The tobacco chloroplast YCF4 gene is essential for transcriptional gene regulation and plants photoautotrophic growth

Muhammad Sarwar Khan1*, Rimsha Riaz1, Muhammad Majid1, Kashif Mehmood2, Ghulam Mustafa1 and Faiz Ahmad Joyia1

1Center of Agricultural Biochemistry and Biotechnology (CABB), University of Agriculture, Faisalabad, Pakistan, 2Department of Molecular and Cellular Biology, Summerlee Science Complex, University of Guelph, Guelph, ON, Canada

A tobacco chloroplast hypothetical open reading frame 4 (YCF4) has been reported as a non-essential assembly factor for photosynthesis based on an incomplete knockout of YCF4, just 93 of 184 amino acids from the N-terminus were knocked out. On the other hand, we removed the complete sequence of YCF4 from tobacco chloroplasts and observed that ΔYCF4 plants were unable to survive photoautotrophically as their growth was hampered in the absence of an external carbon supply, clearly showing that the YCF4 is essential for photosynthesis. Initially, the aadA gene was introduced into the tobacco plastome replacing the complete YCF4 gene through homologous recombination events. The replacement of YCF4 with aadA was confirmed by PCR and Southern blot analysis in ΔYCF4 plants. Homoplasmic ΔYCF4 plants had a light green phenotype, and the leaves became pale yellow as the plants grew older. The structure of chloroplasts of ΔYCF4 mutants of light green phenotype was studied using a transmission electron microscope (TEM), and the micrographs demonstrated structural anomalies in the chloroplasts; including shape, size, and grana stacking compared to the wild-type plants. Further, transcriptome analysis revealed that the expression of PSI, PSII, and ribosomal genes remained unchanged in ΔYCF4 plants. On the other hand, transcriptome levels of rbcL (Ribulose 1,5-bisphosphate carboxylase/oxygenase large subunit), LHC (Light-Harvesting Complex), and ATP Synthase (atpB and atpL) decreased, indicating that the YCF4 has the function(s) in addition to assembling the photosynthetic complex. This was confirmed by in-silico protein-protein interactions of full-length YCF4 as well as 93 and 91 of 184 amino acids from N- and C-termini of the full-length protein, which revealed that the C-terminus (91 aa) of YCF4 is important in interacting with other chloroplast proteins. These findings provide genetic support for the plastid YCF4 gene’s critical role in regulating the plastid gene expression and assembling the photosynthetic complex.

KEYWORDS
YCF4, heterotrophic, knockout, transcript analysis, null alleles, homoplasmic
Introduction

Plastid origin can be traced back over a billion years when a fossil red alga was discovered, providing substantial proof for the theory (Embley and Martin, 2006; Song et al., 2018) that plastids arose from endosymbiotic photosynthetic bacteria (Schimper, 1883). Plastids are the defining traits of plants and green algae, yet they maintain many prokaryotic properties. The bacterial genome has reduced dramatically over time, owing to gene loss and large-scale gene transfer to the nuclear genome. Thus, plastid genomes referred to as plastomes contain around 120-130 genes, the majority of which encode components of the organelle’s gene expression mechanism and photosynthetic apparatus and are structured in nucleoids. Plastids, on the other hand, have far more proteins than their plastomes can code for. As a result, most plastid proteins are now encoded by the nuclear genome and must be transported into the organelle after translation. Plastids have an extensive membrane system, thylakoids, which are condensed to form, grana, in addition to two envelope membranes. In thylakoid membranes of plants including cyanobacteria and algae two enormous multimeric chlorophyll-binding protein complexes, photosystems I and II are embedded that performs the first step in the oxygenic photosynthesis this process, responsible for converting sunlight into chemical energy (Blankenship, 2010; Kargul and Barber, 2011).

The plastid genome was first sequenced in tobacco followed by hundreds of higher plants that have been sequenced and characterized. This plethora of plastid genome data insights researchers for the functional characterization of plastid-encoded genes. Most of the chloroplast genes have been characterized for their role in the organellar stability or metabolic activities within the chloroplasts, yet few of these are still to be worked out for their function. They are labeled as ycf genes (hypothetical chloroplast open reading frames). Few of these have been characterized as non-essential genes whereas others as essentials. The YCF4 gene, one of these ycf’s, is a highly conserved protein in cyanobacteria, green algae, and land plants. Previously, it was discovered that the YCF4 gene product is involved in the formation of the PSI complex in Chlamydomonas reinhardtii (Boudreau et al., 1997). In YCF4 mutants, PSI activity was completely lost, resulting in autotrophic growth failure. Inactivation of the YCF4 homolog causes an increase in the PSI-II-to-PSI ratio in the cyanobacterium Synechocystis sp. PCC 6803. (Wilde et al., 1995). The thylakoid membrane-intrinsic YCF4 was found in complexes with the PSI subunits PsaA through PsaF and the opsin-related eyespot protein COP2 in Chlamydomonas (Ozawa et al., 2009). YCF4 has also been found as a protein component of the eyespot in Chlamydomonas chloroplasts (Schmidt et al., 2006), suggesting that this has a second function in the eyespot (in conjunction with COP2). Furthermore, utilizing reverse genetics, YCF4 has been knocked out in tobacco, and the mutants have shown autotrophic growth (Krech et al., 2012). Tobacco YCF4 knockout mutants were able to assemble enough PSI to enable modest autotrophic growth. However, the YCF4 mutants did not grow autotrophically in our investigations, and extremely slow growth was detected under controlled settings using sucrose as a carbon source, but not in wild conditions.

In the present studies, we have removed the complete YCF4 gene sequence encoding 184 amino acids and developed homoplasmic ΔYCF4 plants, exhibiting very slow growth on an artificial medium supplemented with varied sucrose levels. These mutants unlike previously reported YCF4 mutants (Krech et al., 2012) where the partial sequence of YCF4 encoding 93 amino acids from the N-terminal region of YCF4 has been removed leaving 91 amino acids of the C-terminal intact, failed to grow autotrophically under normal conditions in peat moss-containing pots. It was observed by in-silico protein-protein interactions of full-length YCF4 as well as 93 and 91 of 184 amino acids from N- and C-termini, respectively of the full-length protein that the C-terminus (91 aa) of YCF4 is interacting with other chloroplast proteins.

Materials and methods

Plant material for targeted knockout of YCF4

Nicotiana tabacum L. var. Petit Havana was grown at 25 ± 1 °C under 16 hrs light (white light: 100 µmol·m⁻²·s⁻¹) and 8 hrs dark regime in a growth room. The sterilized seeds were cultured on RMOP medium containing MS salts (4.33 g/L), Myoinositol (100 mg/L), BAP (1.0 mg/L), NAA (0.1 mg/L) sucrose (3%) solidified with 0.026% Gelrite. Fully mature dark green leaves of 4-6 weeks old plants were used for targeted knockout of the YCF4 gene (Nazir and Khan, 2013).

Development of chloroplast transformation vector and selection of putative knocked out plants

Considering the location of the YCF4 gene where rbcL, accD, and psal are in the upstream region and ycf10, petA, and psbJ are in the downstream region of the YCF4. To develop chloroplast transformation vector for the targeted inactivation of YCF4, ycf10 sequence was cloned as right border flanking sequence whereas Psal along with few nucleotides of accD were cloned as left border flanking sequences. FLARE-S cassette having aadA

Abbreviations: YCF4, Hypothetical chloroplast open reading frame; PSI, Photosystem I; PII, Photosystem II; Cytb,f, cytochrome b,f complex.
(aminoglycoside 3-adenyltransferase) and gfp (green fluorescent protein) was cloned in between the plastid flanking sequences (Khan and Maliga, 1999). The resultant plastid transformation vector was coated on 0.6 μm gold particles and were bombarded on tobacco leaves using particle gun (Bio-Rad, USA). The bombarded leaves were chopped into tiny slices and cultured on RMOP medium containing 500 mg/L spectinomycin. Antibiotic resistant shoots were rooted on MS medium augmented with different levels of carbon (0, 0.5, 1.0, 1.5, 2.0, 2.5, and 3.0% sucrose). The plants were grown in normal medium augmented with different levels of carbon (0, 0.5, 1.0, 1.5, 2.0, 2.5, and 3.0% sucrose).

**Electron microscopy to study chloroplast ultrastructural variation in ΔYCF4 plants**

Transmission Electron Microscopy (TEM) was carried out following the protocol given by Islam et al., 2008. Leaves from ΔYCF4 and wild-type tobacco plants were cut into 1 to 2 mm² pieces and fixation was carried out using 0.2 M phosphate buffer (PBS, pH 7.2) containing 4.0% glutaraldehyde (v/v) at 4°C for 6-8 hours. Postfixation of samples was carried out in 1% osmium tetroxide (OsO₄) for 1 h at 4°C and then incubated in 0.2M PBS (pH 7.2) for 1-2 hours at room temperature. The dehydration was carried out using a graded series of ethanol and acetone and finally, the leaf pieces were embedded in Spurr’s resin. Ultrathin sections (~70 nm) of embedded leaf samples were prepared on an ultramicrotome (RMC Mt 7000) and mounted on copper grids to be viewed in the TEM (JEOL, Model JEM-1010) at an accelerating voltage of 90.0 kV. Multiple images of each section were recorded by exposing a photographic film and later developed in the darkroom.

**Determination of physiological parameters**

Various photosynthesis-related physiological parameters were determined using IRGA (Infrared gas analyzer). The physiological parameters included photosynthetic rate, transpiration rate, substomatal conductance, substomatal CO₂, and light intensity.

**Heterotrophic/autotrophic nature of growth of YCF4 mutants**

To assess the impact of carbon starvation on the growth, mutant as well as wild-type tobacco plants were cultured on MS medium augmented with different levels of carbon (0, 0.5, 1.0, 1.5, 2.0, 2.5, and 3.0% sucrose). The plants were grown in normal light. Data were recorded to see the impact of carbon starvation on mutant tobacco plants.

**Molecular docking of YCF4 protein with other photosynthesis-related proteins**

The molecular docking of full-length and truncated versions of YCF4 with other photosynthesis-responsive proteins was performed using an online web server ClusPro 2.0. It predicts the interaction between the candidate proteins through rigid-body docking based on Fast Fourier Transform, followed by clustering, and minimizing the docked complex to get the highly populated cluster having the least-energy conformation (Kozakov et al., 2017). The first 93 amino acids of the YCF4
were knocked out by Krech et al., 2012 in their study leaving 91 amino acids intact. Whereas, we have used 93 amino acids from the N-terminal and 91 amino acids from the C-terminal side of the YCF4 as well as full-length YCF4 for studying their interaction with YCF10, ribosomal proteins or RNA (rps16, rps2, rrrn16), subunits of PS-I (psaA, psaB, psaC, psaH), subunits of PS-II (psbA, psbB, psbC, psbD, psbE), alpha and beta chains of ATP synthase (atpB, atpI) and other photosynthetic proteins (rbcL, clpP, rpoA, rpoB, accD, petA, Light-harvesting complex (LHC) using ClusPro. The docked complexes with the maximum clustering members and minimum energy scores were selected. DIMPLLOT program of Lignplot+ v.4.5.3 was used to find the number of hydrogen bonds and the bond lengths between the interacting residues (Wallace et al., 1996).

Results

Development of YCF4 mutants and their purification to homoplastic level

The YCF4 mutants were developed by targeting the FLARE-S (aadA and gfp) cassette with flanking sequences of ycf10 and psal. Sequential amplification and cloning of ycf10 as right border flanking sequence and Psal together with a few nucleotides of acclD as left border flanking sequence resulted in the development of a plasmid transformation vector. The FLARE-S and its regulatory sequences were cloned in between the plastid flanking sequences. The resulting cassette was inserted into the plastid genome to inactivate YCF4 specifically. Leaf sections were cultivated in an RMOP medium containing a selective agent, spectinomycin (500 mg/L), after particle bombardment. The antibiotic-resistant shoots were putative transfectants, but they were indistinguishable from untransformed tobacco plants growing in comparable conditions. For the purification of transplastomic cells, they were exposed to additional rounds of selection and screening. Leaves were chopped into small pieces and cultured on the selective agent for this purpose. All of the plants had a light green to yellow phenotype after several rounds of continuous selection and screening, showing homotransplasmicity (Figure 1). Antibiotic-resistant shoots were putative transformants, but they were indistinguishable from untransformed tobacco plants growing in comparable conditions. For the purification of transplastomic cells, they were exposed to additional rounds of selection and screening.

Impact of YCF4 deletion on phenotype and mode of nutrition

Homoplastic ΔYCF4 plants appeared to have a distinct phenotype. The newly emerged younger leaves were green which gradually bleached out with the attainment of maturity. Lowermost leaves were almost white with minimum chlorophyll whereas top-most leaves were green with comparatively higher content of chlorophyll when plants were grown under normal light. The leaves continued to bleach until the whole plant was discolored and showed stunted growth with an inability to grow autotrophically. To eliminate the possibility that standard light (60 µmol m-2 s-1) may cause photobleaching, the plants were grown under low light (30 µmol m-2 s-1), and a light green phenotype was maintained in the ΔYCF4 plants. Further, to assess the impact of the autotrophic mode of nutrition, plants were grown at different levels of sucrose (0, 0.5, 1.0, 1.5, 2.0, 2.5, and 3.0%). Mutant plants cultured on 0, 0.5, and 1% sucrose were unable to survive and no leaf development was observed. However, plants cultured on 1.5, 2.5, and 3.0% sucrose showed growth, and leaf development was observed with increased sucrose concentration. The ΔYCF4 plants cultured on 3% sucrose were light leaves green in color compared with the wild-type plants which were green in color with normal growth (Figure 4). Further, several mutant plants growing on 3% sucrose were shifted to compost-containing pots for acclimatization, but they were unable to survive. This confirmed that YCF4 mutants are unable to survive autotrophically rather they require an additive carbon source for sustainability and growth.

Electron microscopy revealed ultrastructural variations between normal and ΔYCF4 plants

To investigate the ultrastructural variations in chloroplasts due to the absence of YCF4, leaf tissues from ΔYCF4 mutant and wild-type plants were analyzed using a transmission electron
microscope (TEM). Ultrastructural studies revealed that chloroplasts in knockout plants underwent substantial structural changes that may be correlated with the absence of YCF4 protein. Figure 5 revealed distinct variations in chloroplast size and shape between knockout plants and wild-type (normal) plants. The TEM results showed that chloroplasts in wild-type plants were oblong in shape and larger in size than those of mutant knockout plants which were almost rounded. Further, the thylakoid membranes were densely packed in chloroplasts of wild-type plants as compared to those of knockout plants. In knockout plants, the grana thylakoids were less discrete, and their stacks exhibited a loss of their orderly structure. As the thylakoid membranes became less organized some vesicular structures appeared in mutant chloroplasts (Figure 5A).
FIGURE 2
Confirmation of transgene integration into plastid genome and determining the plasmy level by Polymerase Chain Reaction. (A) Physical map showing the positions of the primers used. (B) Entire pool of spectinomycin-resistant plants regenerated on regeneration medium were screened for the presence of marker gene padA-specific primer set (A19/A20). M is 1 kb marker DNA, WT is the untransformed tobacco, and T1-T7 are putative transplastomic plants. (C) Out of screened plants four as T1-T4 were randomly selected for the determination of plasmy level using S19/S20 primer set that lands on flanking sequences: M represents 1.0 kb DNA ladder, WT represents wild-type tobacco plant whereas T1-T4 represents deletion level of ycf4 from the chloroplast transgenic plants regenerated on spectinomycin-containing regeneration medium. Amplification of a fragment of 4.0 kb indicates transgene integration whereas amplification of a fragment of 2.0 kb represents wild-type plastid DNA.

FIGURE 3
Confirmation of plasmy level of ycf4 plants through Southern Blot Analysis. (A) Physical map showing location of primers used to amplify the probe. (B) Total cellular DNA of ycf4 and wild-type tobacco plants was digested with BamHI, transferred onto the nitrocellulose membrane, and hybridized against a biotin-labeled probe prepared from ycf10 that amplifies both transformed and wild-type chloroplast DNA since ycf10 gene-specific probe detects all plastome copies irrespective to their genotype whether transformed or wild-type (untransformed). However, these plants were selected from an already screened pool (T1-T4 as shown in Figure 2C) of transgenic plants.
Physiological and photosynthetic performance of YCF4 mutants

The YCF4 gene deletion appeared to have profound effects on the photosynthetic and physiological performance of mutant plants (Figure 6A). The mutants were unable to attain normal contents of total chlorophyll as the topmost young leaves of mutant plants accumulated 2.6 mg/g of chlorophyll compared to the wild-type plant leaves (3.1 mg/g). The levels were decreased up to 99.98% in non-photosynthetic cells of mutants as the plant matures, from top to bottom (Figure 6B). Likewise, physiological parameters including photosynthetic rate (A), transpiration rate (E), stomatal conductance (gs), sub-stomatal CO₂ (Ci), and photosynthetic photon flux density (q also lux) also revealed that YCF4 plants were physiologically incompetent as compared with normal tobacco plants (Figure 7).

Transcript analysis of plastid-encoded genes in ΔYCF4 plants

The effect of YCF4 deletion on the expression of plastid-encoded genes was determined by transcript analysis of mutant plants. No transcripts were detected in purified ΔYCF4 plants since the gene has been completely replaced with a marker gene. The transcripts levels of psaA, psaB, psaC and psaH encoding photosystem-I proteins appeared the same in the ΔYCF4 plants and in normal untransformed tobacco plants. Likewise, the accumulation of psbA, psbB, psbC, psbD, psbE transcripts encoding Photosystem-II proteins were not decreased significantly in these mutants suggesting that YCF4 deletion did not have a direct role in the transcriptional regulation of PS-I and PS-II genes. Further, the transcript levels of ribosomal protein-encoding genes rps16, rps2, and...
rn16, and of ycf10, rpoA, rpoB, aacD and petA remained unchanged. Interestingly, the transcript levels of rbcL and lhc genes were significantly reduced, suggesting that YCF4 deletion affects the accumulation of RUBISCO and LHC1, eventually the photosynthesis (Figure 8).

**YCF4 interaction with other photosynthesis proteins**

The molecular interaction of full-length YCF4 with PS-I subunits indicated strong interaction with psaB, psaC, and psaH, each having seven hydrogen bonds with YCF4. But YCF4+psaC complex showed the most stable interaction among them with bond lengths of 2.62-2.93Å (Supplementary Figure S1). The bond length corresponds to the strength of interaction between interacting molecules and should be <4Å as the greater the bond length, the weaker will be the interaction. Likewise, among the PS-II subunits psaE showed a peculiar binding pattern and formed five hydrogen bonds with YCF4 (Supplementary Figure S2). The ATP synthase consists of two chains namely the alpha chain and beta chain. Both chains are known to interact with YCF4 to carry out photosynthesis (Yamori et al., 2011). Only the beta chain (atpB) revealed an effective docking pattern with YCF4 forming twelve hydrogen bonds with a bond length of 2.56-3.15Å (Supplementary Figure S3). Ribosomal proteins were also investigated as they are reported to play an essential role in the development of plant phenotypes.

![Figure 5](image-url)  
**Figure 5**  
Transmission electron microscopy of ycf4 and untransformed tobacco leaves shows the presence of defective plastids in the leaves of bleached plants. (A, B) Chloroplasts in light green leaves of ycf4 plants (C, D) Chloroplasts in lush green leaves of wild-type plants. The micrographs showed that ycf4 knockout caused structural anomalies including smaller size and rounded shape of chloroplasts in leaves.
Horiguchi et al. (2012). rrn16 showed a strong affinity with YCF4 protein forming ten hydrogen bonds within the bond length range of 2.63-3.19 Å. Likewise, the interaction of YCF4 with other plastidic proteins was also determined that showed the maximum binding affinity with rbcL among all the candidate proteins.

These photosynthetic proteins were further docked with carboxyl and amino terminus of YCF4 protein to elucidate which portion of the protein is more essential for photosynthesis. Among PS-I proteins, the amino terminus of YCF4 revealed maximum interaction with psaB forming five hydrogen bonds while twelve hydrogen bonds were observed in the carboxyl terminus of the YCF4+psaH complex. The psbE of PS-II bonded with the amino terminus of YCF4 effectively as compared to other proteins but formed only six hydrogen bonds while psbC strongly docked making thirteen hydrogen bonds with the terminus of YCF4. The carboxyl terminus of YCF4 also demonstrated durable interaction with other PS-II proteins but weaker than psbC. The docked complexes of the carboxyl terminus of YCF4 with rps2, rps16 and rrn16 showed 7, 11, and 6 hydrogen bonds respectively, while the amino terminus of YCF4 bonded with these proteins comparatively stronger showing 14, 18, 18 hydrogen bonds with rps2, rps16 and rrn16 (Supplementary Figure S4). Furthermore, the beta chain of ATP synthase is supposed to interact better compared to the alpha chain. The beta chain formed only eight hydrogen bonds with the amino terminus of YCF4 however twenty-eight hydrogen bonds were found in the carboxyl terminus of YCF4+atpB (Supplementary Figure S3). Likewise, assessing other proteins, rpoB found to be well-interacted with the truncated versions of YCF4 than other proteins revealing nine bonds with amino terminus and twenty-five bonds with carboxyl terminus of YCF4 (Supplementary Figure S5).

The YCF4 was also docked with the four core subunits of LHC of PS-I and a nuclear-encoded small subunit (RBCS) of RUBISCO to further validate the findings. Their carboxyl
terminus showed stronger interaction with a high number of hydrogen bonds forming between them. The relevance of the carboxyl terminus of YCF4 is once again demonstrating its strong connection with LHCA1, LHCA2, LHCA3, LHCA4, and RRBC (Table 1). The number and length of hydrogen bonds in the docked complexes of full-length and truncated YCF4 variants with many other proteins are shown in Table 1. Thus, these in-silico investigations support the current study’s notion that the carboxylic terminus of YCF4 is more crucial for photosynthesis than the amino terminus.

Discussion

The photosystem I complex requires both plastid and nuclear-encoded proteins for biosynthesis. PSI assembly requires the association of several redox cofactors, chromophores, and Fe-S clusters. Ycf4 is a crucial auxiliary element in the PSI assembly process. It has previously been shown to be important in photosystem I (PSI) formation in the unicellular green alga *Chlamydomonas reinhardtii*, with ycf3- and YCF4-deficient mutants unable to develop photoautotrophically and accumulate PSI (Boudreau et al., 1997). They concluded that Ycf3 and Ycf4 are not essential for PSI subunit synthesis but are most likely involved in PSI complex assembly. Orf184 mutants of *Cyanobacterium synechocystis* grew normally like wild-type cells, according to Wilde et al. (1995). However, the pigment content of mutant cells (especially the phycocyanin to chlorophyll ratio) differed significantly from that of wild-type cells. Another study found that YCF4 mutants could maintain photoautotrophic growth. They believed the YCF4 gene product was not necessary for photosynthesis. Despite the fact that mutants had lower PSI levels, this was not due to a deficit in plastid gene expression. They concluded that Ycf3 and Ycf4 are not required for the synthesis of PSI subunits but are most likely involved in the assembly of the PSI complex. However, pigment composition (particularly phycocyanin to chlorophyll ratio) of mutant cells was distinctly different from those of wild-type cells. Another research group also concluded that YCF4 mutants were able to sustain photoautotrophic growth. They were of the view that the YCF4 gene product is not essential for photosynthesis. Though mutants were deficient in PSI contents, this deficiency was not caused by a defect in plastid gene expression. Rather, it was suggested that YCF4 plays a key role at the post-translational stage, resulting in faulty PSI assembly or reducing its stability (Krech et al., 2012).

Contrarily, we establish that tobacco homoplastic YCF4 mutants are heterotrophic and cannot survive in an autotrophic environment. Carbon starvation appeared to inhibit plant growth, as mutants were unable to grow on MS medium containing sucrose up to 10 mg/L. Plants appeared to survive at 15-30 mg/L, although their growth was limited. When switched to compost-containing pots, these plants were photosynthetically incompetent and could not grow photoautotrophically. Our findings contradict that of...
Transcript analysis of Δycf4 and wild-type tobacco plants. Transcripts show the expression level of genes in both wild-type and Δycf4 plants. The expression of various subunits of photosystem-I (psaA, psaB, psaC, psaH) and photosystem-II (psbA, psbB, psbC, psbD, psbE) remained unaltered. The absence of a transcript in Δycf4 indicates the purification of the Δycf4 plants to homoplasmy.

**FIGURE 8**

Transcript analysis of Δycf4 and wild-type tobacco plants. Transcripts show the expression level of genes in both wild-type and Δycf4 plants. The expression of various subunits of photosystem-I (psaA, psaB, psaC, psaH) and photosystem-II (psbA, psbB, psbC, psbD, psbE) remained unaltered. The absence of a transcript in Δycf4 indicates the purification of the Δycf4 plants to homoplasmy.
Krech et al. (2012), who claimed that photoautotrophic growth was possible in YCF4 mutant plants.

Our findings, however, are consistent with those of Boudreau et al. (1997), who knocked out the nearly entire YCF4 gene from *C. reinhardtii* and concluded that the YCF4 gene product is essential for photosynthesis. The expression of the plastid-encoded genes encoding for the key subunits of photosystem I (*psaA*, *psaB*, *psaC*, and *psaH*) and Photosystem-II (*psbA*, *psbB*, *psbC*, *psbD*, *psbE*) were unaffected in YCF4 plants, indicating that Ycf4 deletion did not appear to have any direct role in photosystem biogenesis. However, YCF4 plants had much lower levels of *lhc* (light-harvesting complex) and *rbcL* (large subunit of ribulose-1,5-bisphosphate carboxylase-oxygenase) expression than wild-type normal plants. The LHC is important for the formation of a super complex photosystem (PSI) whereas rbcL is critical for RUBISCO. Decreased expression of LHC and rbcL may affect the conformation of the photosystem and accumulation of functional RUBISCO respectively, resulting in defective photosynthesis.

In the knockout plants, the microscopic studies revealed that chloroplast structure was abnormal, presumably due to the lack of YCF4 protein. Wild-type chloroplasts were substantially larger and oblong in shape, but knockout chloroplasts were much smaller and spherical. The thylakoid membranes appeared to be less organized with certain vesicular structures. Chloroplasts of non-green senescing Broccoli florets have shown similar disorganization and subsequent formation of vesicular structure (Terai and Watada, 2000) that has been attributed to the disorganization and disintegration of thylakoid membranes.

The size of the deleted YCF4 section may be a fundamental difference in the performance of YCF4 tobacco mutants generated by Krech et al. (2012) and mutants reported in the current investigations. The carboxyl terminus spanning 91 amino acids of the YCF4 protein had not been

### TABLE 1 Number of hydrogen bonds and the range of bonds length in docked complexes.

| Docked complex | Hydrogen bonds (full-length range bonds of YCF4) | Hydrogen bonds (amino terminus of YCF4) | Hydrogen bonds (carboxyl terminus of YCF4) |
|----------------|-----------------------------------------------|---------------------------------------|--------------------------------------------|
| PS-I           |                                               |                                       |                                            |
| ycf4+psaA      | 3 2.85 - 3.07                                 | 3 2.84 - 2.93                         | 5 2.74 - 3.02                              |
| ycf4+psaB      | 7 2.58 - 3.22                                 | 5 2.75 - 2.89                         | 12 2.61 - 3.20                             |
| ycf4+psaC      | 7 2.62 - 2.93                                 | 1 2.50                                | 8 2.62 - 3.19                              |
| ycf4+psaH      | 7 2.57 - 3.01                                 | 4 2.73 - 3.05                         | 17 2.57 - 3.26                             |
| ycf4+LHC       | 3 2.62 - 2.91                                 | 4 2.72 - 2.85                         | 9 2.83 - 3.20                              |
| ycf4+LHCA1     | 5 2.03 - 2.76                                 | 4 2.35 - 3.76                         | 9 2.45 - 3.76                              |
| ycf4+LHCA2     | 1 2.87 - 3.22                                 | 3 2.78 - 3.43                         | 12 2.01 - 2.92                             |
| ycf4+LHCA3     | 4 2.56 - 3.48                                 | 6 2.46 - 3.88                         | 10 2.64 - 3.67                             |
| ycf4+LHCA4     | 4 2.09 - 3.22                                 | 5 2.55 - 2.76                         | 9 2.67 - 3.66                              |
| PS-II          |                                               |                                       |                                            |
| ycf4+psbA      | 4 2.73 - 3.21                                 | 3 2.67 - 3.15                         | 8 2.73 - 3.28                              |
| ycf4+psbB      | 4 2.73 - 3.05                                 | 2 3.04 - 3.08                         | 6 2.47 - 3.07                              |
| ycf4+psbC      | 2 2.77 - 2.79                                 | 2 2.85 - 2.92                         | 13 2.68 - 3.12                             |
| ycf4+psbD      | 3 2.66 - 3.08                                 | 5 2.69 - 3.12                         | 6 2.61 - 2.99                              |
| ycf4+psbE      | 5 2.59 - 3.01                                 | 6 2.66 - 3.16                         | 6 2.72 - 3.17                              |
| Ribosomal      |                                               |                                       |                                            |
| proteins       |                                               |                                       |                                            |
| ycf4+rrn16     | 10 2.63 - 3.19                                | 6 2.72 - 3.18                         | 18 2.55 - 3.29                             |
| RNA            |                                               |                                       |                                            |
| ATP synthase   |                                               |                                       |                                            |
| ycf4+atpB      | 12 2.56 - 3.15                                | 8 2.62 - 3.12                         | 28 2.54 - 3.27                             |
| Other genes    |                                               |                                       |                                            |
| ycf4+ycf10     | 5 2.57 - 3.33                                 | 1 2.85                                | 7 2.69 - 3.25                              |
| ycf4+rbcL      | 13 2.56 - 3.15                                | 8 2.75 - 3.31                         | 17 2.54 - 2.98                             |
| Ycf4+rbcS      | 10 2.75 - 3.10                                | 9 2.64 - 3.22                         | 14 2.58 - 2.96                             |
| ycf4+clpP      | 6 2.52 - 2.99                                 | 6 2.63 - 2.92                         | 13 2.64 - 2.32                             |
| ycf4+rpoA      | 12 2.45 - 3.34                                | 7 2.72 - 3.09                         | 12 2.64 - 2.32                             |
| ycf4+rpoB      | 9 2.64 - 3.29                                 | 9 2.65 - 3.33                         | 25 2.62 - 2.33                             |
| ycf4+accD      | 10 2.62 - 3.29                                | 4 2.70 - 3.29                         | 7 2.71 - 2.88                              |
| ycf4+petA      | 5 2.88 - 3.22                                 | 5 2.71 - 2.83                         | 10 2.63 - 3.00                             |
knocked out, and the resultant mutants were able to grow photoautotrophically whereas, we have knocked out the entire open reading frame and found that knocked plants could not survive photoautotrophically and grew stuntedly at 30 g/L sucrose. The results of the in-silico studies supported our findings since the interactions between the photosystem-I subunits psaB, psaC, psaH, and LHC and the carboxyl terminus of the YCF4 were stronger than those between the amino and carboxyl termini. Similar to this, the interaction of YCF4 with the large (encoded by chloroplast) and small (encoded by nuclear) subunits of RuBicCO also supported the significance of the carboxyl terminus. Hence, this study is direct evidence of the fact that deletion of full-length YCF4 has made the plants unable to survive photo-autotrophically as interaction with all the proteins engaged directly or indirectly in photosynthesis has revealed strong binding patterns with the carboxyl terminus of YCF4.

We came to the conclusion that deletion of the entire YCF4 open reading frame prevents tobacco plants from growing autotrophically, even though photosystem biogenesis is a complex process and various co-factors and related proteins still need to be investigated. Under heterotrophic circumstances (30g/L sucrose), plants could thrive, but they grew stuntedly and died when placed in pots with peat moss.

Data availability statement

The original contributions presented in the study are included in the article/Supplementary Materials. Further inquiries can be directed to the corresponding author.

Author contributions

MK conceived the idea, supervise the students, and wrote the manuscript. MM, RR, KM, and GM performed data analysis. Whereas, FJ handled microscopic studies and helped in the manuscript write-up. All authors contributed to the article and approved the submitted version.

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Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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Supplementary material

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fpls.2022.1014236/full#supplementary-material

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