Figure 18). Proteins from wild-type and Δycf9 were separated on a 13% Schägger gel and then immunoblotted with polyclonal Ycf9 antibodies. After Western analysis, a signal of approximately 4 - 5 kDa was detected in wild-type, but not in the Δycf9 material, demonstrating that the Ycf9 protein is an integral constituent of thylakoid membranes. For an exact localization of Ycf9 within thylakoid membrane protein complexes, several knock-out tobacco mutants were analyzed using Ycf9 antiserum. Thylakoid membranes isolated from the wild-type, Δycf9 and PSII tobacco mutants, encompassing an insertion of the *aadA* cassette in *psbE, psbF, psbL* and *psbJ* genes, respectively (Swiatek et al. manuscript in preparation), were immunoblotted with Ycf9 antiserum (Figure 19). The Ycf9 signal was present in the wild-type sample, strongly reduced in ΔpsbL and ΔpsbJ plants, which possess
RESULTS

partially active PSII, but absent in Δycf9, ΔpsbE and ΔpsbF mutants, suggesting its association with photosystem II.

Fig. 18. Immunodetection of Ycf9 in tobacco plants grown under 100 µE m⁻² s⁻¹ at 25°C. Antiserum elicited against Ycf9 protein was reacted with the proteins of intact chloroplasts (lanes 1 and 2) and various subchloroplast fractions: stroma (lanes 3 and 4) and thylakoid membranes (lanes 5 and 6).

Fig. 19. Accumulation of PsbE (cytochrome b₅₅₉) and Ycf9 in wild-type, ΔpsbE, ΔpsbF, ΔpsbL, ΔpsbJ and Δycf9 plants. Thylakoid membrane proteins were separated in a 14% Schägger/von Jagow gel, electroblotted onto a PVDF membrane and incubated with antisera raised against Ycf9 or PsbE, as indicated at the right.
3.1.11 Changes in the polypeptide content in thylakoid membranes from Δycf9 mutants

Since immunological analyses of Ycf9 demonstrated its localization in the thylakoid membrane, this chloroplast subfraction was further investigated to reveal possible differences between wild-type and Δycf9 protein content. First analyses included SDS-PAGE gel assay, including silver staining of proteins and serological analyses. Subsequently, more refined techniques, such as two-dimensional gel electrophoresis or sucrose density gradient centrifugation, were applied.

3.1.11.1 SDS-PAGE analysis of the polypeptide content of Δycf9 plants

Figure 20 shows a comparison of thylakoid membrane proteins from heterotrophically grown wild-type and Δycf9 tobacco plants. After denaturing SDS-PAGE and silver staining, the Δycf9 material revealed a polypeptide deficiency in the low molecular weight region of the gels. By analogy to the serological analyses presented in Section 3.1.10, the thylakoid protein content of photosystem II tobacco mutants, encompassing an insertion of the aadA cassette in the psbE, psbF, psbL and psbJ genes, respectively (produced in our group by R. Regel), was examined. All photosystem II mutants, similarly to Δycf9 tobacco, also failed to accumulate an approximately 4 kDa protein. The deficient polypeptide corresponded to the protein immunodetected with Ycf9 antiserum and was attributed to the ycf9 product. In comparing the wild-type and Δycf9 tobacco protein samples, an additional quantitative difference was seen in a region of higher molecular weight, at approximately 26 kDa (arrow at the right part of Figure 20A). Immunological analysis identified this component as CP26, one of the apoproteins of the minor LHC antenna.
Fig. 20. Separation of thylakoid membrane polypeptides from tobacco ∆ycf9, ∆psbE, ∆psbF, ∆psbL and ∆psbJ mutants. Proteins were revealed with silver staining. Tobacco plants were grown under 10 µE m⁻² s⁻¹ at 25°C. In Panel A arrows indicates the positions of polypeptide deficiencies in the ∆ycf9 transformant. The bold arrow indicates the immunologically confirmed Ycf9 band, whereas the lighter arrow indicates CP26. Panel B shows a lower region of the gel with compared wild-type, ∆ycf9, and photosystem II mutants, as indicated at the top of the figure. Arrow indicates the Ycf9 band. Molecular weights in kDa are indicated at the left of each figure.

3.1.11.2 Serological analysis of the ∆ycf9 thylakoid proteins

To examine the content and the distribution of the major photosynthetic complexes in the ycf9 mutant in comparison to wild-type tobacco, several antisera were tested with total thylakoid proteins as well as with fractions from the sucrose gradients (Table 2). Immunodetection was performed using the enhanced chemiluminescence system (ECL). The immunoblots were then scanned and the intensities of the individual signals estimated relative to these of the wild-type using the TINA software package v2.08 beta (Raytest, Spröckhovel).
Table 2. Immunological analysis of the Δycf9 mutants and wild-type tobacco. Signals of intensity identical as in wild-type material are marked (+++), and decreasing intensities are shown as (++) or (+). OEC: oxygen evolution complex.

<table>
<thead>
<tr>
<th>Gene</th>
<th>Product</th>
<th>wild-type</th>
<th>Δycf9 mutants</th>
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<tr>
<td>PsbP</td>
<td>23 kDa protein of OEC</td>
<td>+++</td>
<td>+++</td>
</tr>
<tr>
<td>PsbQ</td>
<td>16 kDa protein of OEC</td>
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</tr>
<tr>
<td>psbA</td>
<td>D1 protein of PSII core</td>
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<td>+++</td>
</tr>
<tr>
<td>psbC</td>
<td>D2 protein of PSII core</td>
<td>+++</td>
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</tr>
<tr>
<td>psbD</td>
<td>CP43</td>
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<td>+++</td>
</tr>
<tr>
<td>psbE</td>
<td>subunit 1 of cyt. b₅₅₉</td>
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<td>+++</td>
</tr>
<tr>
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<tr>
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<td>P700 chl-a protein 1 of PSI core</td>
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<tr>
<td>psaB</td>
<td>P700 chl-a protein 2 of PSI core</td>
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<td>Lhcb1-3</td>
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<td>+++</td>
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<tr>
<td>petB</td>
<td>cytochrome b₆</td>
<td>+++</td>
<td>+++</td>
</tr>
<tr>
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<td>+++</td>
</tr>
<tr>
<td>atpA</td>
<td>α subunit of ATPase complex</td>
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<td>+++</td>
</tr>
<tr>
<td>rbsL</td>
<td>large subunit of RuBisCo</td>
<td>+++</td>
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</tr>
</tbody>
</table>

No difference in the amount of proteins from most photosynthetic complexes was observed in ycf9 mutants compared to wild-type. Nevertheless, a clear deficiency in the amounts of CP29 and CP26 proteins was noted in the mutant.

3.1.11.2.1 The effect of Ycf9 deletion on minor antenna protein complexes

The accumulation of the minor antenna proteins CP29 and CP26 was further investigated by immunoanalyses of thylakoid membrane proteins isolated from plants grown under different light and temperature regimes. As shown in Figure 21, the content of the two minor antenna proteins strongly depends on the growth conditions. In agreement with the report from Ruf et al. (2000), it was found that under most growth conditions, less CP26 accumulated in the ycf9
mutant relative to the wild-type control. The deficiency in CP26 was most pronounced at 20°C in dim light, conditions under which the plants exhibited a dwarf phenotype (Figure 8). In most cases, CP29 content did not vary, with the notable exception of the dwarf Δycf9 plants in which CP29 decreased by about 50%. In contrast, the PsbS protein, an antenna-like subunit associated with PSII, was present in unaltered quantities in both mutant and wild-type plants. Since CP43, a major PSII core antenna subunit, showed no difference between wild-type and Δycf9 plants under each of the chosen growth conditions, it can be concluded that the absence of PsbZ had a selective effect on the accumulation of minor antenna proteins, that is independent of the overall accumulation of PSII cores.

Fig. 21. Effects of the growth conditions on the minor antenna in thylakoid membranes from wild-type (WT) and Δycf9 tobacco. Polypeptides were separated by SDS-tricine PAGE, electroblotted onto nitrocellulose and incubated with the specific antisera indicated at the right of each panel. DL, 10 μE m⁻² s⁻¹; HL, 100 μE m⁻² s⁻¹
3.1.11.3 Isolation of the major protein complexes from the thylakoid membranes

3.1.11.3.1 Isolation of the thylakoid protein complexes by sucrose density gradient centrifugation

Thylakoid protein complexes were fractionated by sedimentation through sucrose gradients, following solubilization of the membrane with β-dodecylmaltoside. Gradients from Δycf9 and wild-type tobacco are compared in Figure 22. The heaviest green band corresponds to PSI supercomplexes, followed by slower sedimenting PSII-LHCII supercomplexes, PSII dimer, PSI+LHCI, PSII monomers, LHCII trimers, LHCII monomers with the minor antenna complex, a yellow, slowly sedimenting fraction enriched in carotenoids, and free proteins at the top of the gradient. All protein complexes were examined immunologically. A clear difference in the accumulation of PSII supercomplexes was observed in Δycf9 material compared to wild-type, as discussed in Section 4.1.

Fig. 22. Comparison of the sucrose gradients from β-dodecylmaltoside solubilized thylakoids from wild-type and Δycf9 plants. The biochemical identification of each chlorophyll-containing band, indicated at the right, was assessed by collecting 0.3 ml fractions for silver staining and analysis using antisera against constituents of the major photosynthetic complexes.

After separation of protein complexes by sucrose density gradient centrifugation, proteins were subsequently separated in 13% Schägger/von Jagow gels (Figure 23). Silver staining of the gels uncovered two small peptides in the molecular weight range of 4 - 6 kDa which were detected in wild-type, but not in the mutant fractions. The band of 4 kDa, corresponding to the
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fractions 19 - 32 containing the PSII complexes, was found in wild-type, but not in Δycf9 material. Therefore, this protein was considered to correspond to the ycf9 gene product. A second band of 6 kDa was also missing from Δycf9 thylakoids. This protein was found in upper regions of the gradient, with a distribution profile that did not match that of LHCII proteins, neither that of the minor antenna complex. The 6 kDa protein may be an unprocessed form of Ycf9 not associated with PSII, or it might be an unrelated protein that is unstable in the absence of Ycf9.

**Fig. 23.** Thylakoid membranes from ycf9 mutant and wild-type plants grown at 25°C under 100 μE m⁻² s⁻¹ were solubilized with β-dodecylmaltoside and separated in a linear 0.1 - 1.0 M sucrose gradient. Proteins from sucrose gradient fractions (600 μl each) were separated in a 13% Schägger/von Jagow gel and revealed by silver staining. Two polypeptide deficiencies, corresponding to bands at 6 kDa and 4 kDa, respectively (arrows), are observed in Δycf9 mutant compared to wild-type. Fraction numbers and protein complexes are indicated above the figure. (wt) = wild-type, (m) = Δycf9 tobacco. Molecular weights in kDa are indicated at the left of the figure; m: monomer, d: dimer, t: trimer, s: supercomplexes.
3.11.3.1.1 Serological analysis of sucrose gradient fractions

Sucrose gradients of thylakoid lysates of Δycf9 and wild-type tobacco are compared in Figure 22. It is evident that in the Δycf9 plants there is little or no material at the position of the PSII-LHCCI supercomplexes (Figure 24). This conclusion was substantiated by immunological analyses. The sucrose gradients were fractionated and the proteins were separated by electrophoresis in denaturing 12% polyacrylamide gels, electroblotted and incubated with antisera raised against D1, LHCCI and the reaction center proteins PsaA and PsaB of PSI. No changes of the PSI RC were noted between Δycf9 and wild-type material, whereas signals of D1 and LHCCI proteins were clearly absent in the fractions corresponding to the PSII supercomplexes in the wild-type tobacco (Figure 24).

![Fig. 24. Immunological analyses of all sucrose gradient fractions with antisera directed against D1, LHCCI and PsaA/PsaB proteins of PSI reaction center. Identities of the complexes and the corresponding pigmented bands in the sucrose gradients are indicated at the bottom of the panels. Fraction numbers are indicated at the bottom of each blot. In the Δycf9 panel (Panel B), vertical arrows mark fractions with a lack of immunoreactivity with D1 and LHCCI antibodies different from the results with wild-type material (Panel A).](image-url)
3.11.3.2 Isolation of the major protein complexes from the thylakoid membranes by two-dimensional gel electrophoresis

3.11.3.2.1 Deriphat-PAGE gel analysis

A two-dimensional gel analysis of the thylakoid membrane proteins from wild-type and Δycf9 material grown under different light and temperature conditions (10 μE m⁻² s⁻¹, 25°C and 20°C; 100 μE m⁻² s⁻¹, 25°C and 20°C) proved the Δycf9-specific deficiency in the minor antenna protein CP26, and the impaired ability of Δycf9 plants to form stable PSII-LHCII supercomplexes.

The Deriphat-PAGE method, in contrast to sucrose density gradient centrifugation, or Blue-Native gel system, allows to separate the entire minor antenna complex, which otherwise comigrates with LHCII monomer. Wild-type and Δycf9 thylakoid membrane preparations were solubilized with 1% α-dodecylmaltoside, and applied to a non-denaturing Deriphat-PAGE. After separation of the protein complexes, slices of the gel, each with separated thylakoid protein complexes, were cut out and loaded onto a denaturing 12% Laemmli gel. After the electrophoretical separation protein spots were visualized using Coomassie staining. For all growth conditions reduction of the signal corresponding to CP26 protein was observed, as shown in Figure 25. No remarkable reduction of CP29 and CP24 signals was detected in the mutant material. Another deficiency observed in ycf9 mutants compared to the wild-type was absence, or strong reduction, of the slowly migrating protein complex representing PSII-LHCII supercomplexes, best detectable in material grown at 25°C under 100 μE m⁻² s⁻¹.
Fig. 25. 2D gel analysis of wild-type and Δycf9 plants grown under different light and temperature regimes. 12.5 µg chlorophyll of the wild-type and the Δycf9 thylakoid membrane preparations were solubilized with 1% α-dodecylmaltoside, applied onto a non-denaturing Deriphat-PAGE gel, and, after separation of the protein complexes, loaded onto a denaturing Laemmli gel with 2nd dimension. The molecular weight markers are indicated for each gel, the upper part of the figure shows the 1st dimension of the non-denaturing PAGE. CP26, CP29 and PSII supercomplexes (s) proteins are indicated by arrowheads. Light and temperature regimes are indicated on the top of the left panel.
3.1.11.3.2.2 Blue-native PAGE analysis

After solubilization by β-dodecylmaltoside, the membrane complexes of wild-type and Δycf9 material were separated by blue-native gel electrophoresis and gel stripes with separated protein complexes were subsequently loaded on denaturing 13% Schägger/von Jagow gels. Silver staining of the gels uncovered two small peptides in the molecular range of 4 - 6 kDa which were seen in wild-type, but not in mutant fractions. The band at 4 kDa, corresponding to fractions containing the PSII dimer and PSII supercomplexes, was located in wild-type but not in Δycf9 material. A second band of 6 kDa was also missing from Δycf9 thylakoids. This protein was found in the upper region of the gel, with a distribution profile that did not match that of LHCII proteins (Figure 26).

![Fig. 26](image)

**Fig. 26.** Thylakoid membrane proteins from Δycf9 (right) and wild-type (left) plants grown at 25°C under 100 µE m⁻² s⁻¹ were separated by two-dimensional gel electrophoresis and revealed by silver staining. Two polypeptide deficiencies, corresponding to 6 kDa and 4 kDa, are observed in the ycf9 mutant compared to wild-type, as indicated by light and bold arrowheads, respectively. The upper part of the figur shows schematically the 1st dimension of the non-denaturing PAGE. Protein complexes are indicated: (m)- monomer, (d)- dimer, (t)-trimer, (s)- supercomplexes. The molecular weights marker in kDa is indicated at the right.
3.1.12 Alterations of protein phosphorylation in the Δycf9 mutant

3.1.12.1 Alterations of protein phosphorylation in the Δycf9 mutant

It is well documented that the interaction between peripheral antenna and PSII cores is controlled by phosphorylation of antenna subunits in a process known as state transitions, by which a portion of LHCII detaches from the PSII cores due to its increased phosphorylation by a redox-controlled kinase (Allen 1992). Thus, the loss of PSII-LHCII supercomplexes in the Δycf9 mutants could result from changes in the phosphorylation patterns of PSII components as well as, or instead of, a direct steric contribution of Ycf9 to the interaction of the PSII cores with their peripheral antenna. To monitor the phosphorylation status of the thylakoid membranes, antibodies raised against phosphothreonine residues were used for immunoblot analyses.

The thylakoid membranes isolated from the tobacco mutant grown under conditions that produced a marked phenotype, displayed two notable changes (Figure 27). Phosphorylation of two PSII core subunits, CP43 and D1, was markedly decreased in Δycf9 material compared to wild-type under the four growth conditions tested (D1 and CP43 phosphorylation patterns in Figure 27). In parallel, LHCII phosphorylation was strongly increased in Δycf9 compared to wild-type when plants were grown at 25°C in high and dim light, whereas the opposite effect was observed at 20°C under dim light. Since the overall accumulation of PSII (as monitored by D1 amounts) and LHCII were similar in wild-type and mutant leaves grown under similar conditions, the differences observed in the phosphoprotein patterns can be attributed to changes in the steady-state phosphorylation levels between PSII and LHCII subunits. The ratio of protein phosphorylation in LHCII vs. PSII was greatly enhanced in the Δycf9 plants under all conditions studied.
Fig. 27. Phosphorylation of thylakoid membrane proteins in wild-type and \( \Delta \text{ycf9} \) mutants. Thylakoid membrane proteins from plants grown either at 25°C and 20°C under 100 \( \mu \text{E m}^{-2} \text{s}^{-1} \) light (HL) or under 10 \( \mu \text{E m}^{-2} \text{s}^{-1} \) light (DL) were separated by SDS-tricine 16.5% polyacrylamide gel electrophoresis, electroblotted onto nitrocellulose and incubated with anti-phosphothreonine, D1, CP43 and LHCII antisera (only samples grown at 25°C, 100 \( \mu \text{E m}^{-2} \text{s}^{-1} \) and 20°C, 10 \( \mu \text{E m}^{-2} \text{s}^{-1} \) light are shown). The proteins were isolated as described in Methods and the protein concentrations in the samples were normalized to the chlorophyll concentration.

3.1.12.2 Influence of light intensities on protein phosphorylation patterns of wild-type and \( \Delta \text{ycf9} \) thylakoids

The influence of the light intensities on phosphorylation patterns of wild-type and \( \Delta \text{ycf9} \) thylakoid proteins was examined by immunoblotting of thylakoid membrane proteins from plant material illuminated by 0 - 1000 \( \mu \text{E m}^{-2} \text{s}^{-1} \) using anti-phosphothreonine antibodies. The results of this Western analysis were scanned and signal intensities were estimated using the TINA software package v2.08 beta (Raytest, Spröckhovel). The phosphorylation of LHCII proteins increased with light intensity until it reached a maximum at approximately 400 \( \mu \text{E m}^{-2} \text{s}^{-1} \).
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2 s\(^{-1}\), then slowly decreased (Figure 28, as shown also by Rintamäki et al. 1997). In \(\Delta yc9\) material, LHCII phosphorylation increased faster and reached a higher maximum level compared to wild-type. An opposite effect was observed on the D1 phosphorylation, which increased faster in wild-type material until it reached a maximum. In \(\Delta yc9\) material the phosphorylation of D1 protein increased slower compared to the wild-type, but finally reached an identical maximal level.

![LHCII phosphorylation at 25°C](image)

![D1 phosphorylation at 25°C](image)

**Fig. 28.** Influence of light intensities on the phosphorylation patterns of wild-type and \(\Delta yc9\) thylakoid proteins. Freshly cut leaves from wild-type and \(\Delta yc9\) tobacco were illuminated for three hours with 0, 100, 200, 400, 600, 800, and 1000 \(\mu\)E m\(^{-2}\) s\(^{-1}\), and, after freezing in liquid nitrogen, membrane proteins were isolated, separated electrophoretically in a 15% PAA gel, electroblotted, and incubated with anti-phosphothreonine antibody. Additionally, the positions of the D1 and LHCII proteins were localized using specific antibodies. For the quantification of phosphoproteins, the immunoblots were scanned and the signal intensities were estimated using the TINA software package v2.08 beta. Upper panel: Phosphorylation pattern of the LHCII proteins at light intensities indicated on the X-axis, lower panel: Phosphorylation pattern of D1 protein at light intensities indicated on the X-axis.
3.1.12.3 Phosphorylation pattern of fractionated thylakoids from the wild-type and $\Delta ycf9$ mutants

For a detailed examination of the amount and localization of phosphoproteins in wild-type and $\Delta ycf9$ plants (grown at 25°C and 100 µE m$^{-2}$ s$^{-1}$) proteins from the sucrose gradient fractions (see Section 3.1.11.3.1) were immunoblotted with anti-phosphothreonine antibodies. As previously observed, stronger phosphorylation of LHCII was noted in the corresponding fractions of the $\Delta ycf9$ mutant compared to wild-type (fractions 4 - 7 in Figure 29). Strong phosphorylation of D1 protein was noted in all wild-type fractions containing PSII (fractions 9 - 10 and 13 - 16 on Figure 29), which was absent in the corresponding $\Delta ycf9$ fractions. Additionally, 24 kDa phosphorylated D1 breakdown product (Kruse et al. 1997, Singh and Singhal 1999) was detected in the D1 monomeric fraction of the wild-type.

**Fig. 29.** Immunoblot analyses of all sucrose gradient fractions (600 µl per fraction) with antisera directed against phosphothreonine. Identities of the membrane complexes, as proved by D1 and LHCII-specific antisera and silver staining of the PAA gel, and fraction numbers are indicated at the top of the figure. Tobacco was grown on B5 medium containing 20 g/l sucrose under 100 µE m$^{-2}$ s$^{-1}$ at 25°C; m: monomer, d: dimer, t: trimer, s: supercomplexes. Arrowhead: phosphorylated degradation product of D1 protein.
To examine the correlation between the phosphorylation of D1 protein and its structural organization in the various PSII complexes, the very same sucrose gradient fractions as presented above were separated by SDS-PAGE, electroblotted and probed with anti-D1 antibodies (Figure 30). For quantification the immunoblots were scanned and the intensity of signals was estimated using the TINA software package v2.08 beta.

**Fig. 30.** Immunoblot analyses of all sucrose gradient fractions (600 µl per fraction) with antisera directed against D1. Identities of the complexes and fraction numbers are indicated at the top of the figure. Tobacco was grown on B5 medium containing 20 g/l sucrose under 100 µE m⁻² s⁻¹ at 25°C; m: monomer, d: dimer, t: trimer, s: supercomplexes. Arrows: degradation products of D1 protein. For comparison see Figure 29.

After comparison of Figure 29 and 30 it is evident that the phosphorylation of D1 protein in wild-type plants correlates with an increased formation of the dimeric form of D1, which is necessary to form supercomplexes (Barbato et al. 1992, Kruse et al. 1997). Lack of D1 phosphorylation in Δycf9 may be one reason for an increase of the monomeric D1 amount and an absence of the PSII-LHCII supercomplexes. Interestingly, D1 degradation products were detected by both anti-D1 and anti-phosphothreonine antibodies in the monomeric D1 fraction of wild-type tobacco, but not in the mutant material, even when an increased amount of the monomeric D1 protein was observed.
3.1.12.4 In vitro phosphorylation of thylakoid proteins

To examine possible qualitative or quantitative differences in the $\Delta ycf9$ and wild-type protein phosphorylation, an in vitro phosphorylation of thylakoid membranes was performed as described in Methods. Plants grown at all previously described light and temperature regimes were analysed in this experiment, although no phosphorylation differences were observed between the different phenotypes of $\Delta ycf9$. No qualitative notable difference was detected between wild-type and $\Delta ycf9$ phosphoproteins (Figure 31). However, a clear quantitative in vitro phosphorylation difference was noted between wild-type and $\Delta ycf9$ material. Under all conditions tested an approximately three times stronger phosphorylation of $\Delta ycf9$ proteins was observed as compared to wild-type.

Fig. 31. In vitro phosphorylation of thylakoid membrane proteins from wild-type and $\Delta ycf9$ plants grown under 100 µE m$^{-2}$ s$^{-1}$ at 25°C. The phosphorylation assay was performed as described in Methods and the protein concentrations in the samples were normalized to the chlorophyll concentration. For each wild-type and $\Delta ycf9$ sample 20, 10 and 7 µg of chlorophyll were loaded onto the gel (as indicated at the top of the panel: lanes 1, 2 and 3, respectively). The molecular weights in kDa are indicated at the right. Arrowheads: D1 and LHCII phosphorylation.
3.1.13 Non-photochemical quenching in the Δycf9 plants

For protection against photooxidative damage, occurring when the light energy absorbed becomes excessive relative to the capacity of photosynthesis, photosynthetic organisms developed non-radiative pathways that dissipate excess light energy absorbed by the antenna system. The xanthophyll cycle and the non-photochemical quenching of chlorophyll fluorescence (NPQ) represent processes preventing a photooxidative damage of the photosynthetic apparatus induced under high light. Non-photochemical quenching provides non-photochemical thermal dissipation of the absorbed excess photons. Thermal dissipation of the energy competes with the fluorescence emission, thus NPQ induction can be measured as a function of fluorescence quenching. Comparative measurements of the maximum level of fluorescence in dark-adapted samples (Fmax) and after a period of high light treatment (F’re), allows to calculate NPQ as (Fmax – F’re)/F’max (Horton 1996). Figure 32 shows the induction and relaxation of NPQ in wild-type and Δycf9 tobacco plants grown under different light and temperature conditions. Initially, 10 minutes exposure under bright light (2200 µE m⁻² s⁻¹) caused a typical NPQ induction in both wild-type (black circle) and Δycf9 (red circles). This process has two kinetic components, a fast phase that was completed within the first minute of illumination, and a slow phase that developed steadily over the next 9 minutes. Recovery took place in darkness with a half-life of about 6 minutes. The amplitude of the NPQ developed upon light stress was dependent on the light and temperature conditions applied for the wild type and mutant plants (Table 3 and Figure 32). In the wild type the adaptation to high light or low temperature stress induced an increase in the non-radiative energy dissipation (NPQ amplitude). In contrast, the Δycf9 mutant displayed an opposite NPQ behaviour under the same experimental conditions, namely the decrease of the NPQ amplitude under the higher stress conditions (Table 3). At constant growth temperature, increasing the light intensity decreased the NPQ amplitude. Similarly, decreasing the temperature under constant light intensity also decreased the NPQ amplitude. For Δycf9 plants grown under 100 µE m⁻² s⁻¹, the NPQ amplitude was 37% lower at 17°C than at 25°C. The upper panel of Figure 32 also shows that at 17°C the rate of NPQ relaxation in the dark was strongly affected in the mutant. After 10 minutes of recovery in darkness, the quenching in the wild-type leaves was reversed by 85%, whereas only 30% recovery was recorded for Δycf9. Moreover, 50% of the recovery in the wild-type occurred within the first 2 min of dark adaptation, representing the rapidly relaxing ΔpH-dependent (qE) component of NPQ (Horton 1996). The following slow phase of recovery reflects the slowly relaxing, photoinhibition
component (qI). Only the second, slow phase of recovery was recorded in the mutant suggesting that Δycf9 plants may have lost part of the adaptation mechanism which dissipates excess light energy under stress conditions (higher light, sub-optimal growth temperature; Huner et al. 1998). The ratios of NPQ amplitudes between Δycf9 and wild-type plants are presented in Table 3.

**Table 3.** Maximal amplitude of quenching following 10 min exposure to supersaturating light. Measurements were performed at room temperature with plants grown at different light and temperature regimes. Before exposure to supersaturating light, plants were dark-adapted for 10 min. DL: 10 µE m⁻² s⁻¹, HL: 100 µE m⁻² s⁻¹.

<table>
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<th>17°C</th>
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Fig. 32. Non-photochemical quenching in intact leaves of wild-type (black circles) and Δycf9 (red circles) tobacco. Plants were grown on B5 medium containing 20 g/l sucrose under 100 μE m$^{-2}$ s$^{-1}$ (right) or 10 μE m$^{-2}$ s$^{-1}$ (left), and 17°C (upper panel), 20°C (middle panel) or 25°C (lowest panel). Non-photochemical quenching was induced with 2200 μE m$^{-2}$ s$^{-1}$ continuous light for 10 min, and the recovery took place in darkness.
3.1.14 Pigment composition and xanthophyll cycle in the $\Delta ycfs$ plants

As mentioned above, NPQ has been correlated with the activation of the xanthophyll cycle, which promotes a reversible, $\Delta$pH-induced deepoxidation of the xanthophyll violaxanthin to zeaxanthin via the intermediate antheraxanthin catalyzed by the enzyme violaxanthin deepoxidase. Deepoxidation of violaxanthin is involved in the conversion of PSII to a state of high thermal energy dissipation and low chlorophyll fluorescence emission (Demmig-Adams 1990). The composition of the xanthophyll cycle pigments was investigated in the wild-type and in $\Delta ycfs$ tobacco (Table 4). During the experiment plants were first exposed to dim light, and a growth regime of 20°C under 10 $\mu$E m$^{-2}$ s$^{-1}$, which was followed by an exposure to increased light intensity (900 $\mu$E m$^{-2}$ s$^{-1}$) for 30 minutes (HL), and then by a transfer to darkness at 20°C for 10 minutes (HL+dark). The wild-type plants initially contained more than 95% violaxanthin which was subsequently converted into zeaxanthin under high light. This pigment modification corresponded to a de-epoxidation state (DS) of 0.33, with DS = $(Z+1/2A)/(Z+V+A)$ (Demmig-Adams and Adams 1992). After recovery in darkness for 10 minutes, the zeaxanthin pool decreased to about 20% of total, with a DS of 0.14. The high-light treatment of $\Delta ycfs$ plants caused a strong increase of the de-epoxidation state, with only 25% of the pigments from the xanthophyll cycle remaining as violaxanthin. Significant amounts of de-epoxidized forms, including both zeaxanthin and antheraxanthin, accumulated in parallel to the disappearance of violaxanthin (DS = 0.63). After 10 minutes recovery in the dark, the re-epoxidation was much slower than in the wild-type and the mutant leaves still contained an excess (65%) of de-epoxidized xanthophylls, corresponding to a DS of 0.56. Additionally, a higher amount of total carotenoids was detected in $\Delta ycfs$ plants as compared to the wild-type. This effect was dependent on the applied light intensities, and varied from 16% under dim light (10 $\mu$E m$^{-2}$ s$^{-1}$), to 48% after 30 minutes illumination under high light. No such effect was observed for wild-type plants. Interestingly, the newly synthesized carotenoids belonged to the xanthophyll cycle pool while the content of other carotenoids, such as lutein, neoxanthin and $\beta$-carotene, remained constant. The increase in carotenoid content was not reversible during the following incubation for 10 minutes in the dark. Thus, light stress conditions specifically increased the xanthophyll pool available for deepoxidation in the mutant plants.
### Table 4. HPLC analyses of changes in the pigment composition induced by high light.

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<td>β-carotene</td>
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<td>4.3</td>
</tr>
<tr>
<td>lutein</td>
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<td>14.2</td>
</tr>
<tr>
<td>VAZ</td>
<td>6.8</td>
<td>10.9</td>
<td>6.8</td>
</tr>
<tr>
<td>(Z+1/2A)/(VAZ)</td>
<td>0.044</td>
<td>0.050</td>
<td>0.33</td>
</tr>
<tr>
<td>Chl a</td>
<td>71.2</td>
<td>73.8</td>
<td>75.2</td>
</tr>
<tr>
<td>Chl b</td>
<td>28.8</td>
<td>26.2</td>
<td>24.8</td>
</tr>
<tr>
<td>Total Carotenoids</td>
<td>31.2</td>
<td>36.4</td>
<td>31.7</td>
</tr>
</tbody>
</table>

1 Pigment composition was normalized to 100 moles of total (\(a+b\)) chlorophyll. V: violaxanthin; A: antheraxanthin; Z: zeaxanthin; Total Car: total carotenoids; VAZ: V+A+Z. The ratio (Z+1/2A)/(VAZ) refers to the de-epoxidation state as defined in Demmig-Adams and Adams (1992).

2 Pigment composition was measured in leaves of tobacco plants grown at 20°C, 10 µE m\(^{-2}\) s\(^{-1}\) (DL), exposed to high light (900 µE m\(^{-2}\) s\(^{-1}\)) for 30 min (HL), and allowed to recover in darkness for 10 min (HL+dark). Analyses were performed by HPLC of 80% acetone leaf extracts.

### 3.1.15 Chlorophyll fluorescence analysis

#### 3.1.15.1 77K chlorophyll fluorescence analysis

#### 3.1.15.1.1 Analysis of PSI and PSII supercomplexes assembly; excitation spectra by emission at 735 and 690 nm

Measurement of excitation spectra by emission at 735 or 690 nm allows to distinguish the amount of pigments, including LHC antenna, attached to PSI and PSII, respectively, reflecting the formation of PSI and PSII supercomplexes in the thylakoid membrane. Both
wild-type and Δycf9 thylakoid membranes isolated from plants grown under different light and temperature conditions (10 and 100 µE m⁻² s⁻¹, 20°C and 25°C) were examined. Representative results of the analysis of plant material grown at 25°C, 100 µE m⁻² s⁻¹ are shown in Figure 33.

Fig. 33. 77K analyses of the wild-type and Δycf9 plants. The excitation spectra were measured with emission wavelengths of 690 nm (PSII) and 735 nm (PSI). The peaks at 440 nm represent reaction centers (chlα) of PSI (Panel A) and PSII (Panel B) and the peaks at 475 nm represent PSI and PSII reaction centers with associated LHC antenna (chlβ), corresponding to PSI and PSII supercomplexes (Panels A and B, respectively).

In agreement with results presented in Section 3.1.11, the excitation spectra of PSI and PSII of wild-type and Δycf9 plants show differences in the assembly/stability of PSII and PSI supercomplexes. While a decreased amount of PSII reaction centers with associated antenna was observed in the Δycf9 material compared to wild-type under all conditions tested, an opposite effect was true for the PSI.

3.1.15.1.2 Chlorophyll b emission spectra

Chlorophyll emission spectra after excitation at a wavelength of 475 nm enable to calculate the ratio of PSI/PSII as well as the amount of LHC antenna attached to both reaction centers. This analysis of plants grown under different light and temperature regimes offers the possibility to investigate the process of state-transitions. Wild-type and Δycf9 thylakoid
membranes isolated from plants grown under different light and temperature conditions (20°C and 25°C, 10 and 100 µE m² s⁻¹) were investigated.

Fig. 34. Chlorophyll b emission spectra of the wild-type and Δycf9 tobacco. Excitation of chlorophyll b at 475 nm results in two peaks representing PSII at 690 nm and PSI at 735 nm. Plants grown under light and temperature regimes as indicated above and on the left side of the graphs were analysed. The calculated PSI/PSII ratio is presented in the Table 5.

Table 5. PSI/PSII ratio of wild-type and Δycf9 tobacco grown under different light and temperature regimes.

<table>
<thead>
<tr>
<th></th>
<th>25°C, 100 µE m² s⁻¹</th>
<th>25°C, 10 µE m² s⁻¹</th>
<th>20°C, 100 µE m² s⁻¹</th>
<th>20°C, 10 µE m² s⁻¹</th>
</tr>
</thead>
<tbody>
<tr>
<td>PSI/PSII ratio</td>
<td>WT</td>
<td>Δycf9</td>
<td>WT</td>
<td>Δycf9</td>
</tr>
<tr>
<td></td>
<td>2.1</td>
<td>2.4</td>
<td>1.0</td>
<td>2.3</td>
</tr>
</tbody>
</table>

Measurements at 25°C with wild-type tobacco showed a correlation of the redistribution of LHC antenna from PSII to PSI with increasing light intensity, a process which can be
attributed to the state-transition mechanism. No such effect was observed for \( \Delta ycf9 \) material. Moreover, under all growth conditions tested the amount of LHC antenna attached to PSII was reduced in \( \Delta ycf9 \) compared to the wild-type. No clear correlation between the distribution of LHC antenna and light intensities was observed at lower temperatures. It has been previously demonstrated (Verhoeven et al. 1999) that low temperature may cause the same photoinhibitory effect as high light treatment. Thus, it can not be excluded that the conditions applied additionally affected the size or distribution of the LHC pool in both, wild-type and \( \Delta ycf9 \) material.

3.1.15.2 Chlorophyll fluorescence measurements at room temperature

For the spectroscopic analysis of chlorophyll fluorescence wild-type and \( \Delta ycf9 \) tobacco grown under different light and temperature regimes (10 and 100 µE m\(^{-2}\) s\(^{-1}\), 20°C and 25°C) were analyzed. The maximum fluorescence (\( F_M \)) and the ratio \( (F_M-F_0)/F_M = F_V/F_M \), as well as the fluorescence quenching parameters \( q_P \) (photochemical quenching) = \( (F_M-F_S)/(F_M-F_0) \), and \( q_N \) (non-photochemical quenching) = 1- \( (F_M-F_0)/(F_M-F_0) \) (Schreiber 1986) were determined for all samples. The averages of at least three measurements are presented in Table 6.

**Table 6.** Spectroscopic analyses of wild-type and \( \Delta ycf9 \) mutant plants.

<table>
<thead>
<tr>
<th></th>
<th>WT 25°,100µE</th>
<th>( \Delta ycf9 ) 25°,100µE</th>
<th>WT 25°,10µE</th>
<th>( \Delta ycf9 ) 25°,10µE</th>
<th>WT 20°,100µE</th>
<th>( \Delta ycf9 ) 20°,100µE</th>
<th>WT 20°,10µE</th>
<th>( \Delta ycf9 ) 20°,10µE</th>
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</thead>
<tbody>
<tr>
<td>( F_V/F_M )</td>
<td>0.80</td>
<td>0.73</td>
<td>0.76</td>
<td>0.75</td>
<td>0.75</td>
<td>0.78</td>
<td>0.79</td>
<td>0.76</td>
</tr>
<tr>
<td>( q_P )</td>
<td>0.96</td>
<td>0.93</td>
<td>1.00</td>
<td>0.94</td>
<td>1.05</td>
<td>1.01</td>
<td>0.99</td>
<td>0.99</td>
</tr>
<tr>
<td>( q_N )</td>
<td>0.26</td>
<td>0.24</td>
<td>0.28</td>
<td>0.18</td>
<td>0.34</td>
<td>0.36</td>
<td>0.29</td>
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</tbody>
</table>

Measurements of the maximum fluorescence (\( F_M \)) and the ratio \( F_V/F_M \) allow to estimate the electron flow through PSII and the photosynthetic activity of plants. No differences between wild-type and \( \Delta ycf9 \) was observed, indicating unaltered photosynthetic activity of \( \Delta ycf9 \) tobacco.
3.2 Functional analysis of the ycf10 (cemA) gene product

3.2.1 Comparison of the Ycf10 sequences

The plastid-encoded ycf10 gene (ORF 229, cemA) from photosynthetically active organisms encodes a highly conserved polypeptide of 229 - 231 amino acid residues (Figure 35). The amino acid sequence deduced for the Ycf10 protein seems to contain four membrane spanning domains (see hydrophathy plot of the Ycf10 protein in Figure 36), and a putative cleavage site for a signal peptidase between the residues Ser-24 and Phe-25 (Willey and Gray 1990, Sasaki et al. 1993)

<table>
<thead>
<tr>
<th>YCF100E1</th>
<th>YCF100E4</th>
<th>YCF10NT</th>
<th>YCF10AT</th>
<th>YCF10SO</th>
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<td>SLEMIQLFLI</td>
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<th>YCF10AT</th>
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<td>ASLQYIGCLFL</td>
<td>FIPLGV8SF</td>
<td>QKCFL5WPQ</td>
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<tr>
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<td>---<em>.</em>.</td>
<td>---<em>.</em>.</td>
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<td>---<em>.</em>.</td>
<td>---<em>.</em>.</td>
<td>---<em>.</em>.</td>
</tr>
</tbody>
</table>
RESULTS

Fig. 35. Comparative analysis of Ycf10 amino acid sequences. The consensus is indicated at the bottom: 80 - 100% of identity among all sequences is marked by asterisks, identity from 60% - 80% by dots, identity below 60% by a dash. Abbreviations: OE1,4: *Oenothera elata* plastome I and IV, NT: *Nicotiana tabacum*, AT: *Arabidopsis thaliana*, SO: *Spinacia oleracea*, ZM: *Zea mays*, OS: *Oryza sativa*, PT: *Pinus thunbergii*, MP: *Marchantia polymorpha*. The predicted membrane helices are underlined, the putative cleavage site is marked in bold letters.
Fig. 36. Hydrophathy profile of Ycf10 (CemA; according to Kyte and Doolittle 1982). Amino acid positions are indicated at the bottom of the graph. The scan window size was 13 amino acid residues. Four hydrophobic, trans-membrane α-helices, designated (I), II, II and IV are indicated above the zero line. The putative cleavage site is marked by arrow.

3.2.2 Construction of a recombinant plasmid for ycf10 inactivation

An inactivation vector for the ycf10 gene was constructed by introducing the promoterless aadA cassette into the BbsI restriction site in the ycf10 gene. The correct integration of the aadA gene onto the pGEMycf10 construct was monitored by PCR, restriction and nucleotide sequence analyses.

Fig. 37. Construction of the plastid transformation vector for disruption of the ycf10 gene. The KpnI/SalI fragment of the plasmid pTB22 (Shinozaki et al. 1986), containing the ycf10 gene and approximately 1 kb flanking regions was cloned into the pGEM vector. The inactivation of the ycf10 gene was performed by insertion of the aadA cassette into a gene-internal BbsI site. A plasmid clone with the aadA cassette inserted in the same orientation as ycf10 was used for plant transformation. The transcription direction of the aadA gene is indicated by an arrow.
3.2.3 Plant transformation, selection and regeneration of mutants

Twenty leaves of *Nicotiana tabacum* were bombarded with gold particles carrying the recombinant plasmid pGEMycf10. After three days of cultivation on non-selective RMOP medium transformed leaves were cut and transferred to RMOP medium supplemented with spectinomycin. Resistant green colonies (calli) began to appear after approximately three weeks and were transferred onto fresh selective RMOP medium. The transformation yielded three independent mutant lines. Segregation of the mutant plastomes was enhanced by transferring shoot tops onto fresh selection medium in three-week intervals. For molecular analyses, the mutant lines underwent at least five cycles of shoot regeneration.

3.2.4 Growth phenotypes of the Δycf10 plants

Δycf10 plants were regenerated from the green calluses on antibiotic-supplied medium, and further cultivated parallel *in vitro* and in the greenhouse. All three mutant ycf10 lines were able to survive on soil and produced fertile seeds. This confirmed that their phenotype was photoautotrophic. However, the material showed a clear non-wild-type phenotype. Under standard heterotrophic conditions (100 µE m⁻² s⁻¹, 25°C, medium supplemented with sucrose) Δycf10 plants developed dark green leaves with numerous pale green to white regions, generally present at the margins of the leaves (Figure 38). The leaves of the Δycf10 plants have a prolonged shape and longer petioles compared to those of wild-type material. This effect was less dominant when plants where grown on soil in the greenhouse.
**Fig. 38.** Growth phenotypes of Δycf10 plants. Panels A and C: Δycf10 plants grown *in vitro* (100 µE m⁻² s⁻¹, 25°C, growth medium supplemented with sucrose) are compared with wild-type tobacco. White regions of Δycf10 leaves are indicated by arrows in panel C. Panel B: wild-type and Δycf10 plants grown photoautotrophically under greenhouse conditions.
3.2.5 Ultrastructure of the Δycf10 chloroplasts

Electron microscopy of wild-type and Δycf9 plants was performed in the laboratory of Prof. G. Wanner (Botanisches Institut, München). As previously described, the Δycf10 plants displayed a characteristic phenotype of dark green leaves with numerous pale regions. Both areas were examined by electron microscopy. The green parts of Δycf10 leaves showed a chloroplast structure indistinguishable from wild-type, with numerous stroma and grana thylakoids and evenly distributed plastoglobules, whereas the bleached regions of the leaves displayed a substantially altered structure, without defined stroma and grana thylakoids (Figure 39). Further experiments (see Section 3.2.12) demonstrated that pale leaf regions lacked photosynthetic activity.

Fig. 39. Electron micrographs of plastids from wild-type and Δycf10 plants of Nicotiana tabacum. (a) Typical lens-shaped wild-type chloroplast with numerous stroma (ST) and grana (GT) thylakoids, (b) chloroplast from green regions of the Δycf10 mutant indistinguishable from wild-type, (c) plastid from a pale leaf region of the Δycf10 mutant: no grana or stroma thylakoid structures were detected. Large plastoglobules (Pg) show a uniform distribution.
3.2.6 Investigation of the homoplastomic state of the \( \Delta ycf10 \) mutants

To confirm the homoplastomic state of \( \Delta ycf10 \) plants the strategy outlined in Section 3.1.6 was employed, including nucleotide sequence analysis, Southern analysis and PCR with total, plastid and PFGE - purified DNA.

3.2.6.1 Nucleotide sequence analysis

To confirm the correct integration into the tobacco plastid chromosome of the construct carrying the \( aadA \) gene, the segment of plastid DNA encoding \( ycf10 \) was sequenced in transformants. For this purpose, an amplification product of \( ycf10 \)for and \( ycf10 \)rev primers was purified from an agarose gel that served as a template for nucleotide sequencing (Hupfer et al. 2000). This analysis established the successful disruption of \( ycf10 \). Moreover, insertion of the \( aadA \) causes a frame-shift in the gene, preventing a translational re-initiation of the transcript containing the C-terminal gene segment.

```
63347    TGTGTGTAAAG AAATATTCGA TCGCATAGAG TGTACGAATG GGTTGATTAA
63397    CAATTCACAG ATGAAAAAAAT GGCAAAAAAAG AAAGCATTCA CTCTCTCTTTTT
63447    CTATCTTGCA TCTATAGTAT TTTTGCCCTG GTGGATTTCT TTCTCAGTTA
63497    ATAAATGTCT GGAATCTTGG GTTACCAATT GGTGAATAC TGGGCAATCC
63547    GAAATTATTT TGAATAATAT TCAAGAAAAA ATTCGCCGTC GTTCAATGAG
                AATGGATAAG AGGCTCGTGG GATGGACGTG AGGGGCAGG GATGGCTATA
                TTTCTGGGAG CGAACTCCCG GCCATATCAG TAGTTGTAGG GAGGGATCCA
                TGGCTCGTGA AGCGTTATAC GCCGAAGTAT CAACTCAACT ATCAGAGGTA
                GTTGGCGTCA TCGAGCCGCA TCTCGAACCAG AGCTTGTCTG GCGTGACTTTT
                GTACGGCTCC GCAGTGGATG GCGGCCTGAA GCCACACGAT GATATTTGATT
                TGCTGTTTAC GTGAGGCRTA AGGCTTGATG AAACAACCGC GCCAGCTTTT
                ATCAACGACC TTTTGGAAAC TCTCCGTCCC CCTGGAGAGA GCGAGGCTTA
                CCGGTCTGTA GAAGTCACCA TTGGGCTGAA CTGCAATTTG GAGAATGGCA GCGCAATGAC
                ATTCTTGCAG GTATCTTCGA GCCAGCCGAA CTGCAATTTG GAGAATGGCA GCGCAATGAC
                ATTCGTCAG GTATCTTCGA GCCAGCCGAA CTGCAATTTG GAGAATGGCA GCGCAATGAC
```
RESULTS

CTTGCTGACA AAAGCAAGAG AACATAGCGT TGCTTGGTA GGTCCAGCGG
CGGAGGAACT CTTTGATCGG GTTCTGAAC AGGATCTATT TGAGGCCGCTA
AATGAAACCT TAAGCCTATG GAACGCTGCG CGCGACTGGG GTGGCGATGA
GGCAAAATGTA GTGCTTACGT TGCTCCGACAT TGGTACAGC GCAGTAACCG
GCAAAACTCGC GCCGAAGGAT GTGCTTGCGG ACTGGGCAAT GGAGCGCCTG
CCGGCCCCAGT ATCAGGCCGCT CATACTTGAA GCTAGACAGG CTTATCTGAG
ACAAGAAGAA GATCGCGTGG CTTGCGCGGC AGATCAGTTG GAAGAATTTG
TCCACTACGT GAAAGCGGAG ATCAGCTAAGG GCTACATCGA AATAACTGCAG

63576 GCATGCAAGC TAAAAAGGTC TTCTAGAAAA ATTCATAGAA TTAGAGGAAC
63611 TCCTCTTCTT GGACGAAATG ATCAAGGAAT ACTCGGAAAC ACATCTCGAA
63661 GAGTCTTGGGA TAGGAATCCA TAAAGAAGCG TACGAATTTAA TCAAGGATAC
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63811 ATTCCTAATG CTGGGCTGCA GGAATTTCTA TATAACTTAA GTGACACAGT
63861 AAAAGCTTTT TCTATCTCTT TATTAACCTGA TTTATGTTAT GGAATCCATT
63911 CACCCCAAGG TTGGGAATTA ATGATTGCTG CTATCTTCAA CAGATTTTGA
63961 TTGGCTCATA ATGATCAATA TATATCGTCT TCTCTTCTCA CTCCTCTGAGT
64011 CATCTCGAT ACAATTCTTAA AATATTGGAT TTTCCGTAT TTAGATCGTC
64061 TGTCTCGAGTC ACTTGTTGGG ATTTATCGAT CAAATGAAATG CTGATAAAAGG
64111 ATCCATTGGAT ATTAATCTAA TCCAATTAGA ATGCCTTGGTA CTTTGTAGTT
64161 GTACATAAGC AGAGTATTTG

Fig. 40. Nucleotide sequence of the ycf10 gene from a transformed tobacco lines. Start and stop codons of orf229 are indicated in bold letters. The aadA sequence is underlined.
3.2.6.2 Southern analysis

Plant material after 6 - 8 regeneration cycles on selective medium was chosen to test the proportion of transformed/wild-type plastome copies by Southern analysis. The isolated plastid DNA was used for RFLP analysis and subsequent hybridization with a radiolabeled ycf10 probe. In all transgenic lines no hybridization signal for wild-type plastome copies was detected.

![RFLP analysis](image)

**Fig. 41.** RFLP analysis to test homoplastomy of the ycf10 knock-out mutants. Plastid DNA of wild-type and transformed plants was digested with EcoRI and hybridized to a radiolabeled ycf10 probe (PCR product). Restriction yields a 585 bp fragment for the WT plastome and a 1.4 kb fragment for the ycf10 transformant. The lack of the wild-type band in the transformant DNA establishes the homoplastomy of the ycf10 knock-out mutant; 1: wild-type plastid DNA, 2: Δycf10 plastid DNA.

3.2.6.3 PCR analysis

To confirm the homoplastomic state of the Δycf10 plants the more sensitive PCR method was applied. The reaction was performed using a primer pair designed for the regions flanking the ycf10 gene (ycf10forPF and ycf10revPF). This primer combination allowed to distinguish between the signals originating from transformed and wild-type DNA sequences. The signal expected from the wild-type plastome was apparent when total DNA was used as a template. To reduce the possibility of contamination of the sample by nuclear and/or mitochondrial DNA sequences, Percoll-gradient purified plastid DNA was subsequently used as template for PCR. A strong reduction of the wild-type plastome product was detected, although the signal arising from the hybridization between the transformed and wild-type amplification products was still present. Subsequent analyses of plastid DNA purified by pulsed-field gel electrophoresis lacked the wild-type signal in Δycf10 material, thus establishing its homoplastomy (Figure 42).
3.2.6.3.1 Hybridization analysis of Δycf10 PCR products

To confirm the Δycf10 PCR analyses, reaction products were hybridized to a radiolabeled probe. The PCR products were electrophoretically separated in an agarose gel, visualized by ethidium bromide staining, as presented in Figure 42, subsequently blotted onto nylon membrane, and hybridized with a radiolabeled ycf10 specific probe (Figure 43). In contrast to the total cellular DNA and DNA from gradient-purified plastids, no wild-type-specific signal was found in PFGE-purified plastid DNA sample (Figure 43). The detection level of this approach was estimated at one wild-type DNA copy detectable in one plant cell.

Fig. 42. An example of PCR analysis of Δycf10 plants. The ycf10 fragment was amplified from wild-type and transformed plants using the ycf10forPF and ycf10revPF primers. The wild-type signal should appear as a 200 bp band, and the transplastomic fragment should be 900 bp larger (1100 bp). Different preparations of plant DNA were used to investigate the homoplastomic state of the mutant, as indicated at the top of the picture. 0: zero control, t: total DNA, pt: plastid DNA, pf: pulsed-field gel isolated DNA.
3.2.7 Northern analysis

In tobacco, the ycf10 gene was shown to be located upstream of the petA gene encoding cytochrome f (Shinozaki et al. 1986), and being cotranscribed with that gene in pea (Willey and Gray 1990; Nagano et al. 1991). The complex pattern of the transcripts suggests that in tobacco the ycf10 gene is a part of a gene cluster including also psaI, encoding PsaI subunit of PSI, ycf4 upstream and petA downstream of ycf10. The presence of at least two promoters can be deduced from the transcription map of the psaI-ycf4-ycf10-petA operon (Figure 44). RNA filter hybridizations performed with all relevant probes (ycf10, ycf4, petA, aadA) were consistent with the above described genome structure. Four different transcripts emanating from the psaI-ycf4-ycf10-petA cluster were identified, which were about 0.9 kb (aadA cassette) larger in the Δycf10 plants. Two additional transcripts, arising from the aadA gene promoter, were localized in the Δycf10 material.
**Fig. 44.** Deduced maps of the *psaI-ycf4-ycf10-petA* gene cluster in wild-type (left) and ∆ycf10 (right) tobacco chloroplasts. Potential promoters are shown as bent arrows. Below each map, horizontal lines indicate the positions of the probes used for RNA filter hybridization. The deduced transcripts are shown below the maps. Transcript sizes in wild-type are: T1: 4.4 kb, T2: 1.9 kb, T3: 1.7 kb, T4: 1.4 kb. In the transformant the sizes of the transcripts modified by the *aadA* insertion (indicated by asterisks) are: T1*: 5.3 kb, T2*: 2.8 kb, *aadA*-specific transcripts in the mutant are T5: 2.6 kb and T6: 1.4 kb.
3.2.8 Localization of the $ycf10$ product

To localize the Ycf10 protein polyclonal monospecific antibodies, generated against Ycf10 from pea (kindly provided by Dr. Sasaki), were tested on thylakoid membranes, stroma and inner chloroplast envelope fractions isolated from chloroplasts of wild-type and the $\Delta ycf10$ tobacco. The pea inner envelope preparation served as a positive control and showed cross-reaction of two proteins of approximately 30 and 26 kDa with the antiserum. A single band of 30 kDa was observed in tobacco wild-type, but not $\Delta ycf10$ inner envelope fraction. No cross-reaction was obtained with thylakoid and stroma subfractions of chloroplasts (Figure 45, Panel A). To estimate the purity of chloroplast subfractions, the samples from the wild-type and $\Delta ycf10$ tobacco were reacted with an antiserum directed against mixed envelope proteins (Figure 45, Panels B and C).

Fig. 45. Panel A: Localization of the Ycf10 protein in chloroplast subfractions; Lane 1: wild-type stroma proteins, lane 2: $\Delta ycf10$ stroma proteins, lane 3: wild-type thylakoid proteins, lane 4: $\Delta ycf10$ thylakoid proteins, lane 5: pea inner envelope proteins, lane 6: wild-type tobacco inner envelope proteins, lane 7: $\Delta ycf10$ inner envelope proteins. Panel B: Immunoblot analysis of chloroplast subfractions with antiserum directed against inner and outer chloroplast envelope proteins; lane 1: wild-type thylakoid membranes, lane 2: $\Delta ycf10$ thylakoid membranes, lane 3: wild-type stroma fraction, lane 4: $\Delta ycf10$ stroma fraction. Panel C: Immunoblot analysis of inner envelope preparations from wild-type (Lane 1) and $\Delta ycf10$ (Lane 2) tobacco with antiserum directed against inner and outer chloroplast envelope proteins. Molecular weights in kDa are indicated at the right of each panel.
3.2.9 Isolation of thylakoid protein complexes from Δycf10 mutants by sucrose density gradient centrifugation

The isolation and characterization of thylakoid protein complexes from wild-type and Δycf10 plants was performed as described in Section 3.1.10. As expected, no qualitative nor quantitative differences in thylakoid protein composition were detected in the Δycf10 mutants compared to wild-type, as shown on Figure 46.

**Fig. 46.** Comparison of the sucrose gradients from β-dodecylmaltoside solubilize thylakoids from wild-type and Δycf10 plants. The biochemical identification of each chlorophyll-containing band, indicated at the right, was assessed by collecting 0.3 ml fractions for silver staining and analysis using antisera against constituents of the major photosynthetic complexes (see Section 3.1.11.3.1).

3.2.10 Serological analysis of the Δycf10 mutants

To examine the content and distribution of the major photosynthetic complexes in the Δycf10 mutants in comparison to wild-type tobacco, several antisera were tested on membranes containing total thylakoid proteins of wild-type and Δycf10 tobacco (Figure 47). The immunodetection was performed using the enhanced chemiluminescence system (ECL). The immunoblots were scanned and the intensities of signals were estimated using the TINA software package v2.08 beta (Raytest, Spröckhovel; Table 7). No quantitative differences of the proteins tested were detected in Δycf10 tobacco as compared to the wild-type.
RESULTS

Fig. 47. Serological analyses of Δycf10 mutants. Thylakoid proteins from the wild-type and Δycf10 tobacco were separated by SDS-PAGE, electroblotted onto nitrocellulose and incubated with the specific antisera indicated at the right or left of the panels. Two concentrations of the wild-type (WT) were tested to quantify possible differences in protein amounts in Δycf10 mutants.

Table 7. Immunological analyses of the Δycf10 mutants and wild-type tobacco. Signals of intensity identical as in wild-type material are marked (+++). OEC: oxygen evolution complex.

<table>
<thead>
<tr>
<th>Gene</th>
<th>Product</th>
<th>wild-type</th>
<th>Δycf10 mutants</th>
</tr>
</thead>
<tbody>
<tr>
<td>psbP</td>
<td>23 kDa protein of OEC</td>
<td>+++</td>
<td>+++</td>
</tr>
<tr>
<td>psbA</td>
<td>D1 protein of PSII core</td>
<td>+++</td>
<td>+++</td>
</tr>
<tr>
<td>psbD</td>
<td>CP43</td>
<td>+++</td>
<td>+++</td>
</tr>
<tr>
<td>psaA</td>
<td>P700 chl-α protein 1 of PSI core</td>
<td>+++</td>
<td>+++</td>
</tr>
<tr>
<td>psaB</td>
<td>P700 chl-α protein 2 of PSI core</td>
<td>+++</td>
<td>+++</td>
</tr>
<tr>
<td>psaD</td>
<td>ferredoxin-binding protein of PSI</td>
<td>+++</td>
<td>+++</td>
</tr>
<tr>
<td>Lhcb4</td>
<td>CP29</td>
<td>+++</td>
<td>+++</td>
</tr>
<tr>
<td>petA</td>
<td>cytochrome f</td>
<td>+++</td>
<td>+++</td>
</tr>
<tr>
<td>atpA</td>
<td>α-subunit of ATPase complex</td>
<td>+++</td>
<td>+++</td>
</tr>
<tr>
<td>rbsL</td>
<td>large subunit of RuBisCo</td>
<td>+++</td>
<td>+++</td>
</tr>
<tr>
<td>Phosphothreonine</td>
<td></td>
<td>+++</td>
<td>+++</td>
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</tbody>
</table>
3.2.11 Chlorophyll fluorescence analysis

Chlorophyll fluorescence measurements under continuous light (Kautsky effect in continuous light, measurement performed using a FluorCam device) provides a fast test for the photosynthetic performance of plants in vivo. Wild-type and Δycf10 tobacco plants of two distinct phenotypes, namely young green leaves and older leaves, displaying bleached sections (see Section 3.2.7), were analyzed. The young green leaves of Δycf10 plant showed a similar fluorescence induction as the wild-type, while the isolated white tissue regions showed a lack of photosynthetic activity (Figure 48).

![Chlorophyll fluorescence in continuous light](image)

**Fig. 48.** Chlorophyll fluorescence under continuous light (Kautsky effect) of wild-type and Δycf10 mutant. The graph presents data proven by at least 5 independent experiments. Legend: wild-type, Δycf10 green leaf (G), Δycf10 white leaf regions (W). All plants were grown in vitro at 25°C under 100 µE m⁻² s⁻¹. Presented data are not normalized for chlorophyll amount.
For a more detailed spectroscopic analysis of the chlorophyll fluorescence using a PAM fluorometer (Walz, Germany), plants of wild-type and Δycf10 tobacco were grown under 25°C and 100 µE m⁻² s⁻¹. The maximum fluorescence (F_M) and the ratio (F_M-F₀)/F_M = F_V/F_M, as well as the fluorescence quenching parameters q_P (photochemical quenching) = (F_M'-F_S)/(F_M'-F₀'), and q_N (non-photochemical quenching) = 1- (F_M'-F₀')/(F_M'-F₀) (Schreiber 1986) were determined for all samples. The average results of at least three measurements are presented in Table 8.

**Table 8.** Spectroscopic analyses of wild-type and Δycf10 mutant plants.

<table>
<thead>
<tr>
<th></th>
<th>WT 25°C, 100 µE m⁻² s⁻¹</th>
<th>Δycf10 25°C, 100 µE m⁻² s⁻¹</th>
</tr>
</thead>
<tbody>
<tr>
<td>F_V/F_M</td>
<td>0.80</td>
<td>0.80</td>
</tr>
<tr>
<td>q_P</td>
<td>0.90</td>
<td>0.95</td>
</tr>
<tr>
<td>q_N</td>
<td>0.20</td>
<td>0.20</td>
</tr>
</tbody>
</table>

As presented above, the ratio F_V/F_M (reflecting the photosynthetic activity of a plant), and the fluorescence quenching parameters were identical for wild-type and Δycf10 plants, indicating an unaltered photosynthetic activity of Δycf10 tobacco.

**3.2.12 Cᵢ-dependent O₂ evolution**

Figure 49 shows the estimated O₂ evolution as a function of the external Cᵢ concentration in protoplasts and chloroplasts of wild-type and ycf10-deficient tobacco. In each case, 1 ml of protoplast or chloroplast suspension with a chlorophyll concentration between 0.03 and 0.07 mg/ml was measured with an increase of the external HCO₃⁻ concentration from 0 to 5 mM. Since the isolated intact tobacco chloroplasts remained stable for approximately 20 minutes under the chosen experimental conditions, only two measurement points, for 0 and 5 mM HCO₃⁻, were taken. The graphs represent an average of at least 5 independent experiments.
Fig. 49. Substrate dependence of O₂ evolution in wild-type and Δycf10 plants. For the measurements, the isolated intact chloroplasts or protoplasts were resuspended in isolation medium or PCN medium, respectively (see Methods), with a chlorophyll concentration that ranged between 0.03 - 0.07 mg/ml.

An efficient C₁-dependent oxygen evolution was observed with both isolated protoplasts and chloroplasts from wild-type, and, to a lesser degree, from Δycf10 plants. Both wild-type and Δycf10 cells obtained a maximum O₂ evolution with an external C₁ concentration of about 2 mM, although the maximum photosynthetic rate of the mutant was 40 - 50% lower than in the wild-type.
3.3 Pulsed-field electrophoresis as a new method for the determination of the homoplastomic status of transformed plants

3.3.1 DNA separation in pulsed-field agarose gel electrophoresis

Pulsed-field gel electrophoresis (described 1984 by Schwartz and Cantor) allows to separate DNA fragments with the lengths of up to 10 Mbp. Changes of the direction of an electrical field cause a continuous reorientation of DNA molecules, which results in their separation. A schematic picture of the pulsed-changed electrical field electrodes is presented in Figure 50. As presented below, the method of pulsed-field gel electrophoresis served as an efficient tool for the isolation of uncontaminated plastid DNA molecules, and was sucessfully applied for the investigation of the homoplastomic state of transformed plants.

Fig. 50. A schematic drawing of the pulsed-field gel electrophoresis system used. The gel is represented by the white square, the main electrode as a grey hexagon, the electrodes of the pulsed-changed electrical field: squares represent (E/W) direction, circles (S/N) direction.
3.3.2 Isolation of monomeric plastid DNA from the agarose gel

Plastid samples from wild-type and Δycf10 plants and pulsed-field gel electrophoresis were performed as described in Methods. After the electrophoresis, the agarose gels were stained with ethidium bromide and the results visualized under UV light. Several distinct DNA bands were detected, corresponding to the different oligomerization states of the plastid DNA. The monomeric DNA bands were excised from the gel and the DNA was eluted from agarose using the QIAEX kit. The extracted DNA was further used for PCR analysis.

![Fig. 51. Purified intact plastids were immobilized in low-melting agarose blocks and their DNA was separated in a pulsed electrical field. Different forms of plastid DNA are visible: (from the bottom to the top) monomer (1), dimer (2), trimer (3), tetramer (4). Molecular weights in kbp are indicated at the right.](image)

3.3.3 Investigation of the purity of plastid DNA isolated by pulsed-field gel electrophoresis

3.3.3.1 Presence of contaminating nuclear DNA in different plant DNA preparations

The presence of nuclear DNA contamination was investigated with NEP (nuclear-encoded RNA polymerase)-specific primers in a PCR reaction with total plant DNA, DNA isolated from Percoll gradient-purified intact chloroplasts, and plastid DNA isolated by pulsed-field electrophoresis as templates. DNA preparations from wild-type and from Δycf10 tobacco were used in these analyses. As shown by Figure 52A, PCR products derived from nuclear
DNA were detected in both total and DNA from Percoll-purified plastids, but no such signal was detected when the pulsed-field electrophoresis isolated plastid DNA was used as PCR template.

To reinforce and improve this result, the PCR products were blotted onto a nylon membrane and hybridized with a radiolabeled, nested oligonucleotide specific for the NEP primer-amplified DNA fragment. No amplification product was detected in the sample containing the pulsed-field isolated plastid DNA (Figure 52B).

**Fig. 52.** Analysis of the purity of different DNA preparations from wild-type and Δycf10 tobacco. The NEP-specific primer pair was used in a PCR reaction to amplify the DNA fragment originating from nuclear DNA contamination; (t): total DNA, (pt): DNA isolated from Percoll gradient purified plastids, (pf): plastid DNA purified by pulsed-field gel electrophoresis. Molecular weights of products in kbp are indicated at the left. The amplification products were separated electrophoretically and visualized by ethidium bromide staining (Panel A), and hybridization with a radiolabeled NEP-specific oligonucleotide (Panel B).

### 3.3.3.2 Presence of contaminating mitochondrial DNA in different plant DNA preparations

To investigate the presence of mitochondrial DNA contamination in different plant DNA preparations an analogous procedure as described in Section 3.3.3.1.1 was applied, with mitochondrial coxII gene-specific primers.
RESULTS

3.3.4 Analysis of the Δycf10 homoplastomic state by pulsed-field gel electrophoresis

The homoplastomic state of the Δycf10 mutants was examined by a PCR reaction with ycf10-specific primers using total DNA, DNA isolated from the Percoll-purified intact plastids and plastid DNA isolated by pulsed-field gel electrophoresis as a template. Because the pulsed-field electrophoresis isolated plastid DNA is virtually free of promiscuous DNA, any wild-
type signal amplified from \textit{ycf10} DNA will, with high probability, be derived from wild-type plastome copies still present in the transformed cells. The results of the PCR (Figure 54A) showed the presence of the wild-type size amplification product in samples where total DNA and DNA from Percoll gradient-purified plastids served as templates. No such amplification product was detected in the pulsed-field isolated DNA sample by staining of the gel with ethidium bromide.

To improve this result, radioactive probe hybridization was applied. The gel presented in Figure 54A was blotted and hybridized with a radiolabeled, \textit{ycf10}-specific oligonucleotide (Figure 54B). No visible PCR products of the wild-type plastome were detected in the samples with \(\Delta\textit{ycf10}\) plastid DNA purified by pulsed-field electrophoresis.

\textbf{Fig. 54.} PCR analysis of wild-type and \(\Delta\textit{ycf10}\) tobacco. The PCR reaction was performed using \textit{ycf10}-specific primers and different DNA preparations as templates; (t): total DNA, (pt): DNA isolated from Percoll gradient-purified plastids, (pf): plastid DNA isolated by pulsed-field gel electrophoresis. The molecular weights of products in kbp are indicated at the left. Amplification products were separated electrophoretically and visualized by ethidium bromide staining (Panel A), and hybridization with a radiolabeled \textit{ycf10}-specific oligonucleotide (Panel B).
3.3.5 Quantitative estimation of wild-type chromosome copies detected by PCR analysis

To analyze the number of wild-type chromosome copies detectable by PCR analysis of heteroplastomic material, PCR with plasmid DNA templates containing an insertion of ycf10 (wild-type) or the ycf10aadA (mutant) gene was performed. Subsequently, a decreasing amount of wild-type template was mixed with the ycf10aadA plasmid DNA to estimate the ratio of wild-type/mutant DNA copies resulting in a detectable wild-type signal. For PCR analysis $5 \times 10^9$ wild-type and mutant plasmid DNA molecules served as templates (100%). The PCR products were separated electrophoretically and visualized by ethidium bromide staining, subsequently blotted onto a nylon membrane and hybridized with a radiolabeled, nested, ycf10-specific oligonucleotide (Figure 55, Panel A and B, respectively).

![Fig. 55. PCR analysis of plasmid DNA containing ycf10 (wt) or the ycf10aadA (mutant) gene insertion. Lanes: (0) zero control, (1) wt DNA template (100%; $5 \times 10^9$ plasmid DNA molecules), (2) mutant DNA template (100%; $5 \times 10^9$ plasmid DNA molecules), (3) mixed templates: mutant/wt = 1/1, (4) 1/0.1, (5) 1/10$^{-2}$, (6) 1/10$^{-3}$, (7) 1/10$^{-4}$, (8) 1/10$^{-5}$, (9) 1/10$^{-6}$, (10) 1/10$^{-7}$. The molecular weights of the products are indicated at the left. The amplification products were separated electrophoretically, visualized by ethidium bromide staining (Panel A), and hybridized with a radiolabeled ycf10-specific oligonucleotide (Panel B).]

3.3.6 Sensitivity level of the PCR

The detection level of the PCR analysis under the chosen conditions (described in Methods) was tested using a decreasing amount of plasmid DNA template. Both the ethidium bromide
staining and the hybridization with radioactive ycf10 probe detected a PCR product when a template containing 50 plasmid DNA molecules was used.

**Fig. 56.** PCR analysis of plasmid DNA containing the ycf10 gene insertion. Lanes: (0) zero control, (1) $5 \times 10^9$ plasmid DNA molecules serving as a template, (2) $5 \times 10^8$ DNA molecules, (3) $5 \times 10^7$ DNA molecules, (4) $5 \times 10^6$ DNA molecules, (5) $5 \times 10^5$ DNA molecules, (6) $5 \times 10^4$ DNA molecules, (7) $5 \times 10^3$ DNA molecules, (8) $5 \times 10^2$ DNA molecules, (9) 50 DNA molecules, (10) 5 DNA molecules. Molecular weights of the products are indicated at the left. The amplification products were separated electrophoretically and visualized by ethidium bromide staining (Panel A), and hybridization with a radiolabeled ycf10-specific oligonucleotide (Panel B).
4. DISCUSSION

The existence of a unique genetic system of plastids has first been suggested at the beginning of the 20th century. From analyses of variegated plants Correns (1909) and Baur (1909) deduced independently the hypothesis of plastid inheritance that did not obey Mendelian rules. Erwin Baur concluded in 1909 that (i) the plastids are carriers of hereditary factors which are able to mutate, (ii) in variegated plants the random process of plastid sorting-out takes place, and (iii) genetic plastid factors are inherited biparentally. In contrast, Carl Correns (1909), using a different model, described maternal inheritance of factors localized in the cytoplasm (summarized in Hagemann 2000). Only fifty years later, the specific cytochemical staining of plastid-localized DNA in the green algae Chlamydomonas moewussii (Ris and Plaut 1962), as well as the isolation and description of a unique DNA species of Chlamydomonas reinhardtii, enriched in chloroplasts (Sager and Ishida 1963) proved Baur’s hypothesis. The subsequent isolation and microscopic characterization of chloroplast DNA from higher plants (Wells and Birnstiel 1969, Herrmann et al. 1975, Kolonder and Tewari 1975) provided an insight into the structure, conformation and coding capacity of plastid DNA. The development of techniques for identification and isolation of genes as well as sequence analyses of entire plastid chromosomes (Ohyama et al. 1986, Shinozaki et al. 1986) allowed to assess the coding potential of plastid chromosomes. Recently, the development of chloroplast transformation methods first established in Chlamydomonas reinhardtii (Boynton et al. 1988) and then in tobacco (Svab et al. 1990, Golds et al. 1993) as well as the availability of additional techniques for analysis, such as protein sequencing or two-dimensional gel electrophoresis, allowed to identify and characterize most of the plastid-encoded genes and their products. Nevertheless, higher plant plastomes still comprise several open reading frames (ORFs) of unknown function. Conserved ORFs, which are found in almost all known plastid chromosomes are designated ycf$s$, and at present are subject of intense molecular, biochemical and biophysical analyses.

The work presented in this thesis has concentrated on the functional characterization of two ycf$s$ of unknown function, namely ycf9 (psbZ, ORF62), and ycf10 (cemA, ORF229). With respect to chloroplast transformation, a PFGE-based plastid DNA isolation protocol was established which allowed to verify the status of homoplastomy of transformed plants.
4.1 Functional analysis of the ycf9 gene product

A reverse genetic approach was applied to analyze the function of the conserved ycf9 gene in *Nicotiana tabacum* and, in cooperation with the group of Prof. F.-A. Wollman (Université Paris, France), in *Chlamydomonas reinhardtii*.

**Inactivation of the ycf9 gene in Nicotiana tabacum, assessment of the homoplastomic status.** The ycf9 gene was an interesting case for the reverse genetic approach not only because of its enigmatic function, but also because parallel attempts to inactivate the gene had yielded controversial results. Disruption of ycf9 in *Chlamydomonas* generated a mutant line displaying a wild-type phenotype (Higgs et al. 1998). A more recently described ycf9 knock-out mutant in tobacco did not reach the homoplastomic state and the gene product was assigned to the PSI complex (Mäenpää et al. 2000). Studies of Ruf et al. (2000), in turn, had demonstrated homoplastomic Δycf9 mutants, and the gene product was proposed to be a member of a small antenna protein family. The biolistic transformation performed in our laboratory yielded four independent mutant lines, which, after six cycles of regeneration, reached homoplastomy. The homoplastomic state of all transformed lines was established by sequence analysis, PCR, Southern and Northern analyses, as well as by a combination of pulsed-field analysis and PCR of relevant regions of the DNA band containing intact plastid chromosomes. Northern and Southern analyses demonstrated the absence of wild-type transcripts and DNA copies, respectively, in Δycf9 plants, although wild-type signals were detected in PCR analyses. The pulsed field electrophoresis approach followed by PCR of relevant chromosomal regions and hybridization with radiolabeled probes, developed in this study, demonstrated that those wild-type signals arose, in all likelihood, from promiscuous DNA, i.e. copies of plastid DNA residing in nucleus or chondriome, and proved homoplastomy at a level lower than 0.00001%.

The estimation of the homoplastomic state is a crucial, but also a controversial issue in the analysis of plants with transformed plastids. Due to about 100 chloroplasts per cell and approximately 100 chromosome copies per chloroplast (Herrmann and Possingham 1980), the primary transformant is heteroplastomic, containing both wild-type and mutant plastid chromosomes. The homoplastomic state, in which all wild-type copies are replaced by mutant ones, is generally achieved after several rounds of regeneration, but plants can be “fixed” in the heteroplastomic state, if the transformed gene is essential for vital functions (Drescher et
al. 2000). Multiple copies of chloroplast DNA per cell, as well as the existence of promiscuous plastid DNA fragments in both the nuclear genome and the chondriome (Ellis 1982, Stern and Lonsdale 1982, Ayliffe and Timmis 1992, Brennicke et al. 1993) complicate the analysis of the homoplastomic state of transformed plants.

Generally, two strategies are applied to verify the presence of residual wild-type plastid DNA in transplastomic plants: (i) Southern and Northern analysis of plastid DNA and RNA, respectively, and (ii) PCR using sequence specific oligonucleotides flanking the insertion. Southern and Northern analyses give general information about the presence of the wild-type DNA copies (RFLP) or wild-type transcripts, respectively, but both methods lack sensitivity to unambiguously determine the homoplastomic status of transformed plants. On the other hand, the PCR approach is more sensitive and allows the detection of minute quantities of wild-type DNA copies undetectable by Southern analysis, but it bears the risk of (co)amplifying promiscuous plastid DNA sequences residing in the nucleus or mitochondrial genome. This risk can be reduced, but not eliminated, by using DNA extracted from intact plastids purified by isopycnic, isotonic gradient centrifugation (Schmitt and Herrmann 1977, Herrmann 1982) as a template for amplification. However, as a consequence of the mechanical disruption of plant material prior to gradient centrifugation, and/or insufficient phase separation, traces of nuclear and/or mitochondrial DNA generally copurify with intact chloroplasts, thus mimicking genuine plastid chromosomes in the subsequent PCR reaction.

To ensure the absence of wild-type plastid chromosome copies left in a transplastomic plant, an additional, more reliable method is required. One alternative to judge the homoplastomic state of fertile mutant plants is to screen the progeny for the presence of a marker gene by germinating them on selective medium (Maliga and Nixon 1998). In this thesis, pulsed-field gel electrophoresis (PFGE) was applied to obtain seemingly contamination-free plastid DNA from transplastomic mutants. Without the PFGE purification step the Percoll-gradient purified plastid DNA yielded a 200 bp amplification product characteristic for wild-type chloroplast chromosomes in the subsequent PCR. Its wild-type origin was unambiguously proven by the use of plastid-, mitochondria- and nucleus-specific oligonucleotides, and DNA templates derived from different extraction methods, including PFGE. In wild-type plants, the 200 bp amplification product was obtained with total DNA isolated from leaves as well as with plastid DNA extracted from intact, Percoll-purified chloroplasts before and after PFGE. The insertion of the marker cassette into the mutant plants, in turn, was detected in three
subfractions of DNA from transplastomic plants. Wild-type-derived amplification products were obtained, whenever DNA extracted from leaves and Percoll-purified chloroplasts served as a template. In striking contrast, no such signal was detected, if a PFGE step was included into the DNA extraction procedure. The finding that neither mitochondrial nor nuclear sequences were amplified with gene-specific oligonucleotides using PFGE-purified plastid DNA as a template demonstrates the high purity of the extracted DNA, free of any cross-contamination with mitochondrial and/or nuclear DNA. In contrast, both mitochondrial and nuclear DNA fragments were amplified from DNA extracted from leaves or even Percoll gradient-purified plastids. The sensitivity of the PCR was estimated with a plasmid DNA containing either the wild-type gene (ycf10) or the gene with insertion of the aadA cassette. With a ratio of one wild-type gene copy to 10,000 transformed gene copies (1/10^4) the wild-type derived 200 bp amplification product was clearly detectable in ethidium bromide-stained agarose gels. The sensitivity of detection was increased by transfer of amplified products onto nylon membrane and hybridization with a radiolabeled probe. Taking into consideration that in a fully developed tobacco leaf cell, the ratio of nuclear DNA copies (4 haploid copies in the allotetraploid nucleus) to plastid DNA copies (100 per chloroplast and 50 - 100 chloroplasts per cell) is less than 1/10^3, and much stronger signals are generated for the wild-type fragment with the non-pulsed field gel purified fractions than with the 1/10^3 molar WT/mutant mixture, the promiscuous DNA copies most probably do not only originate in the nucleus, but also in mitochondria (several hundreds per cell). To sum up, the absence of a wild-type-derived signal in PFGE-purified plastid DNA isolated from transplastomic plants can either be due to (i) their complete segregation (homoplastomy), or (ii) to few wild-type plastid chromosome copies (less as one wild-type gene per 10,000 transformed chromosomes), which corresponds to less than one wild-type plastid chromosome per plant cell containing 100 chloroplasts with approximately 100 plastid chromosome copies per chloroplast.

**Phenotypes of the Δycf9 plants.** Δycf9 plants grown under different light and temperature regimes differed in their phenotypes. These variations did not arise from the heterotrophic *in vitro* conditions but were as well visible under photoautotrophic growth conditions using the same light and temperature regimes. As recently reported, similar phenotypes were observed in Δycf9 plants grown in the greenhouse under regulated light conditions (Baena-González et al. 2001). To understand how growth conditions and Δycf9 phenotypes are interconnected, detailed biochemical analyses of transformed plants were required.
Localization of the ycf9 gene product. The co-transcription of ycf9 together with the psbD and psbC genes, encoding subunits of the PSII core, suggested its localization in PSII complexes as well. However, the first attempt to localize the ycf9 gene product serologically revealed co-migration with PSI (Mäenpää et al. 2000). Subsequent analyses of Ruf et al. (2000) demonstrated co-purification of the Ycf9 protein with the LHCII complexes. Analyses of tobacco and Chlamydomonas reinhardtii demonstrated compellingly that the Ycf9 protein is a new subunit of the PSII core complex (Swiatek et al. 2001). By eliciting specific antibodies against the Chlamydomonas and tobacco proteins it was shown that in both organisms the accumulation of Ycf9 is strictly correlated with the other PSII core subunits (Swiatek et al. 2001). Furthermore, recent analyses of plants lacking the PSII core proteins PsbE and PsbF, but with an unaltered level of minor antennae or LHCII (Swiatek et al. manuscript in preparation), demonstrated that those plants also failed to accumulate the Ycf9 protein, implying its close relationship and dependence on other PSII compounds. In Chlamydomonas, the protein co-purified with PSII after sucrose gradient centrifugation of detergent-solubilized thylakoid membranes. Interestingly, in tobacco silver staining of solubilized thylakoid membranes revealed two protein components of 4 and 6 kDa detectable in wild-type, but lacking in mutant fractions. The 4 kDa component co-migrated clearly with PSII complexes, while the second protein of approximately 6 kDa, which was also missing in Δycf9, resided in the upper region of the sucrose gradient containing carotenoids and free protein fraction, and its distribution profile did not match that of LHCII complexes, as suggested by Ruf et al. (2000). Isolation of thylakoid protein complexes by native Deriphat-PAGE, followed by denaturing SDS-PAGE revealed the existence of the minor antenna protein complex (CP24, CP26 and CP29) localized above the trimeric LHCII band. The sucrose-gradient centrifugation of β-dodecylmaltoside-solubilized thylakoid membranes demonstrated comigration of the minor antennae complex with the monomeric LHCII complex, which was not resolved using the Deriphat-PAGE technique. The 6 kDa protein comigrated with those complexes, although its maximal amount was detected above, in fractions enriched in carotenoids. It has been described that, on denaturing polyacrylamide gels, strongly hydrophobic proteins can migrate faster than expected from their molecular weight. On the basis of SDS-PAGE with total thylakoid proteins, the deficiency of only one protein of approximately 4 kDa was observed in Δycf9 plants. The size of this component is identical to the protein immunodetected by Ycf9 antibodies. These results strongly suggest that the lack of the 4 kDa protein in PSII complexes of transformed plants is due to the disruption of the ycf9 gene. Since the Ycf9 is localized in PSII cores, it was designated PsbZ
(Swiatek et al. 2001). The 6 kDa component, missing in Δycf9 tobacco, could represent the PsbZ precursor form unassociated with PSII, or an unrelated protein which is unstable in the absence of PsbZ. It remains unclear, why only one protein deficiency was detected on basis of total thylakoid protein SDS-PAGE. To answer this question an improved gel system for resolution of small proteins in overloaded gels has to be developed. Recent analyses of several transformed plants lacking small proteins of PSII (Swiatek et al., manuscript in preparation) demonstrated the presence of numerous proteins in the range of 6 - 8 kDa, which might not be individually separated even using the Schägger/von Jagow gel system, optimized specifically for the resolution of small proteins.

Sequence analyses of PsbZ from various species favour a putative cleavage site for a peptidase directly following the first of the two transmembrane helices. Although Ruf et al. (2000) have shown a virtual co-purification of this protein with LHCII complexes, the refinement of the separation techniques of thylakoid protein complexes in sucrose density gradients as well as in 2D gels clearly demonstrated its localization in the carotenoid/free protein fraction and thus argues against its co-migration with LHCII.

In recent years, the application of advanced electron microscopy and single-particle image techniques solved the PSII structure and, based on biochemical work, allowed to divide PSII into various subcomplexes, including PSII reaction centers (RCII), PSII core complexes (CCII), which contain the core antenna proteins CP47 and CP43 together with the reaction center and other subunits, and several antenna complexes whose association strengthens the functional PSII unit (Green and Durnford 1996, Barber et al. 1997, Wollman et al. 1999, Barber and Kühlbrandt 1999, Nield et al. 2000, Zouni et al. 2001). Although RCII and CCII have been purified and characterized, the function of most of the subunits in the formation of a functional PSII unit remains poorly understood. During the past decade, several small PSII subunits have been identified, generally by reverse genetic approaches. Although these subunits have been tentatively attributed to functions, their physiological importance is still elusive (Wollman et al. 1999). Recent studies of the supramolecular organization of PSII, based on mild detergent solubilization procedures and refined electron microscopic analysis, including single particle averaging (Boekema et al. 1999a), cryoelectron microscopy (Nield et al. 2000), and two-dimensional crystal electron diffraction (Barber and Kühlbrandt 1999, Hankamer et al. 1999, Rhee et al. 1998), have shown various PSII forms ranging from a large protein complex retaining its peripheral antenna complement to subcore particles.
Furthermore, the three-dimensional structure of a cyanobacterial PSII core preparation yielded molecular insight into the positions of the major PSII transmembrane helices (Zouni et al. 2001). Studies of the supramolecular organization of PSII revealed existence of minimum 36 transmembrane helices (Barber and Kühlbrandt 1999, Hankamer et al. 1999, Zouni et al. 2001) from which 24 have been attributed to D1, D2, CP43, CP47, PsbE, and PsbF. Several small subunits, including PsbI, PsbL, PsbH, PsbK and PsbX have been tentatively assigned in PSII core, and seven helices could not be correlated with known subunits. Of particular interest is the structural domain within a PSII monomer where D1 stands in close contact to CP43, which, in turn, is suggested to associate with CP26. Adjacent to D1 and CP43 are several transmembrane helices of unknown origin (Nield et al. 2000, Zouni et al. 2001). PsbZ is regarded as a potential candidate for providing one or two of these transmembrane helices, which are in close contact to D1, CP43 and CP26. As will be discussed below, the interaction of PsbZ with the PSII core and peripheral antennae strongly supports this localization.

**Fig. 57.** Molecular reconstruction of the PSII-LHCII supercomplex showing the putative localization of PsbZ. The presentation of PSII and LHCII proteins are based on data published by Zouni et al. (2001) and Kühlbrandt et al. (1994), respectively. The location and orientation of the peripheral antenna complexes with respect to the PSII core are based on Harrer et al. (1998), Boekema et al. (1999b), and Nield et al. (2000). Protein moieties are shown by space-filling models of the C-α carbon atoms, and chlorophylls are shown as dark green frame models. The helices belonging to the PSII core that have no attribution by Zouni et al. (2001) are shown in dark yellow. The putative localization of PsbZ is among five helices circled in yellow. Model kindly provided by R. Kuras and R. Bassi.
**The effect of PsbZ deletion on minor antenna protein complexes.** The analyses of the protein pattern of wild-type and ∆ycf9 plants demonstrated the absence of a protein of approximately 25 kDa in mutant material. Subsequent Western analyses indicated that, in fact, the amount of two proteins of similar masses, namely the CP26 and CP29 minor antennae proteins, was reduced in transformed plants. The application of different growth conditions allowed further analyses of minor antenna accumulation in ∆ycf9 plants. In agreement with data from Ruf et al. (2000), mutant plants displayed a reduced amount of CP26 under most growth conditions. The deficiency in CP26 was most pronounced at 20°C in dim light, conditions at which plants exhibited a dwarf phenotype. Under the same conditions the amount of CP29 protein, otherwise comparable to wild-type levels, was decreased by approximately 50%. The amount of the third minor antenna apoprotein, CP24, remained unchanged under all conditions applied. Thus, the absence of PsbZ enhanced the light- and temperature-dependent loss of two minor antenna proteins. CP29, CP26 and CP24, together with LHCII, form the light-harvesting complex of PSII. Detailed electron transfer studies, as well as single-particle image techniques, localized CP26 in close proximity with the CP43 protein of PSII core center (Zouni et al. 2001). As illustrated by Figure 57, PsbZ possibly represents one or two unidentified transmembrane helices identified in this region. This indicates a role of PsbZ in attachment of LHCII complexes *via* CP26 to the PSII core center, and, consequently, in the energy transfer from the LHCII to PSII. This also explains, as will be discussed below, the role of PsbZ in the assembly of PSII supercomplexes and alteration of xanthophyll cycle observed in ∆ycf9 plants. In contrast to CP26, the observed instability of CP29 can not be explained by its direct interaction with PsbZ. CP29 has been shown to reside near the PSII dimer grove, and on the opposite side of PSII monomer as CP26. The impaired stability of CP29 in absence of PsbZ is likely a secondary effect, possibly caused by altered protein phosphorylation observed in ∆ycf9 plants.

**Functional role of PsbZ in the assembly of PSII-LHCII supercomplexes.** As mentioned above, PsbZ is dispensable for photoautotrophic growth and CCII assembly. However, it appears to play a key role in the interaction between CCII and the peripheral antenna complement. When growth conditions were chosen in such a way that limited biochemical defects were detectable in thylakoid membranes of the mutant (heterotrophic and autotrophic growth at 25°C), the stability of PSII supercomplexes, which results from a tight interaction between CCII and peripheral antenna complexes, was severely decreased: these supercomplexes no longer resisted freeze-fracturing (Swiatek et al. 2001) or detergent
solubilization of thylakoid membranes. These observations confirm the idea that PsbZ is a key CCII subunit mediating its interaction with the peripheral antenna complexes.

**Alteration of the protein phosphorylation pattern in tobacco Δycf9 plants.** The interaction between peripheral antennae components and PSII cores is controlled by the phosphorylation of several antennae subunits in an acclimation process known as the state transition (reviewed by Allen 1992). Analyses of thylakoid protein phosphorylation in tobacco and *Chlamydomonas* provided some intriguing insights into the mechanism enabling PsbZ to act as a linker between CCII and peripheral antenna complexes. In Δycf9 mutants the steady state level of CCII phosphorylation relative to LHCII was markedly decreased, as demonstrated by anti-phosphothreonine antibodies and radiolabeling with $^{32}$P. Since an increased phosphorylation of LHCII vs. CCII under state II conditions is accompanied by detachment of peripheral antennae components from PSII (Allen 1992), the reduced stability of PSII-LHCII supercomplexes is not surprising, and very likely induced by the relative increase of LHCII vs. CCII phosphorylation observed in the absence of PsbZ. These results further imply that the content of CP26 may be controlled by the distribution of phosphate groups among the polypeptides that interact in the PSII units. Biochemical analyses of PSII-LHCII subcomplexes indicated an interaction of CP26 with CP43 and LHCII (Harrer et al. 1998), both of which are subject to reversible phosphorylation. Thus, the accumulation of CP26 in PSII units may rely on the presence of its interacting partners in a phosphorylated state, which itself is dramatically reduced at 20°C under dim light in the absence of CP26 in mutant membranes. A putative physiological function of the PSII core subunit phosphorylation as a prerequisite for stable assembly of PSII unit (CCII + chlorophyll a/b binding proteins) has been proposed (de Vitry et al. 1989), and is supported by the observation that BF4, a *Chlamydomonas* mutant devoid of chlorophyll a/b binding antenna proteins as well as of CP26, CP29 and CP24, does not display any PSII core phosphorylation (de Vitry and Wollman 1988). Reversible phosphorylation of D1 and D2 proteins is also part of the PSII repair cycle process. Under stress conditions, dephosphorylation of photodamaged D1 is prerequisite for its degradation (Koivuniemi et al. 1995, Rintamäki et al. 1996, Kruse et al. 1997). It remains unknown if PsbZ could take part in repair cycle processes. The presence of phosphorylatable amino acids residues in PsbZ sequence suggests the possibility of its reversible phosphorylation. The possibility of phosphorylation of PsbZ and its role in regulation of protein turn-over will be an interesting subject for future studies. The work of Baena-González et al. (2001) confirmed an altered thylakoid protein phosphorylation pattern
in Δycf9 plants. These authors observed a similar decrease in the steady state level of CCII phosphorylation, whereas LHCII phosphorylation level contradicted the results presented in this thesis. These differences might be due to a different treatment of the plants prior to the experiment. The studies performed in my thesis relied on plants grown under strictly controlled light and temperature regimes. The transfer of plants to different light conditions induces changes in the phosphorylation state of thylakoid proteins and could therefore result in phosphorylation patterns different from those presented in this thesis.

**Analysis of non-photochemical quenching and xanthophyll cycle in Δycf9 plants.** The PsbZ-dependent interaction of PSII cores with peripheral antenna proteins significantly influences the ability of PSII to regulate the flux of light excitation. Inactivation of the ycf9 gene encoding PsbZ not only reduced the stability of minor antenna proteins, especially CP26, but also impaired adaptation mechanisms in PSII, namely NPQ and the xanthophyll cycle. NPQ protects PSII against photoinhibition by dissipating excess excitation energy as heat (Demmig-Adams 1990, Horton 1996, Ramelli et al. 1999, Croce et al. 1999, Bassi et al. 1999, Verhoeven et al. 1999, Havaux and Niyogi 1999, Ruban et al. 1999, Ruban et al. 2000). The molecular mechanism of NPQ is not yet fully understood. However, its full expression involves de-epoxidation of violaxanthin to zeaxanthin as shown by the marked decrease in NPQ in *Arabidopsis* mutants inhibited in the violaxanthin de-epoxidation process (Niyogi et al. 1998). Furthermore, NPQ seems to require the interaction of xanthophylls with LHCII proteins as indicated by a reduced NPQ in the barley chlorina f2 mutant, lacking Chlorophyll b and functional LHCII complexes (Tardy and Havaux 1997). As mentioned above, the major violaxanthin binding proteins, CP26 and CP29, are located at the interface between the PSII core and the trimeric LHII at the periphery of the PSII complexes (Harrer et al. 1998, Boekema et al. 1999a and 1999b). Minor antenna proteins contain protonatable amino acid residues which might sense a low lumenal pH, thus triggering NPQ (Walters et al. 1994, Walters et al. 1996, Pesaresi et al. 1997, Ruban et al. 1998). The depletion of CP26 can explain the paradoxical behavior of Δycf9 mutants: although CP29 and CP26 are both able to exchange violaxanthin for zeaxanthin, this process is much more rapid with CP26 than with CP29 (Ruban et al. 1994, Croce et al. 1996). At sub-optimal temperatures (17°C and 20°C), the de-epoxidation of violaxanthin to zeaxanthin could be inhibited by the loss of CP26 and a reduced amount of CP29 in Δycf9 mutants, whereas at higher temperatures remaining CP29 and CP26 might be sufficient to maintain this process, thus leading to the recovery of NPQ at 25°C. However, antisense inhibition of the CP26 and CP29 proteins in *Arabidopsis* excluded their direct role in the formation of NPQ (Andersson et al. 2001), even though CP26 and
CP29 antisense plants exhibited variations in quenching, namely a reduction of NPQ at intermediate light levels, especially in CP29 antisense plants (Andersson et al. 2001). Thus, it was proposed that CP26 and CP29 are not directly involved in NPQ, but rather in LHCII antenna coordination. Another possibility is, that either CP29 or CP26 alone is sufficient for NPQ formation. Up to date, no successful generation of CP26/CP29 double mutants was performed to resolve this question. Nevertheless, unaltered levels of other LHCII proteins, as well as a wild-type level of PsbS in Δycf9 plants imply a role of CP26 or/and CP29 in NPQ formation. In addition, the increased amplitude of the quenching process could be due to a much higher content of xanthophyll cycle pigments observed in the mutant as well as an increased zeaxanthin to violaxanthin ratio induced under strong illumination. The dramatic increase of the carotenoid content in mutant plants could, in turn, express their low capacity for excitation energy dissipation under high-light conditions. A similar effect has been observed in Chlorella in response to high-light treatment, which increases the excitation energy pressure on PSII (Maxwell et al. 1995).

The results discussed above are consistent with the dwarf phenotype of Δycf9 plants grown at low temperature, which corresponds to the effect of PSII overexcitation induced by moderate illumination at low temperature in barley (Huner et al. 1998). To sum up, it can be concluded that the inactivation of ycf9 disrupts the PSII-LHCII supercomplex architecture, exerting its effect mainly on the regulation of Chl a-excited states by mainly destabilizing the CP26 subunit, and to a lesser extent CP29, under certain growth conditions. The decreased capacity for energy dissipation leads to a “bleached” phenotype and an over-stimulation of xanthophyll synthesis, required for both the recovery of NPQ and for scavenging of reactive oxygen species in the photosynthetic membrane (Havaux and Niyogi 1999).

An additional influence of PsbZ on carotenoid biosynthesis was reported recently (Baena-Gonzáles et al. 2001). In this work, a reduced amount of the plastid terminal oxidase (PTOX), a key enzyme for carotenoid biosynthesis, has been demonstrated in Δycf9 tobacco plants. PTOX desaturates phytoene, a C40 acyclic precursor, which subsequently leads to the formation of lycopene. Carotenoid biosynthesis itself is redox-regulated and requires quinones as redox mediators. The reduction of plastoquinone is led through the NDH complex at the expense of oxygen in a process known as chlororespiration (Bennoun 1982, Carol et al. 1999, Carol and Kuntz 2001). Thus, in Δycf9 plants, de-regulation of the carotenoid
biosynthesis might be a consequence of the altered ability of PSII to regulate the flux of light excitation.

The functional analyses of Δycf9 tobacco plants presented in this thesis indicate a critical role of PsbZ in NPQ, which previously has been attributed mainly to PsbS and CP29 (Crofts and Yerkes 1994, Pesaresi et al. 1997, Li et al. 2000). Although the content of CP29 varied as well in mutant plants under certain growth conditions, these variations did not correlate convincingly with the amplitude of the major NPQ changes. A direct role of CP29 in the regulation of NPQ-related processes could be as well excluded from the analyses of CP29 antisense Arabidopsis plants (Andersson et al. 2001). Furthermore, the accumulation of PsbS in Δycf9 plants was unaltered. Interestingly, PsbZ is present in phycobilisome-containing organisms which contain neither PsbS, CP29 nor the xanthophyll cycle. Nonetheless, these organisms display NPQ upon exposure to high light distinct from state transitions or photoinhibition (Campbell et al. 1998, El Bissati et al. 2000). It is therefore tempting to speculate that PsbZ is the key player in the formation of NPQ which remained in phycobilisome-containing photosynthetic organisms, although the altered, but still functioning NPQ observed in a Synechocystis PCC6803 ycf9 knock-out strain (S. V. Shestakov and R. Bassi, personal communication) suggested an involvement of additional factors, unknown so far, in the regulation of this process. The PsbZ-CP43 interaction, which was based on its influence on CP26 stability and CP43 phosphorylation, places PsbZ at a position where it could interact with the PSII cores to elicit an adaptation response that dissipates excess light excitation via a molecular mechanism that remains to be investigated. In higher plants, the xanthophyll cycle and NPQ, together with other mechanisms, namely alternate electron-donation pathways, are involved in the regulation of excitation energy in PSII. PSII contains two accessory chlorophylls (ChlZ ligated to D1-His318, and ChlD ligated to D2-His117), carotenoid and cytochrome b559 cofactors that function as alternative electron donors under physiological conditions when the primary electron-donation pathway from the O2-evolving complex to P680+ is inhibited. Oxidized ChlZ is known as a potent quencher of chlorophyll fluorescence that may act together with Cyt b559 in the dissipation of excess excitation energy under high light conditions (Tracewell et al. 2001). ChlZ and ChlD are supposed to be bound near small PSII subunits associated with the D1 and D2 proteins. The proposed localization of the PsbZ protein in the proximity of D1 and CP43 implies its regulative role in an adaptation response that can lead to the dissipation of excess light energy in more than one way.
The role of PsbZ in the photosynthetic electron transport. The localization of PsbZ in PSII complexes as well as the discovery of PsbZ-dependent PSII-LHCII interactions reflected its possible influence on the photosynthetic electron transport. It has been published that inactivation of \textit{ycf9} in tobacco leads to an accelerated electron flux to PSI without a simultaneous change in the maximum electron transfer capacity of PSII (Baena-González et al. 2001). These alterations did not influence carbon fixation in the Calvin-Benson cycle, neither the recycling of electrons from ferredoxin back to PQ (ferredoxin-dependent cyclic electron transport) nor back to plastoquinon via the NDH complex. The NDH complex reduces PQ using stromal NAD(P)H as a substrate, and it has been supposed that it contributes to both the electron flow around PSI in the light and in a putative respiratory chain within the chloroplast (Bennoun 1982, Carol et al. 1999, Cournac et al. 2000, Carol and Kuntz 2001). Baena-González et al. (2001) demonstrated an increased ascorbate peroxidase activity in $\Delta$ycf9 plants, which suggests an increased Mehler reaction, in which superoxide is reduced by superoxide dismutase and ascorbate peroxidase (Asada 1999). The increased amount of superoxide could be the effect of an electron leak resulting from impaired electron transfer from PSII to PSI. In this context, the role of PsbZ in the fine-tuning of the photosynthetic reaction, optimization of the distribution of excitation energy between PSII and PSI, and the protection of the two photosystems against photodamage is of particular interest. As discussed above, the results of non-photochemical quenching (NPQ) and the xanthophyll cycle in $\Delta$ycf9 mutants presented in this thesis correspond to these suggestions.

4.2 Functional analysis of the \textit{ycf10} gene product

Similarly to \textit{ycf9} described above, the \textit{ycf10} open reading frame belongs to the few plastid encoded genes, whose function is still a matter of debate. The putative gene product, first designated HBP (heme-binding-protein; Willey and Gray 1990), was renamed CemA (chloroplast envelope membrane protein A) due to its immunolocalization in the inner envelope membrane of pea chloroplasts (Sasaki et al. 1993). The topology of the protein localization remains unclear because of the uncertainty about the number of membrane-spanning regions in the polypeptide. In their work, Willey and Gray (1990) indicated a putative cleavage site between residues Ser-24 and Phe-25. Cleavage at this side would result in a 24-amino acid signal sequence, corresponding to the first hydrophobic region of Ycf10. However, further analyses of Ycf10 in pea (Sasaki et al. 1993) did not support this hypothesis.
The cyanobacterial \textit{cotA} gene product, which displays high sequence homology to the \textit{ycf10}-encoded polypeptide of higher plants has been shown to be involved in the light-induced proton extrusion (Katoh et al. 1996ab), whereas the inactivation of \textit{ycf10} in \textit{Chlamydomonas} suggested its role in inorganic carbon uptake into chloroplasts (Rolland et al. 1997).

**Inactivation of the \textit{ycf10} gene in \textit{Nicotiana tabacum}**. The reverse genetic approach was applied to analyze the function of the conserved gene in \textit{Nicotiana tabacum}. To this day, no successful inactivation of the \textit{ycf10} reading frame in higher plants was reported, despite several attempts (P. Maliga and J.-D. Rochaix, personal communication). In our case, the biolistic chloroplast transformation yielded three independent mutant lines, which reached homoplastomy after seven cycles of regeneration. The homoplastomic state of the transformed plants, indicated by the absence of wild-type derived DNA copies, was ultimately confirmed by PFGE, PCR and Southern analyses. Results, similar to those obtained for \textit{Δycf9} plants, demonstrated that a wild-type specific amplification product, detectable when total or Percoll-gradient-purified DNA served as a template, arose from promiscuous DNA copies residing in the nucleus or/and in the chondriome, and proved homoplastomy at a level lower as 0.00001%. Northern analysis further supported the homoplastomy of mutants. In addition, co-transcription of \textit{ycf10} with the upstream located \textit{psaI} and \textit{ycf4} genes and the downstream \textit{petA} gene, as well as the existence of at least two promoters were deduced from the transcription map of the \textit{psaI-ycf4-ycf10-petA} operon, displaying similarities to the transcription pattern observed in pea (Willey and Gray 1990; Nagano et al. 1991). The segregation of homoplastomic \textit{Δycf10} plants strongly indicates that the \textit{Δycf10} open reading frame is not required for cell viability, supporting the results observed in the \textit{Chlamydomonas} \textit{ycf10} inactivation mutant (Rolland et al. 1997).

**Phenotypes of the \textit{Δycf10} plants**. Transplastomic \textit{Δycf10} plants were able to grow photoautotrophically in the greenhouse, and did not show any phenotypic alterations upon the \textit{in vivo} cultivation. White tissue regions, characteristic for \textit{Δycf10} plants cultivated \textit{in vitro}, were observed as well in \textit{Δycf10} seedlings grown on medium without antibiotic, thus excluding a spectinomycin effect on the \textit{Δycf10} phenotype. The analysis of the \textit{Δycf10} \textit{Clamydomonas} mutants indicated an increased light sensitivity of the transformed cells, resulting in bleached mutant cultures (Rolland et al. 1997). The bleaching of certain tissue regions in tobacco mutants could be a light-dependent effect as well, however, no such
phenotype was observed when plants were grown in dim light (10 μE m⁻² s⁻¹, 25°C, data not shown).

**Photosynthetic activity of the Δycf10 plants.** In all analyzed plant species so far, the ycf10 reading frame is part of a gene cluster encoding essential photosynthetic genes, for example psaI and petA, suggesting a possible physiological function of the product in the photosynthetic processes. However, the chlorophyll fluorescence spectra of three Δycf10 mutants generated independently were undistinguishable from wild-type plants, thus indicating that photosystems I and II, as well as the cytochrome b₆f complex were not significantly affected. This supports the data of Rolland et al. (1997) reported for the Δycf10 mutant of *C. reinhardtii*. Chlorophyll fluorescence measurements of the Δycf10 mutant plants using the FluorCam device revealed a small decrease in the maximal fluorescence under continuous illumination (F_P) relative to the wild-type, indicating slightly impaired reduction of PSII. However, these observations were not supported by more detailed and standardized analyses of PSII activity using the PAM device. Nevertheless, all fluorescence analyses have shown an absence of photosynthetic activity in bleached tissue regions. This result was supported by electron micrographs of Δycf10 mutant cells, showing structured plastids in the green tissue regions indistinguishable from wild-type cells, whereas plastid structure in bleached regions was dramatically altered, with stroma and grana thylakoids generally lacking. Analyses of protein complexes isolated from photosynthetic membranes of Δycf10 plants, did not uncover any differences between wild-type and transformed tobacco plants, further supporting an unaltered membrane structure and photosynthetic activity of Δycf10 plants.

**CO₂-dependent photosynthesis in Δycf10 plants.** The loss of ycf10 obviously affects the response to the external Cᵢ concentration of mutant plants and causes a reduction of their photosynthetic activity of approximately 50% relative to wild-type plants. The same effect was observed in ycf10-deficient *Chlamydomonas* cells (Rolland et al. 1997). Nevertheless, nature and detailed mechanism of the CO₂ uptake are still unknown so that secondary effects of the gene loss on inorganic carbon uptake can not be ruled out. As mentioned above, the cyanobacterial homologue of ycf10 (cemA), cotA, is supposed to be involved in light-induced proton extrusion rather than CO₂ uptake and/or transport (Katoh et al. 1996ab). The extrusion of protons, observed in *Synechocystis*, causes the acidification of wild-type cell suspension in light and generates a transmembrane pH gradient across the cytoplasmic membrane, which
drives the transport of other ions, including C\textsubscript{i} ions. Interestingly, the \textit{ycf10} reading frame is conserved among all photosynthetically active plant species, except \textit{Euglena} (Hallie et al. 1993), an organism without a CO\textsubscript{2} concentrating mechanism that can raise the CO\textsubscript{2} concentration, e.g. in mesophyll cells (in higher plants), in order to favour the carboxylase activity of RuBisCo, the key CO\textsubscript{2}-fixation enzyme (Badger and Price 1994). The marker enzyme of a CO\textsubscript{2} concentrating mechanism, carbonic anhydrase, converts HCO\textsubscript{3}\textsuperscript{-} into CO\textsubscript{2} (Badger and Price 1994). Both, carbonic anhydrase and C\textsubscript{i} transport systems are present in several organisms, including cyanobacteria, algae, C\textsubscript{3} and C\textsubscript{4} plant species. The interrelation between \textit{ycf10} and the presence of a CO\textsubscript{2} concentrating mechanism in different organisms is shown in Table 9.

\textbf{Table 9.} Conservation of the \textit{ycf10} open reading frame in different plant species and its relation to the CO\textsubscript{2} concentrating mechanism (CCM).

<table>
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<th>species</th>
<th>\textit{ycf10}</th>
<th>CCM</th>
<th>references</th>
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<tr>
<td>\textit{Synechocystis}</td>
<td></td>
<td></td>
<td>Cyanobacteria</td>
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<tr>
<td>PCC6803</td>
<td>+</td>
<td>+</td>
<td>Katoh et al. 1996a, 1996b</td>
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<td>\textit{Euglena gracilis}</td>
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<td>-</td>
<td>Hallie et al. 1993, Yakota et al. 1989</td>
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<tr>
<td>\textit{Chlamydomonas reinhardtii}</td>
<td>+</td>
<td>+</td>
<td>Rolland et al. 1997</td>
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<tr>
<td>\textit{Plasmodium falciparum}</td>
<td>-</td>
<td>-</td>
<td>apicomplexan parasite</td>
</tr>
<tr>
<td>\textit{Epifagus virginiana}</td>
<td></td>
<td>-</td>
<td>Wolfe et al. 1992</td>
</tr>
<tr>
<td>\textit{Cyanophora paradoxa}</td>
<td></td>
<td>?</td>
<td>Stirewalt et al. 1995</td>
</tr>
<tr>
<td>\textit{Marchantia polymorpha}</td>
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<td>+</td>
<td>Bryophyta</td>
</tr>
<tr>
<td>\textit{Pinus thunbergii}</td>
<td>+</td>
<td>+</td>
<td>Gymnospermae (C3) Wadasugi et al. 1994</td>
</tr>
<tr>
<td>\textit{Nicotiana tabacum}</td>
<td>+</td>
<td>+</td>
<td>Angiospermae (C3) Shinozaki et al. 1986</td>
</tr>
<tr>
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<td>Angiospermae (C3) Schmitz-Linneweber et al. 2001</td>
</tr>
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<td>Angiospermae (C3) Hiratsuka et al. 1989</td>
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<tr>
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<td>Angiospermae (C3) Hupfer et al. 2000</td>
</tr>
<tr>
<td>\textit{Zea mays}</td>
<td>+</td>
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<td>Angiospermae (C4) Maier et al. 1995</td>
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</table>

Regarding the Glaucocystophyte \textit{Cyanophora paradoxa} the absence of \textit{ycf10} in the cyanoplast genome was not surprising, because it encodes less than 10\% of the genetic information encoded by the cyanobacterial genome. As most of the organelle-specific proteins are nucleus-encoded, it is likely that the \textit{ycf10} reading frame, similarly to the \textit{ndh} gene cluster,
was transferred to the nuclear genome (Löffelhardt and Bohnert 1994). Unfortunately, detailed analyses of $C_i$ uptake into Cyanophora cells are not available. Therefore, it remains under debate whether the Ycf10 (CemA) protein exerts its effect on inorganic carbon uptake directly or indirectly, although the data of Katoh et al. (1996a and 1996b) suggested a role of cyanobacterial CotA protein as an ion transporter rather than a component of the $C_i$ uptake system. To clarify the functional role of $ycf10$ in the envelope membrane, detailed analyses of proton movements from the chloroplast into the cytosol are required for both Chlamydomonas and tobacco. As $cotA$ was the first gene found to be involved in light-induced proton extrusion in Synechocystis (Katoh et al. 1996a and 1996b), it is tempting to assume a similar involvement of the $cemA$ gene product in the energization and/or regulation of an ion transport system in the chloroplasts.
5. SUMMARY

Plastid chromosomes from the variety of plant species contain several conserved open reading frames of unknown function, which most probably represent functional genes. The primary aim of this thesis was the analysis of the role of two such ORFs, designated ycf9 or hypothetical chloroplast reading frames, namely ycf9 (ORF62) and ycf10 (ORF229, cemA). Both were analyzed in Nicotiana tabacum (tobacco) via their inactivation using biolistic plastid transformation. A new experimental protocol, based on pulsed-field gel electrophoresis (PFGE), was established to reliably assess the homoplasmatic state of transformed plants.

1. Functional analysis of the ycf9 gene product: The inactivation of ycf9 in N. tabacum as well as in Chlamydomonas reinhardtii yielded a homoplasmatic mutant phenotype after several rounds of regeneration under selective pressure. The mutant plants grew photoautotrophically, but displayed two clear phenotypes, a light-sensitive one, increasing with the light intensity, and a dwarf phenotype under low-light combined with temperatures below 20°C. The ycf9 gene product was exclusively located in PSII core complexes. This localization was based on the isolation of protein complexes released from thylakoids by controlled, partial lysis, followed by sucrose density gradient centrifugation or 2D gel electrophoresis. This finding revised data of the literature. Biochemical analysis indicated an involvement of the protein in the interaction of the light harvesting antenna II complex (LHCII) with PSII cores. In particular, PSII-LHCII supercomplexes could no longer be isolated from transplastomic tobacco plants. Furthermore, the minor chlorophyll a/b-binding proteins CP26, and to a lesser extent CP29, were substantially reduced under most growth conditions analyzed, in both, tobacco and photoautotrophically grown Chlamydomonas mutants (Swiatek et al. 2001). The gene was therefore renamed psbZ. The ΔpsbZ-related alterations in the supramolecular organization of PSII complexes were accompanied by considerable modification in (i) the phosphorylation pattern of PSII subunits, (ii) the rate of de-epoxydation of xanthophylls, and (iii) the kinetics and amplitude of non-photochemical quenching. The proposed position of PsbZ in close proximity to CP43 enables the protein to interact with PSII cores to elicit an adaptation process in response to excess light excitation. The molecular mechanism underlying this energy dissipation process remains to be investigated.
2. **Functional analysis of the ycf10 gene product:** Biolistic plastid transformation was also used to inactivate the *ycf10* reading frame in tobacco. After several rounds of regeneration under selective pressure, homoplastomic plants were obtained. Northern analysis uncovered co-transcription of *ycf10* within the *psal-ycf4-ycf10-petA* gene cluster, with at least two promoter regions upstream of the *psal* gene. The mutant plants grew photoautotrophically and developed dark green leaves with numerous pale green to white regions, the latter devoid of photosynthetic activity. The loss of *ycf10* did not affect photosynthetic activity, as indicated by unaltered chlorophyll fluorescence. The tobacco *ycf10* gene product was localized in the chloroplast inner envelope membrane. Neither protein composition of stroma or thylakoid fractions, nor the stability of the photosynthetic protein complexes were affected in the mutant plants. In contrast, CO$_2$-dependent oxygen evolution was strongly reduced, with a maximum rate of C$_1$-dependent photosynthesis being approximately 50% lower than in wild-type plants. Two explanations can account for the observed phenomenon: (i) de-regulation of carbon-concentrating mechanisms in transformed cells, or (ii) an indirect effect on CO$_2$-uptake in ∆*ycf10* plants.

3. **Pulsed-field gel electrophoresis is an ideal tool to verify the homoplastomic state of transformed plants:** To enhance the sensitivity of detection of heteroplastic states, and to distinguish between plastome-located wild-type segments in transplastomic material and promiscuous DNA, a new approach was developed. Customary Southern and PCR techniques are not sensitive enough or not discriminating the latter alternatives, respectively. Pulsed-field gel electrophoresis allows to isolate virtually contamination-free plastid DNA. Plastid DNA isolated this way lacked traces of nuclear and mitochondrial DNA at a detection level of 50 DNA molecules. This excludes that gene-specific PCR amplification products originate from promiscuous nuclear or mitochondrial gene copies. Therefore, PFGE appears to be an ideal tool to investigate the homoplastomic state of transformed plants, especially when combined with radiolabeled probes and Southern techniques.
6. REFERENCES


ACKNOWLEDGMENTS

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