



Synthesis of Chlorophyll-Binding Proteins in a Fully Segregated $\Delta ycf54$ Strain of the Cyanobacterium *Synechocystis* PCC 6803

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In the chlorophyll (Chl) biosynthesis pathway the formation of protochlorophyllide is catalyzed by Mg-protoporphyrin IX methyl ester (MgPME) cyclase. The Ycf54 protein was recently shown to form a complex with another component of the oxidative cyclase, Sll1214 (Cycl), and partial inactivation of the *ycf54* gene leads to Chl deficiency in cyanobacteria and plants. The exact function of the Ycf54 is not known, however, and further progress depends on construction and characterization of a mutant cyanobacterial strain with a fully inactivated *ycf54* gene. Here, we report the complete deletion of the *ycf54* gene in the cyanobacterium *Synechocystis* 6803; the resulting $\Delta ycf54$ strain accumulates huge concentrations of the cyclase substrate MgPME together with another pigment, which we identified using nuclear magnetic resonance as 3-formyl MgPME. The detection of a small amount (~13%) of Chl in the $\Delta ycf54$ mutant provides clear evidence that the Ycf54 protein is important, but not essential, for activity of the oxidative cyclase. The greatly reduced formation of protochlorophyllide in the $\Delta ycf54$ strain provided an opportunity to use ³⁵S protein labeling combined with 2D electrophoresis to examine the synthesis of all known Chl-binding protein complexes under drastically restricted *de novo* Chl biosynthesis. We show that although the $\Delta ycf54$ strain synthesizes very limited amounts of photosystem I and the CP47 and CP43 subunits of photosystem II (PSII), the synthesis of PSII D1 and D2 subunits and their assembly into the reaction centre (RCII) assembly intermediate were not affected. Furthermore, the levels of other Chl complexes such as cytochrome *b₆f* and the HliD–Chl synthase remained comparable to wild-type. These data demonstrate that the requirement for *de novo* Chl molecules differs completely for each Chl-binding protein. Chl traffic and recycling in the cyanobacterial cell as well as the function of Ycf54 are discussed.

Keywords: Ycf54, *Synechocystis* 6803, chlorophyll, photosystem II, protochlorophyllide, Mg-protoporphyrin IX methylester cyclase

INTRODUCTION

Chlorophylls (Chl) and Chl binding proteins are essential components of the photosynthetic apparatus. Together they act as principal light harvesting and energy transforming cofactors in photosynthetic organisms, as demonstrated by the structures of both photosystem I (PSI) and photosystem II (PSII; Jordan et al., 2001; Zouni et al., 2001; Umena et al., 2011). It is likely that at least for large Chl-binding subunits of PSI (PsaA, PsaB) and PSII (CP43, CP47) Chl molecules must be inserted into these proteins co-translationally as a prerequisite for correct protein folding (Chua et al., 1976; Eichacker et al., 1996; Müller and Eichacker, 1999; Chidgey et al., 2014). As demonstrated recently using a cyanobacterial $\Delta chlL$ mutant, which is unable to synthesize Chl in the dark, the availability of *de novo* Chl molecules is ultimately essential for synthesis of all central subunits of both photosystems (Kopečná et al., 2013). Nonetheless, there are unexplained aspects of the assembly of PSII subunits, such as the particular sensitivity of the CP47 subunit to the lack of *de novo* Chl (Dobáková et al., 2009; Kopečná et al., 2015).

The Chls are a group of modified tetrapyrrole molecules distinguished by their fifth isocyclic or E ring, the geranylgeranyl/phytol moiety esterified at C17 and a centrally chelated magnesium ion. The isocyclic ring arises from the cyclisation of the methyl-propionate side-chain at C-13 to the C-15 bridge carbon between rings C and D. In oxygenic phototrophs this biosynthetic reaction is catalyzed by the oxidative Mg-protoporphyrin IX monomethyl ester cyclase (MgPME-cyclase), which incorporates atmospheric oxygen into the C13¹ carbonyl group (Porra et al., 1996). Although studied in some detail, the enzyme responsible for the aerobic cyclisation reaction remains the least understood in the Chl biosynthesis pathway. The first gene identified as encoding an oxidative cyclase component was the *Rubrivivax gelatinosus* *acsF* (aerobic cyclisation system iron containing protein) locus (Pinta et al., 2002). Subsequently, AcsF homologs have been identified in all studied oxygenic photosynthetic organisms (Boldareva-Nuianzina et al., 2013).

The Ycf54 protein (12.1 kDa) has been shown recently to interact with the AcsF homolog Sll1214 (hereafter CycI; Peter et al., 2009) in the cyanobacterium *Synechocystis* PCC 6803 (hereafter *Synechocystis*; Hollingshead et al., 2012). Demonstrations that partial elimination of Ycf54 strongly impairs the formation of PChlide and causes Chl deficiency in both cyanobacteria and plants (Albus et al., 2012; Hollingshead et al., 2012) led to speculations that this protein is a catalytic subunit of the MgPME cyclase (Bollivar et al., 2014). Here, we clarify this issue by achieving the complete deletion of the *ycf54* gene in *Synechocystis*. Although the Chl content in this strain was very low, the MgPME-cyclase was apparently active, which demonstrated that the Ycf54 protein is not an essential subunit of the MgPME cyclase. On the other hand, the mutant contained a very low level of CycI and lacked a high-mass complex associated with the light-dependent PChlide oxidoreductase enzyme (POR). The greatly limited formation of PChlide in the *ycf54* mutant provided an opportunity to assess

the sensitivity of assembly pathways for all known Chl-proteins in cyanobacteria to the availability of *de novo* Chl. Interestingly, the deletion of the *ycf54* gene almost abolished the synthesis of PsaA/B subunits and PSII antennas CP47 and CP43, whereas the accumulation of other Chl-proteins showed little or no defects.

EXPERIMENTAL PROCEDURES

Growth Conditions

Synechocystis strains were grown photomixotrophically in a rotary shaker under low light conditions (5 mmol photons m⁻² s⁻¹) at 30°C in liquid BG11 medium (Rippka et al., 1979) supplemented with 10 mM TES-KOH (pH 8.2) and 5 mM glucose.

Construction of the $\Delta ycf54$ *Synechocystis* Strain

In order to disrupt open reading frame *slr1780* (*ycf54*), we prepared a construct for replacing the most this gene (bp 64–276) by a Zeocin resistance cassette. The sequences up- and down-stream (~300 bp) of the *ycf54* gene were amplified with the relevant primers and fusion PCR in conjunction with megaprimers (Ke and Madison, 1997) were used to anneal these either side of the Zeocin resistance cassette. The resulting PCR product was transformed into the GT-W *Synechocystis* substrain (Bečková et al., submitted) and transformants were selected on a BG11 agar plate containing 2 mg ml⁻¹ Zeocin. Complete segregation was achieved by sequentially doubling the concentration of antibiotic to a final concentration of 32 mg ml⁻¹ Zeocin.

Cell Absorption Spectra and Determination of Chl Content

Absorption spectra of whole cells were measured at room temperature using a Shimadzu UV-3000 spectrophotometer (Kyoto, Japan). To determine Chl levels, pigments were extracted from cell pellets (2 ml, OD₇₅₀ = ~0.5) with 100% methanol and the Chl concentration was determined spectroscopically (Porra et al., 1989).

Analysis of Pigments by HPLC

Pigments were extracted from equal quantities of cells by the method described in Canni e et al. (2013) and separated on a Phenomenex Aqua C₁₈ reverse phase column (5 mM

eluted over a linear gradient of 65 to 75% bu er A over 35 min.

Purification of 3-Formyl-MgPME for NMR Analysis

Pigments were extracted by phase partitioning from 6 L of $\Delta ycf54$ culture grown to an $OD_{750\text{ nm}}$ 1.2. One volume of diethyl ether was added to two volumes of cell culture in a separation funnel and the diethyl ether phase containing 3-formyl-MgPME was separated from the cell culture. Pigments were extracted from the cell culture three times. The diethyl ether was removed by rotary evaporation and the extracted pigments were re-suspended in a small volume of HPLC grade methanol. After centrifugation at $15,000 \times g$ for 10 min, 3-formyl-MgPME was purified by preparative HPLC. Ammonium acetate was removed from the HPLC purified 3-formyl-MgPME by solid-phase extraction on DSC-18 reverse-phase columns (Supelco). Solvents C, D, and E were QH_2O , 50% methanol (v/v) and 100% methanol, respectively. After equilibration of the column with 1.0 ml solvent D, the purified 3-formyl-MgPME, diluted 1/3 with QH_2O , was loaded and allowed to enter the column by gravity flow. The column was washed with 1 ml solvent C, then 1 ml solvent D to remove the ammonium acetate. The pigment was eluted into a glass vial with 300 ml methanol. The purified pigment was completely dried in a vacuum centrifuge and stored at $-20^\circ C$.

NMR Assignment of 3-Formyl-MgPME

The dried pigment from HPLC was re-suspended in 500 ml methanol-d4 (Sigma), centrifuged to remove any insoluble pigment, transferred to a 5 mm NMR tube and sealed. All NMR experiments were carried out on a Bruker Avance DRX 600 instrument equipped with a cryoprobe at an acquisition temperature of 298 K.

The one-dimensional selective Nuclear Overhauser Enhancement (NOE) experiments were recorded using a double pulsed field gradient spin echo selective NOE experiment (Stott et al., 1995) using an 80 ms 180° Gaussian pulse for the selective excitation and a 1 s mixing time, acquiring 1024 transients at each saturation frequency. The Total Correlation Spectroscopy (TOCSY) experiment was recorded using a 45 ms spin lock at a power of 8.3 kHz. Two carbon Heteronuclear Single Quantum Correlation (HSQC) experiments were recorded with carbon o sets of 60 and 140 ppm.

2D Electrophoresis, Immunodetection, and Protein Radiolabeling

Membrane and soluble protein fractions were isolated from 50 ml of cells at $OD_{750\text{ nm}} \sim 0.4$ according to Dobáková et al. (2009) using bu er A (25 mM MES/NaOH, pH 6.5, 5 mM $CaCl_2$, 10 mM $MgCl_2$, 20% glycerol). Isolated membrane complexes (0.25 mg/ml Chl) were solubilized in bu er A containing 1% *n*-dodecyl- β -D-maltoside.

To assess protein levels by immunodetection, the protein content of *Synechocystis* lysates was quantified spectroscopically (Kalb and Bernlohr, 1977), separated by SDS-PAGE (Novagen) and transferred to a nitrocellulose membrane. The membranes

were probed with specific primary antibodies and then with secondary antibodies conjugated to horseradish peroxidase (Sigma). The primary antibodies used in this study were raised in rabbits as described in Hollingshead et al. (2012), with the exception of CHL27 (anti-Cycl), which was purchased from Agrisera (Sweden).

Two-dimensional clear-native electrophoresis was performed essentially as described in Kopečná et al. (2013). Proteins separated in the gel were stained either by Coomassie Blue, or Sypro Orange, followed by transfer onto a PVDF membrane. Membranes were incubated with specific primary antibodies, and then with a secondary antibody conjugated with horseradish peroxidase (Sigma).

Radioactive pulse labeling of the proteins in cells was performed using a mixture of $[^{35}S]$ Met and $[^{35}S]$ Cys (Translabel; MP Biochemicals). After 30 min incubation of cells with labeled amino-acids, the solubilized membranes isolated from radiolabelled cells were separated by 2D-electrophoresis. The stained 2D gel was finally exposed to a phosphor-imager plate, which was scanned by Storm (GE Healthcare) to visualize labeled protein spots.

Relative Quantification of FLAG-Cycl and Captured Proteins in Pulldown Assays

Pulldown assays using N-terminal FLAG-tagged Cycl as bait, with both wild-type (WT) and $\Delta ycf54$ backgrounds, were carried out according to Hollingshead et al. (2012). FLAG eluates were concentrated to 100 ml using Amicon Ultra 0.5 ml 3 kDa MWCO ultrafiltration devices (Millipore). The proteins were then precipitated, reduced and S-alkylated according to Zhang et al. (2015). Proteolytic digestion was carried out with 1:25 w/w (enzyme:substrate) pre-mixed trypsin/Lys-C (1 mg/mL, Promega, mass spectrometry grade) at $37^\circ C$ for 2 h. The samples were then diluted with 75 ml 100 mM Tris-HCl, pH 8.5, 10 mM $CaCl_2$ and the digestion allowed to proceed for a further 18 h at $37^\circ C$. After the addition of 5ml 10% TFA, the samples were desalted using C_{18} spin columns (Thermo Fisher) and analyzed by nano-flow liquid chromatography (Ultimate 3000 RSLCnano system, Dionex) coupled to a mass spectrometer (Maxis UHR-TOF, Bruker or Q Exactive HF Orbitrap, Thermo Scientific). For Maxis data, mass spectra were internally calibrated with the lock-mass ion at m/z 1221.9906 then converted to MGF format using a script provided by Bruker. Q Exactive data-files were converted to MGFs using MSConvert¹. Protein identification was carried out by searching against the *Synechocystis* PCC 6803 proteome database (release date 02-08-2015, 3507 entries² using Mascot Daemon v. 2.5.1 running with Mascot Server v. 2.5 (Matrix Science), specifying trypsin as the enzyme in the search parameters and allowing for one missed cleavage. S-carbamidomethyl-cysteine and methionine oxidation were selected as fixed and variable modifications, respectively. MS and MS/MS tolerances were set to 0.01 Da and false discovery rates determined by searching of a decoy database composed of reversed protein sequences. The data-files and search results have

¹www.proteowizard.sourceforge.net

²www.uniprot.org/proteomes/UP000001425

been uploaded to the ProteomeXchange Consortium³ via the PRIDE partner repository (identifier DOI 10.6019/PXD003149).

Electron Microscopy

Wild-type and $\Delta ycf54$ cells were harvested in the log phase by centrifugation. Cell pellets were loaded into 200 nm deep specimen carriers (Leica Microsystems), pre-treated with 1% lecithin in chloroform and cryo-immobilized by high-pressure freezing using EM PACT2 (Leica Microsystems). Freeze-substitution was carried out as described by van de Meene et al. (2006) using an automatic freeze substitution unit (EM ASF, Leica). Samples were then infiltrated with graded series (1:2, 1:1, 2:1) of Spurr-acetone mixture (6–8 h for each), twice with 100% Spurr's resin (SPI Supplies) and finally embedded in fresh resin. The polymerization was performed at 60°C for 48 h. Ultra-thin 70 nm sections were cut on ultramicrotome (UCT, Leica), collected on formvar-coated copper grids and stained with uranyl acetate (5 min) and lead citrate (3 min). Grids were viewed with a JEOL 1010 transmission electron microscope operating at 80 kV equipped with a Mega View III camera (SIS GmbH).

RESULTS

Ycf54 Is Not Essential for Activity of MgPME Cyclase

In our previous report (Hollingshead et al., 2012) we described a $ycf54^-$ *Synechocystis* strain harboring an insertion of the Erythromycin resistance cassette in the *ycf54* gene. Although prolonged attempts to fully segregate the mutant allele into all copies of the chromosome were unsuccessful, the phenotype of the partially segregated strain was informative nevertheless, and it exhibited an obvious defect in PChlide formation. However, the capability of *Synechocystis* cells to tolerate deletions of important genes also depends on the 'WT' substrain used. For instance, a previous attempt to inactivate *gun4*, another gene crucial for Chl biosynthesis, was achieved in the non-motile *Synechocystis* GT-P substrain but it failed in the motile PCC-M (compare Wilde et al., 2004; Sobotka et al., 2008). Thus, in order to obtain a fully segregated *ycf54* mutant, we prepared a new construct for replacement of the *ycf54* gene and transformed GT-P, GT-S, GT-W and PCC-M substrains; the GT-P substrain has been used in our previous work (Hollingshead et al., 2012). Interestingly, the *ycf54* deletion readily segregated in the GT-W substrain (Figure 1A) under low light (5 mmol photons.m⁻².s⁻¹) and photomixotrophic conditions; all attempts to segregate the *ycf54* deletion in other substrains failed (not shown). For the purposes of the work reported here the GT-W substrain is designated as the WT; a detailed analysis of GT-P and GT-W including genome sequencing is presented elsewhere in this issue (Bečková et al., submitted).

The fully segregated $\Delta ycf54$ strain did not grow photoautotrophically, although supplementation of the growth medium with glucose made photomixotrophic growth possible at

light intensities up to 100 mmol photons.m⁻².s⁻¹. Examination of the absorption spectra from cells normalized for optical density at 750 nm (OD₇₅₀), shows that the Chl absorbance maxima at 439 and 679 nm and the carotenoid absorbance maximum at 494 nm are severely depleted in $\Delta ycf54$, whilst the absorbance maximum of the phycobiliproteins at 624 nm remains unchanged when compared to the WT (Figure 1B). Mutant cells contained only about 13% of WT Chl (Figure 1B) and the whole cell spectrum showed a large absorbance peak at 422 nm, indicating a substantial accumulation of MgPME (Figure 1B) (Hollingshead et al., 2012).

Identification of the Chl Precursors that Accumulate in the $\Delta ycf54$ Mutant

Previously we reported that Chl biosynthesis in a partially segregated $ycf54^-$ mutant was blocked at the MgPME cyclase step, which causes accumulation of high levels of MgPME, the substrate of the cyclase, and lesser levels of an unknown pigment with a Soret peak at 433 nm (Hollingshead et al., 2012). To examine the photosynthetic precursor pigments present in $\Delta ycf54$, methanol extracts from low light grown cells were separated by HPLC (Figure 2A). As with $ycf54^-$, $\Delta ycf54$ synthesized high levels of MgPME and lesser levels of the unknown pigment. Given that the Soret band of this pigment is situated between the Soret peaks of MgPME at 416 nm and PChlide at 440 nm (Figure 2B) we proposed that it could be an intermediate of the cyclase reaction.

Nuclear magnetic resonance spectroscopy was used to determine the identity and structure of this unknown pigment, which was extracted by diethyl ether/water phase partitioning from the medium of $\Delta ycf54$ cells grown under very low light conditions. This pigment was purified to homogeneity by preparative HPLC. The one-dimensional ¹H spectrum (Figure 3A) shows the reasonable degree of purity of the pigment, with impurities indicated by an asterisk; the signals downfield of 5 ppm represent minor contaminants and methanol, whilst the impurity (a)1(steno)JT0tG02eld)-786(of)-556(of5

³<http://proteomecentral.proteomexchange.org>

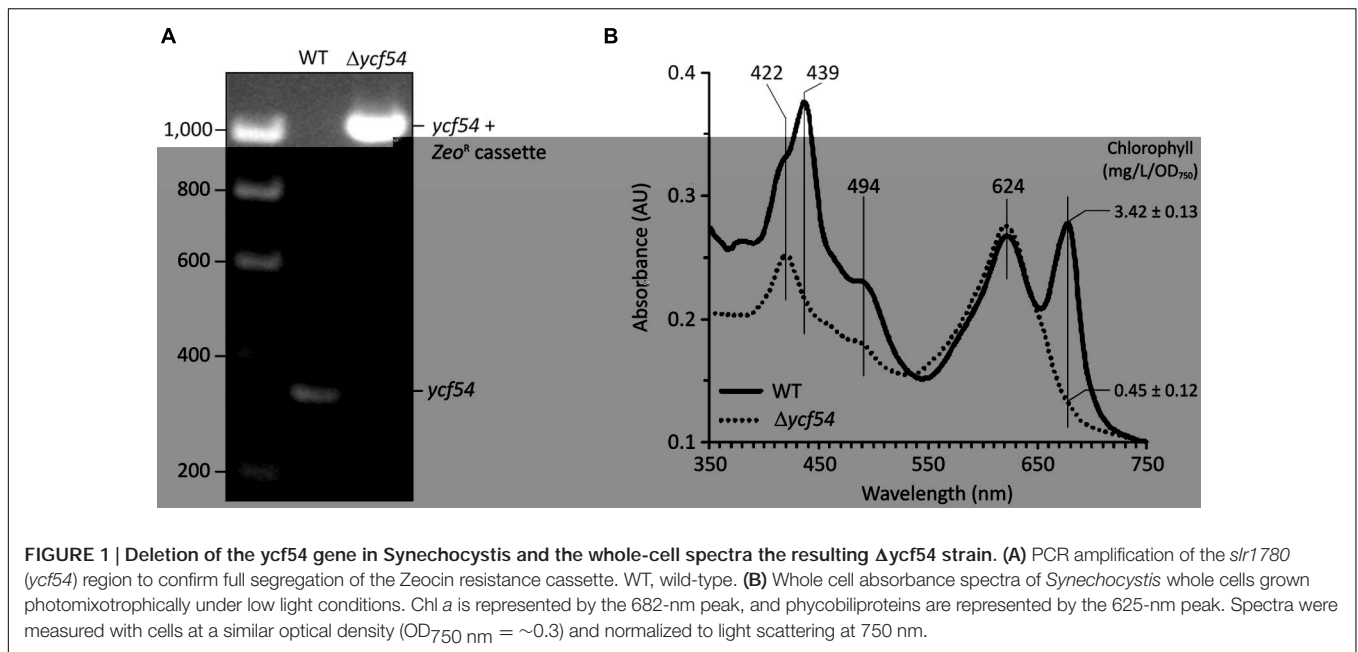


FIGURE 1 | Deletion of the *ycf54* gene in *Synechocystis* and the whole-cell spectra of the resulting $\Delta ycf54$ strain. (A) PCR amplification of the *slr1780* (*ycf54*) region to confirm full segregation of the Zeocin resistance cassette. WT, wild-type. **(B)** Whole cell absorbance spectra of *Synechocystis* whole cells grown photomixotrophically under low light conditions. Chl *a* is represented by the 682-nm peak, and phycobiliproteins are represented by the 625-nm peak. Spectra were measured with cells at a similar optical density ($OD_{750\text{ nm}} \approx 0.3$) and normalized to light scattering at 750 nm.

signal has NOEs to both the 5-meso and 2-methyl protons, both of which were shifted downfield, and no through-bond connectivity in the TOCSY, verifying that this was a 3-formyl group which had replaced the 3-vinyl group of the MgPME. The NMR data are compiled, together with details of the acquisition parameters, in Supplementary Table 1, including Supplementary Figures S1–S3. Further confirmation that this signal represents a 3-formyl group comes from the $^1\text{H-NMR}$ spectra of Chl *d* (Fukusumi et al., 2012), which has a clear signal at ~ 11.5 assigned as the 3-formyl group. Thus, the unknown pigment is magnesium 3-formyl-protoporphyrin IX monomethyl ester (Figure 2D).

Effects of Removal of Ycf54 on Other Chl Biosynthesis Enzymes

In order to investigate levels of Chl biosynthetic enzymes, and to verify the loss of Ycf54 in the $\Delta ycf54$ mutant, lysates from WT and $\Delta ycf54$ cells were fractionated into membrane and soluble components. The appearance of the cell lysate fractions (Figure 4A) reflects their pigment composition; the WT whole cell lysate and membrane fractions are green, and in $\Delta ycf54$ whole cell lysate and membrane fractions are blue and orange, respectively, because of the near-absence of Chl. A western blot of each of these extracts was probed with antibodies raised against a wide range of Chl biosynthesis enzymes (Figure 4B). The immunoblot probed with the antibody to Ycf54 shows this protein is distributed evenly between the soluble and insoluble fractions and is not detected in $\Delta ycf54$, confirming the full segregation of this mutant. The absence of Ycf54 is also accompanied by a decrease in Cycl and geranylgeranyl reductase (ChIP) and increased relative levels of the Mg-chelatase subunits ChII and ChID, although no change was detected in the levels of ChIH (Figure 4B).

Mass spectrometry was used to quantify the effects of *ycf54* deletion, in terms of the ability of Cycl to associate with partner proteins *in vivo*. Pulldown assays with FLAG-tagged Cycl are already known to retrieve Ycf54 from cell extracts (Hollingshead et al., 2012), so this experiment was repeated using FLAG-Cycl in a $\Delta ycf54$ background. The amounts of PChlide oxidoreductase (POR), 3,8-divinyl (proto)chlorophyllide reductase (DVR) and ChIP captured in pulldown assays by FLAG-Cycl/WT and FLAG-Cycl in $\Delta ycf54$ were compared by mass spectrometry. Proteins extracted from FLAG eluates were digested with a combination of endoproteinase LysC and trypsin and the peptide fragments analyzed by nanoLC-MS/MS. The captured proteins were quantified relative to the Cycl bait as shown in Figure 5. Captured POR levels had decreased significantly in the $\Delta ycf54$ strain while DVR was reduced to an undetectable level. ChIP was only just detectable in one $\Delta ycf54$ replicate and relative to Cycl by three orders of magnitude in the other two.

Lack of PChlide Impairs Synthesis of PsaA/B and Inner PSII Antennae but the Accumulation of Other Chl-Binding Proteins Is Not Affected

To evaluate the effects of greatly reduced Chl on the photosystems in $\Delta ycf54$ compared to the WT, photosynthetic membranes isolated from an equal biomass were gently solubilized with b-DDM and the membrane complexes were resolved by clear native electrophoresis (CN-PAGE), followed by SDS-PAGE in the second dimension. The resulting 2D CN/SDS-PAGE (Figure 6A) showed that $\Delta ycf54$ has drastically reduced levels of both photosystems, whilst the levels of other abundant membrane complexes such as ATP synthase, NADH:ubiquinone oxidoreductase and the cytochrome *b₆f* complex (shown by the western blot) are comparable between the two strains.

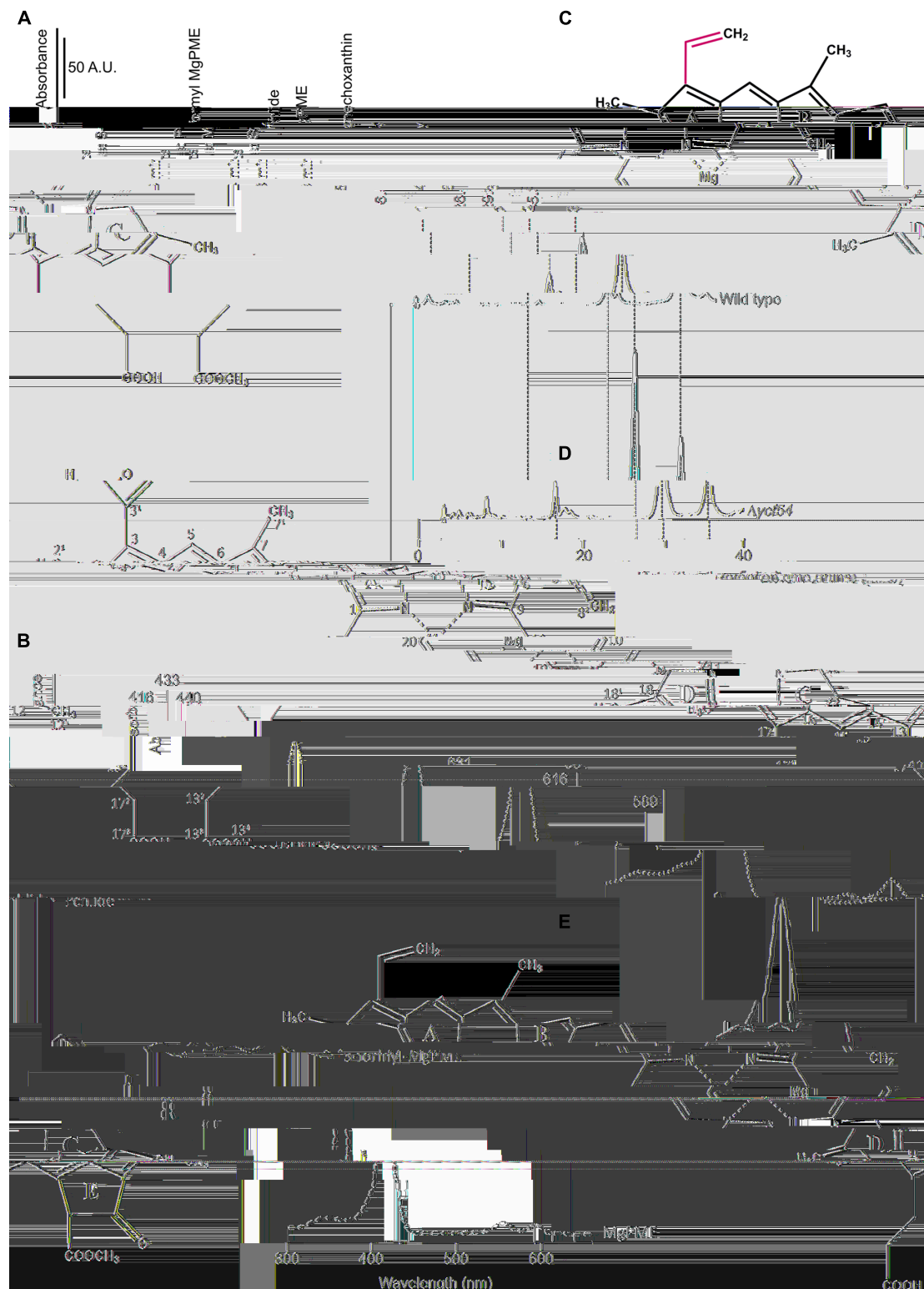


FIGURE 2 | HPLC analyses of pigments from WT and $\Delta ycf54$ cells. (A) Polar pigments were extracted with 80% methanol containing 0.2% (v/v) ammonium from an equal volume of cells at an $OD_{750\text{ nm}} \sim 0.7$ and analyzed on a Phenomenex C₁₈ column. Separation of precursors was detected by a diode array detector set to 432 nm, the Soret peak of 3-formyl MgPME, which is observed in $\Delta ycf54$ cells. The elution times of the pigments of interest are indicated. **(B)** Absorbance spectra of MgPME, Pchl and 3-formyl-MgPME, **(C)** Mg-protoporphyrin IX monomethyl ester (MgPME), **(D)** Mg-3-formyl-protoporphyrin IX monomethyl ester (3-formyl-MgPME), **(E)** protochlorophyllide (Pchl).

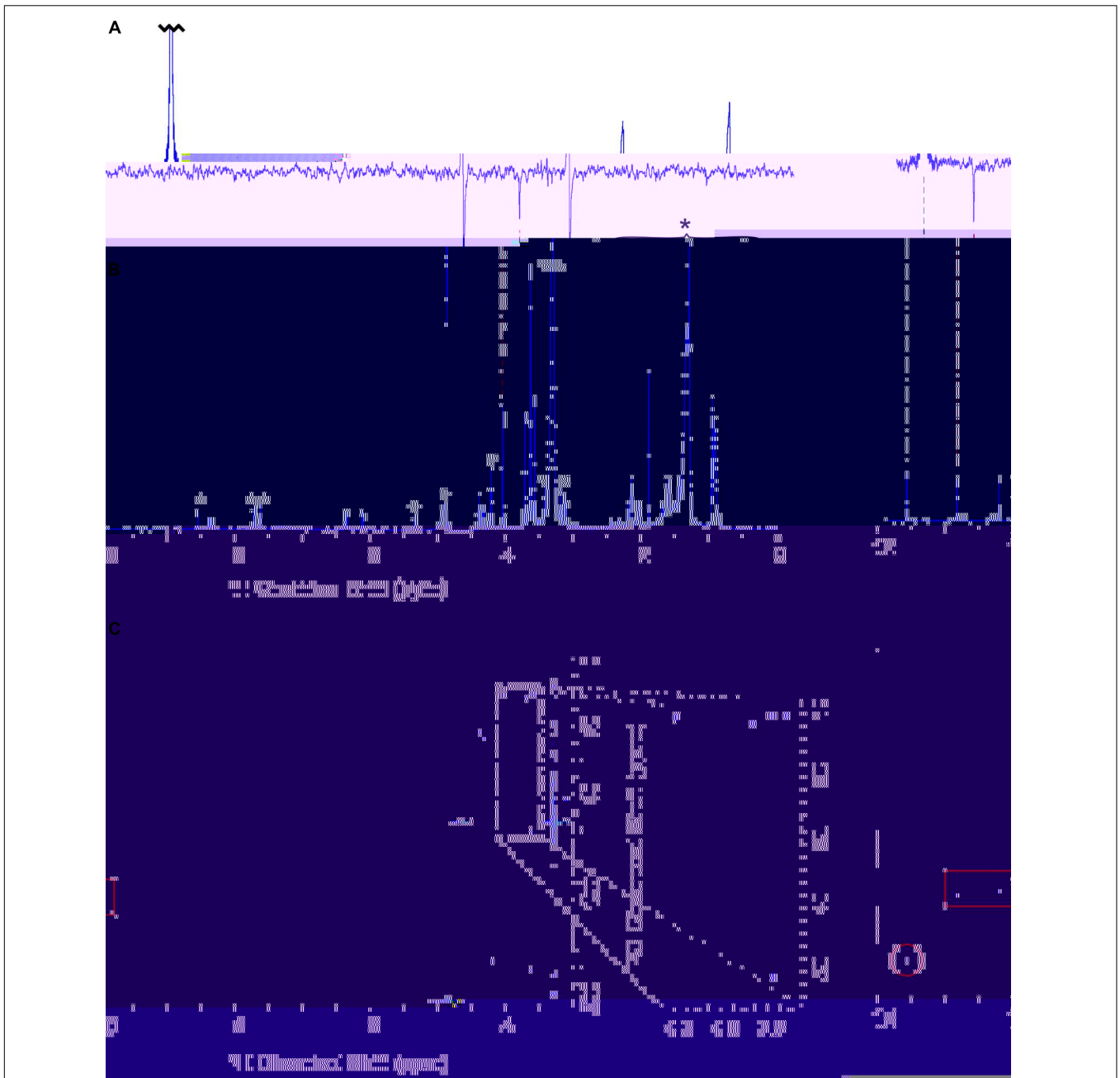


FIGURE 3 | NMR assignment of the A432 pigment accumulating in the $\Delta ycf54$ strain. (A) One dimensional selective NOE spectrum of 3-formyl-MgPME, selectively pulsed at 11.60 ppm (selectively exciting proton 3¹). Negative signals indicate NOE cross signal with protons 5 and 2¹ at 10.86 and 4.07 ppm, respectively. **(B)** ¹H-NMR spectrum of 3-formyl-MgPME. Signals marked with an asterisk are either solvent signals (methanol, water) or impurities (e.g., column matrix). **(C)** ¹³C-HSQC spectrum for 3-formyl-MgPME. The carbon axis is split for clarity. The dashed box indicates methyl signals, expanded on the right. The red box indicates signals from meso protons. The red circle indicates a 3¹ aldehyde signal.

Interestingly, although the fully assembled PSII complexes in the mutant were barely detectable, this strain still accumulated relatively high levels of unassembled CP43 (Figure 6A). This observation suggests a block in formation of the early PSII assembly intermediates, which precedes attachment of the CP43 module and finalization of PSII reaction center core assembly (Komenda et al., 2004).

PSII assembly occurs in a stepwise fashion from four preassembled modules. These consist of one large Chl binding subunit (D1, D2, CP47, or CP43) in addition to several low molecular mass membrane polypeptides, bound pigments and other co-factors (Komenda et al., 2012). Assembly is initiated via the association of D1 and D2 to form the intermediate complex RCII* Knoppová et al. (2014), next the CP47 assembly

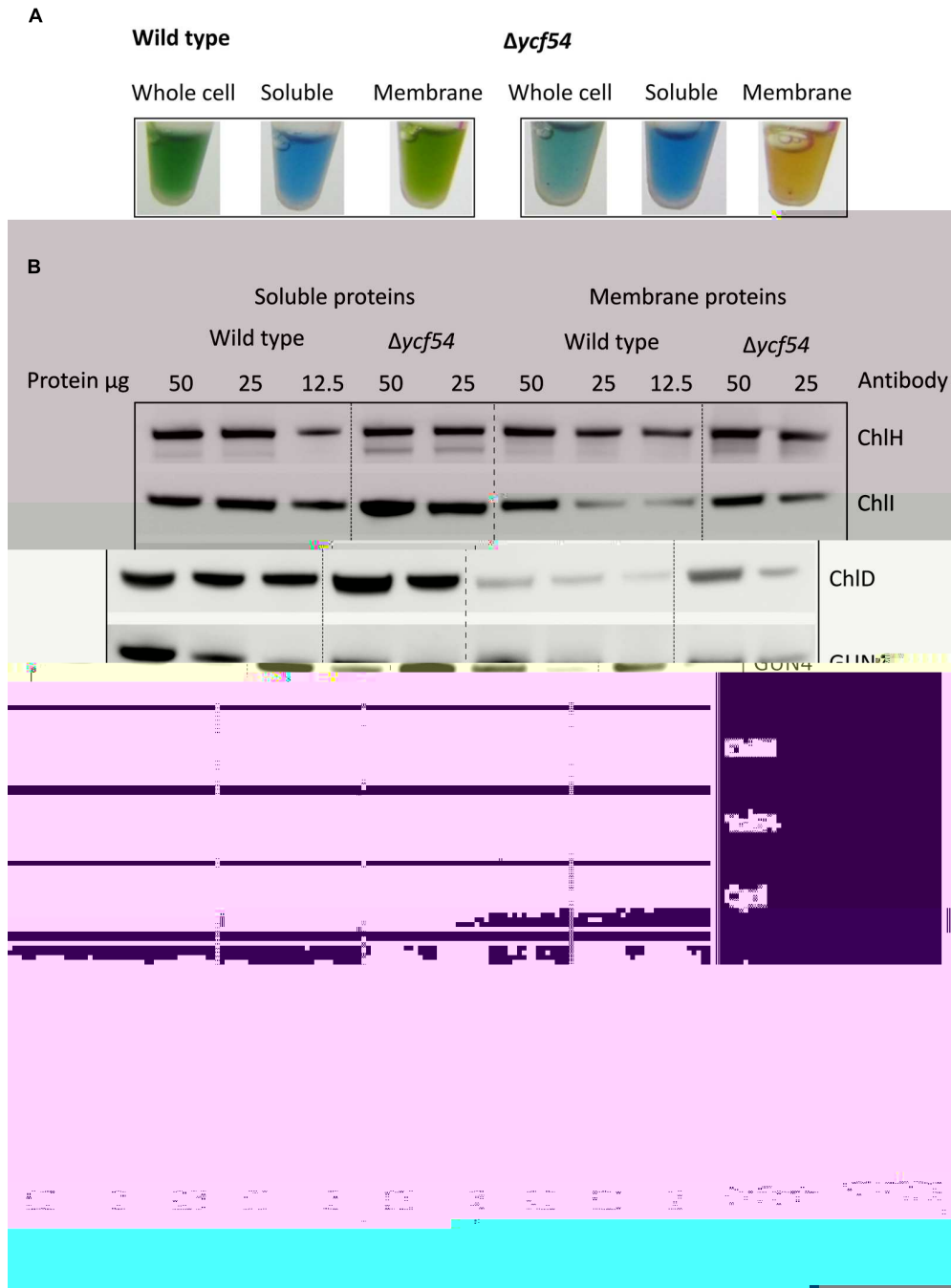
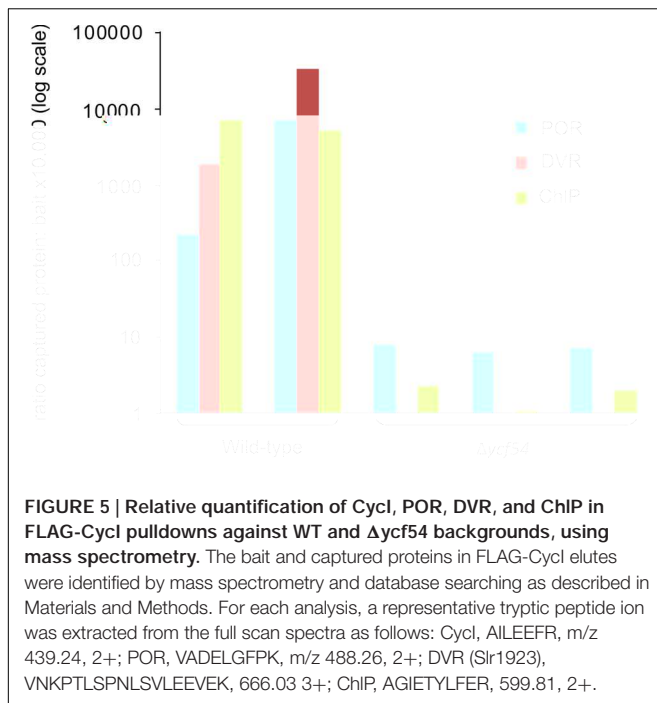


FIGURE 4 | Levels of Chl biosynthesis enzymes in WT and $\Delta ycf54$ cells. (A) Photographs of the whole cell, soluble and solubilized membrane lysate fractions, illustrating the changes in pigmentation between WT and $\Delta ycf54$. **(B)** Western blot analysis of the soluble and membrane fractions from WT and $\Delta ycf54$ *Synechocystis* strains. Samples of known protein co7 -9.465 Td (WT)-(i4ples)-traWT and.465 T3e8(Ycf54)]T 0 0 0 1 k 0 0 578(frs.)]T -by8(oteinDS-electbsence)-Absenc



accumulation of the RCII* complex, the immunoblot was probed with antibodies raised against the RCII* components D1, Ycf39, and HliD (Knoppová et al. (2014). **Figure 6B** shows that the level of RCII* is unaffected by the large reduction in cellular Chl levels in the $\Delta ycf54$ mutant. Next, we investigated if PSII maturation was blocked at CP47 attachment and formation of RC47, by probing the blots with antibodies raised against HliA, a specific component of the CP47 assembly module (Promnarek et al., 2006). We found that HliA, and hence the CP47 assembly module, was readily detectable in WT, but could not be detected in $\Delta ycf54$ (**Figure 6B**), indicating that low Chl abundance in $\Delta ycf54$ is impairing accumulation of the CP47 assembly module.

Our FLAG-pulldown experiments show that the interactions between Cycl, POR, and DVR are significantly reduced in the $\Delta ycf54$ strain (**Figure 5**), therefore we compared the co-migration of these enzymes on a 2D gel (**Figure 6C**). Evident in the WT is a putative high-mass complex of ~400 kDa (highlighted by the green box), which contains both Cycl and POR; this complex was not detectable in the mutant (**Figure 6C**). Interestingly, our 2D gel shows that levels of Chl synthase, ChlG, HliD, and Ycf39, components of a chlorophyll biosynthetic/membrane insertase assembly complex (Chidgey et al., 2014), are unaffected in the $\Delta ycf54$ mutant (**Figure 6C**).

To understand the flux of photosystem biogenesis, we used ^{35}S pulse radio-labeling coupled with 2D CN/SDS-PAGE (**Figure 7**; a Coomassie stained gel is provided as Supplementary Figure S4), to compare the levels of protein synthesis between the WT and $\Delta ycf54$ mutant. As demonstrated in **Figure 7**, the ability of $\Delta ycf54$ to synthesize the Chl-binding PSI subunits PsaA/B is limited and synthesis of CP47 and CP43 subunits is hardly detectable even though 3-times more $\Delta ycf54$ protein was loaded onto the gel (See Supplementary Figure S4 for overexposed

signal of the CP47). In contrast, there were comparable levels of synthesis of the PSII reaction center core subunits D1 and D2 in the WT and $\Delta ycf54$ strains. This observation, coupled with the data from our 2D-immunoblot (**Figure 6B**), shows that the D1 and D2 subunits are rapidly assembled into RCII* in $\Delta ycf54$, but given the lack of assembled PSII complexes, these RCII* are presumably rapidly degraded in the mutant. Interestingly, in the mutant the unassembled CP43 was still detectable on the stained gel, which contrasted to virtually zero level of unassembled CP47 (**Figures 6A,B** and **7**; Supplementary Figure S4). This observation indicates that both synthesis and stability of the CP47 are impaired in the mutant, whereas the structurally similar CP43 antenna can still accumulate though the synthesis is also very weak (Supplementary Figure S4). Taken together, our data suggest that the depleted levels of *de novo* Chl in $\Delta ycf54$ specifically hinder the synthesis of PSI and the inner antennae of the PSII. However, given different stability of CP47 and CP43, it is the lack of CP47 protein that blocks assembly of RC47 and thus PSII maturation, sensitizing the PSII assembly pathway to the availability of *de novo* Chl.

Lack of De Novo Chl Affects Ultrastructure of $\Delta ycf54$ Cells

In order to investigate the effects of removal of 87% of the cellular Chl on the ultrastructure of $\Delta ycf54$ cells, electron microscopy of negatively stained thin cell sections was performed. Electron micrographs are shown in **Figure 8**. In the WT the thylakoids are observed as parallel stacks of two to five membranes that closely follow the contour of the cell membrane (**Figures 8A–C**), but no such organized thylakoid membranes are visible in micrographs of the $\Delta ycf54$ mutant (**Figures 8B–D**). Instead, membrane-like structures are dispersed throughout the cytoplasm of the cell. These results suggest key role of photosystems in the formation of the highly ordered thylakoid structures.

DISCUSSION

The MgPME-cyclase is the least understood component in the Chl biosynthetic pathway, and current knowledge of the individual components of the MgPME-cyclase had been limited to homologs of the *Rubrivivax gelatinosus* AcsF protein. Previous work identified two genes in *Synechocystis*, *slr1214*, and *slr1874*, as *acsF* homologs, which encode a membrane associated component of the MgPME-cyclase (Minamizaki et al., 2008; Peter et al., 2009). AcsF and its homologs contain a putative di-iron site and are thus viewed as the true catalytic subunit of the MgPME-cyclase (Tottey et al., 2003). The discovery of another gene, *ycf54*, that plays an important role in cyclase activity (Albus et al., 2012; Hollingshead et al., 2012) showed that other components are required, but so far it is not possible to assign a catalytic or assembly-related function to the Ycf54 protein. Further work on the role of Ycf54 required a fully segregated $\Delta ycf54$ mutant, which is reported herein.

In our efforts to construct a fully segregated $\Delta ycf54$ mutant, we discovered it was only possible to completely delete the *ycf54* gene in one specific substrain of *Synechocystis* (GT-W).

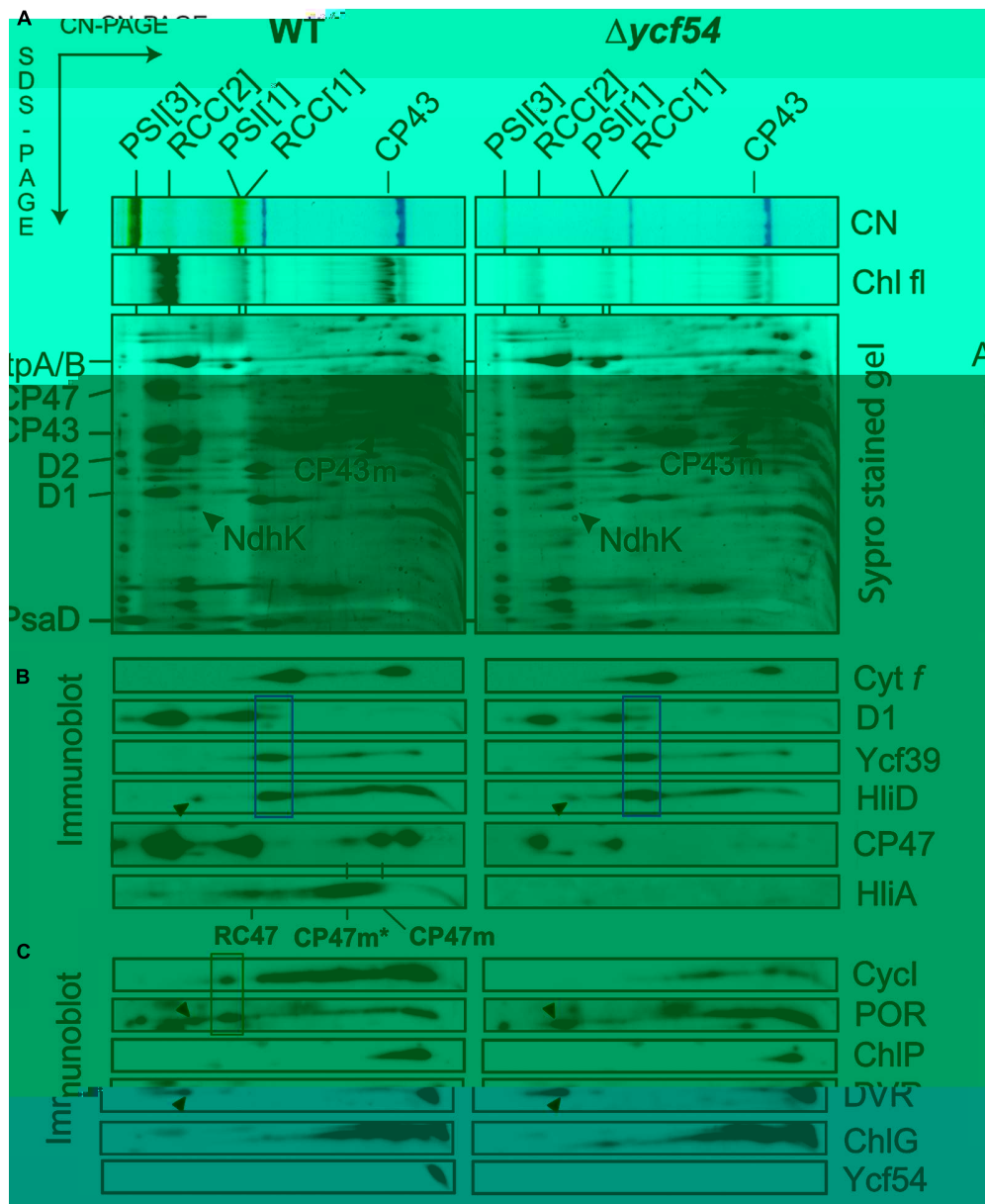
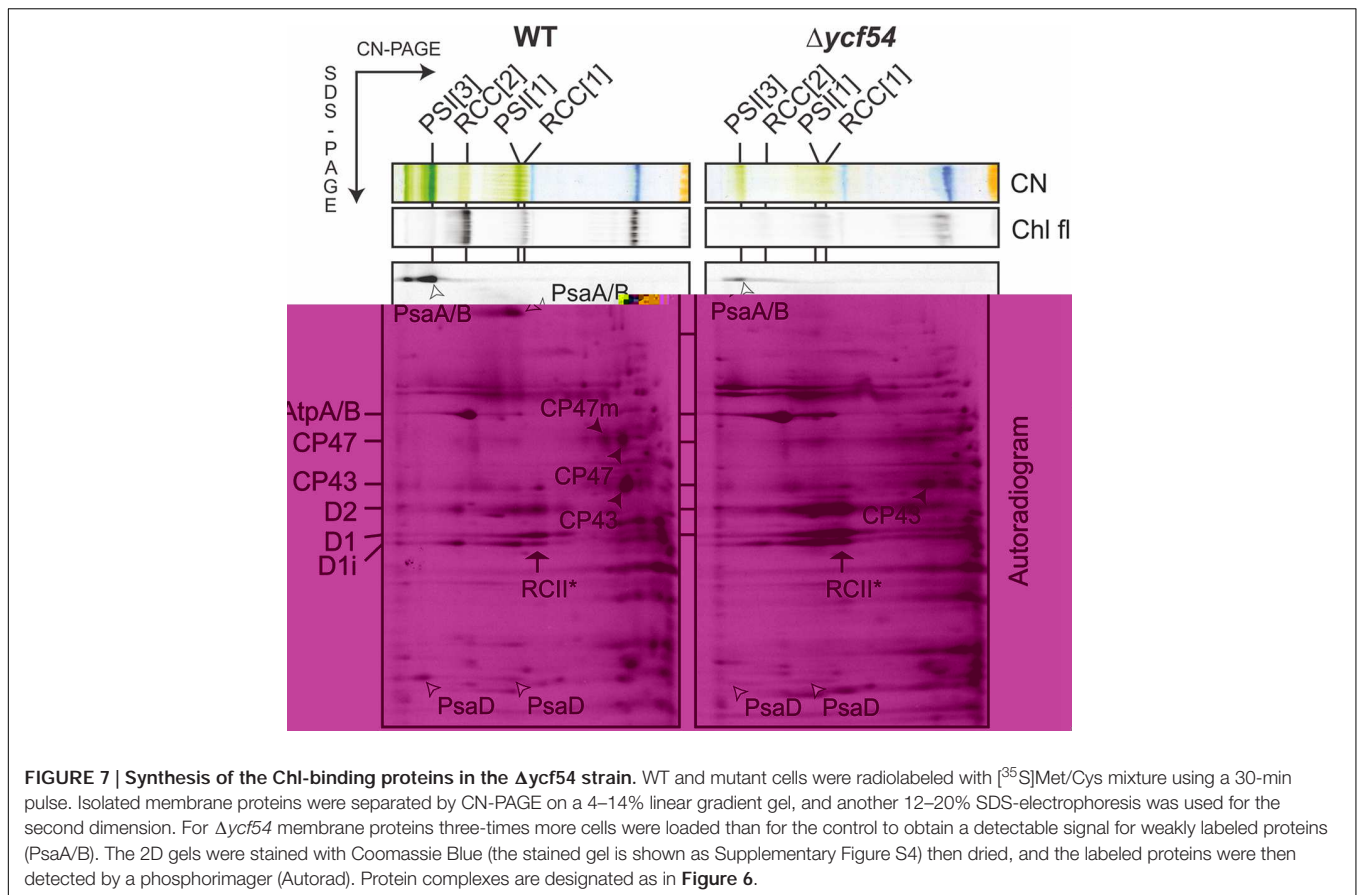


FIGURE 6 | 2D gel-electrophoresis of membrane complexes isolated from WT and $\Delta ycf54$ cells, followed by immunodetection of PSII assembly complexes and enzymes involved in Chl biosynthesis. (A) Membrane proteins were separated by 4–14% CN-PAGE, and then in the second dimension by 12–20% SDS-PAGE. The loading corresponds to the same number of cells. The SDS gel was stained by Sypro Orange, blotted and the cytochrome *f* visualized by heme *f* peroxidase activity. Chl fluorescence emitted by PSII and by unassembled CP43 (Chl fl) was detected by Fuji LAS 4000 after excitation by blue light. Designation of complexes is: PSII[3] and PSII[1], trimeric and monomeric PSII, respectively; RCC[2] and RCC[1], dimeric and monomeric PSII core complexes, respectively. **(B)** Immunodetection of PSII assembly complexes. RCII* complex (boxed in blue) was detected using D1, Ycf39, and HliD antibodies. The positions of the CP47 assembly module CP47m, and CP47m associated with High-Light-Inducible proteins HliA/B (CP47m*), are indicated. The HliA signal also marks the position of the PSII core complex lacking CP43 (RC47). **(C)** Immunodetection of Ycf54 proteins and enzymes involved in Chl biosynthesis in the membrane fraction separated by 2D electrophoresis. Highlighted by the green box is a putative high-mass complex (~400 kDa) containing Cycl and POR; this complex is not detectable in the mutant. Orange triangles indicate unspecific cross-reactions with a subunit of the NDH complex. ChlG is Chl-synthase; other enzymes are designated as in Figure 2.

One possible explanation for this finding was elucidated by analysis of the GT-W genome, which contains a long (~100 kbp) chromosomal duplication that covers one hundred genes, including *cycl* (*sll1214*; Bečková et al., submitted). This

chromosomal duplication is not present in any of the other *Synechocystis* substrains for which a genome sequence is available (Kanesaki et al., 2012; Trautmann et al., 2012). Given the duplication of *cycl* in GT-W, it is likely that *cycl* expression is



increased in this strain. A very low level of *CycI* is a hallmark of the strains in which we partially or completely inactivated the *ycf54* gene. Thus, we hypothesize that the doubled expression of the *slI1214* gene coding for *CycI* may suppress the lethality of inactivating the *ycf54* gene. This hypothesis is in agreement with our observation that in the absence of *Ycf54* the *CycI* cyclase component is destabilized (**Figures 4 and 5**) and the remaining *CycI* content is probably very close to a threshold essential for viability.

Analysis of the pigments that accumulate in the $\Delta ycf54$ mutant could provide clues regarding the role of the *Ycf54* protein, and a previous analysis showed that the partially segregated *ycf54* mutant accumulates MgPME, the substrate of the cyclase. In addition, there was an unknown pigment (**Figure 2**) that was suggested to be an intermediate in the cyclase reaction (Hollingshead et al., 2012), on the basis that the 433 nm Soret absorbance peak falls between the Soret peaks of the cyclase substrate MgPME (416 nm) and the PChlide product (440 nm). Similarly, early work with greening cucumber cotyledons had found pigments with emission maxima between 434 and 436 nm, proposed to be biosynthetic intermediates between MgPME and PChlide (Rebeiz et al., 1975). Identification of the unknown pigment as Mg-3-formyl-PME suggests that this pigment is not an intermediate in the cyclase reaction, as it is highly unlikely that this would produce a Chl pigment modified at the C3 position. Rather, Mg-3-formyl-PME bears a striking resemblance

to Chl *d*, the major light harvesting pigment found in the cyanobacterium *Acaryochloris marina* (Miyashita et al., 1996). The pathway and reaction mechanism of Chl *d* biosynthesis in *Acaryochloris marina* have not yet been elucidated but, based on the genome sequence, Chl *d* is thought to be derived from Chl *a* (Swingley et al., 2008). Previously, Chl *d* has been synthesized in low yields in aqueous acetone from Chl *a* by treatment with papain (Koizumi et al., 2005) and peroxide (Aoki et al., 2011) and in much higher yields from Chl *a* incubated with thiophenol and acetic acid in tetrahydrofuran (Fukusumi et al., 2012). Given the high accumulation of MgPME in $\Delta ycf54$, it is likely that reactive oxygen species, including peroxide, convert the MgPME 3-vinyl group, leading to the formation of Mg-3-formyl-PME.

The identification of Mg-3-formyl-PME as an oxidation product of the substrate rather than a catalytic intermediate is not consistent with a catalytic role for *Ycf54* in the MgPME-cyclase complex. However, the hypothesis by Hollingshead et al. (2012) that *Ycf54* plays a role in the assembly or stabilization of the catalytic MgPME-cyclase enzyme complex remains valid. By using FLAG-*CycI* as bait in pulldown assays, combined with quantitative MS analysis, we demonstrated that the absence of *Ycf54* affects formation of a complex between *CycI* and enzymes further down the pathway (POR, DVR, and ChlP). In particular, the almost complete absence of DVR in the pulldown from the $\Delta ycf54$ strain provides a strong evidence that *Ycf54* facilitates formation of such a complex; in contrast to POR and ChlP the



FIGURE 8 | Transmission electron micrographs of WT and $\Delta ycf54$ cells. Ultrathin sections from *Synechocystis* WT (A–C) and $\Delta ycf54$ cells (D–F) grown photomixotrophically under low light conditions. White arrows indicate thylakoid membranes and white triangles indicate glycogen granules.

stability of DVR in the mutant does not seem to be compromised and thus this result cannot be explained by a hypothetical fast degradation of this enzyme during pull-down assay. Indeed, the present data also provide evidence for an interaction between Cycl1 and POR, DVR, and ChIP, which aligns well with data obtained by Kauss et al. (2012) who performed pull-down experiments with the *Arabidopsis* FLU protein. These analyses found that FLU forms a complex with CHL27, the *Arabidopsis* AcsF homolog, PORB, PORC, and ChIP. In *Synechocystis* at least, it appears that Ycf54 plays no direct catalytic role, and that it is important for the formation and maintenance of a Chl biosynthetic complex, with disruption of this complex possibly triggering degradation of Cycl and consequently Chl deficiency. However, a wider role for Ycf54 in governing the whole pathway appears to be excluded by the lack of effect of the $\Delta ycf54$ on components of the ChIG-HliD-Ycf39 complex that operates at the end of the pathway. This complex likely coordinates Chl delivery to the membrane-intrinsic apparatus for insertion and translocation of apoproteins of the photosynthetic apparatus (Chidgey et al., 2014). It is notable that although the Cycl is almost exclusively associated with the membrane fraction under moderate light conditions Ycf54 is distributed equally between membrane and cytosol under

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modules appears to be more complex, as previous studies show Chl molecules are re-cycled during PSII synthesis and repair (Kopečná et al., 2013, 2015) via re-phytylation of chlorophyllide (Vavilin and Vermaas, 2007). The PSII complex assembles in a modular fashion, starting with the association of D1 and D2 assembly modules, to form the RCII* complex. This is followed by attachment of a CP47 module, then a CP43 module, then the luminal extrinsic proteins and the oxygen-evolving Mn₄CaO₅ complex (reviewed in Komenda et al., 2012). Despite the large decrease in cellular Chl levels in $\Delta ycf54$, all components of the RCII* complex are synthesized in adequate amounts and assembled. We hypothesize that synthesis of the RCII* complex is enabled by the continuous recycling of a relatively stable pool of Chl molecules made available during the RCII* assembly/degradation cycle. Evidence that RCII* contains Chl as well pheophytin, carotenoids and heme cofactors has been shown previously (Knoppová et al., 2014). We cannot exclude the possibility that in the $\Delta ycf54$ mutant there is a pool of the RCII* complex that lacks Chl. However, we did not observe any shift of electrophoretic mobility even for the ³⁵S labeled RCII* that would indicate presence of a hypothetical Chl-less RCII*. Furthermore, $\Delta ycf54$ contains some functional PSII complexes, which requires that at least some RCII* with cofactors has to be synthesized en route to the fully assembled PSII.

Our findings also show that CP43 can accumulate as an unassembled module in $\Delta ycf54$ even though the synthesis is very limited (Figures 6A and 7). In contrast, CP47 seems to be unstable in $\Delta ycf54$, which suggests that CP47 is the *de novo* Chl sensitive component of PSII biogenesis. This observation is consistent with previous work on the accumulation of PSII subunits in Δpor (Kopečná et al., 2013) and $\Delta gun4$ (Sobotka et al., 2008) mutants, disrupted in the PChlide reduction and Mg-chelatase steps, respectively. As also seen for $\Delta ycf54$, the Δpor and $\Delta gun4$ strains accumulate the PSII core complex RCII* and the PSII antenna CP43, but CP47 synthesis is not observed (Sobotka et al., 2008; Kopečná et al., 2013). It is not currently known why CP47 is more sensitive to the availability of *de novo* Chl than the similar CP43 subunit, although it has been recently observed that the newly synthesized CP43, but not CP47, subunit is attached to a PSI complex (Kopečná et al., 2015). We tentatively speculate that the situation in the mutant leads frequently to the synthesis of aberrant CP47 lacking one or more Chl molecules. The synthesis of CP43 might be less error-prone because Chl molecules bound to the periphery of PSI could be used for the assembly of this complex.

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In summary, the role of Ycf54 in the MgPME-cyclase complex has been elucidated further. This work shows that whilst Ycf54 is required for stabilization of Cyc1, the known catalytic component of the MgPME-cyclase, the protein itself is unlikely to play a key catalytic role in the formation of the fifth isocyclic ring. Furthermore, Ycf54 does not appear to be directly implicated in Chl phytolation or Chl insertion into proteins. The construction of a $\Delta ycf54$ mutant has provided a useful tool to investigate the effects of reduced *de novo* Chl on the biosynthesis of cyanobacterial Chl binding proteins, highlighting the differing requirements for Chl exhibited by proteins within the PSI and PSII light harvesting complexes that bind this pigment. Insights into the catalytic cycle of the MgPME-cyclase remain elusive and further work is required to determine the exact molecular mechanisms of this enzyme.

AUTHOR CONTRIBUTIONS

SH, JK, DA, LB, PJ, and GC performed the research; MD, MW, RS, and CNH designed the experiments, and SH, DA, PJ, MD, MW, RS, and CNH wrote the paper.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: <http://journal.frontiersin.org/article/10.3389/fpls.2016.00292>

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