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Evidence for a Role of VIPP1 in the Structural Organization of the Photosynthetic Apparatus in *Chlamydomonas*

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The vesicle-inducing protein in plastids (VIPP1) was suggested to play a role in thylakoid membrane formation via membrane vesicles. As this functional assignment is under debate, we investigated the function of VIPP1 in *Chlamydomonas reinhardtii*. Using immunofluorescence, we localized VIPP1 to distinct spots within the chloroplast. In VIPP1-RNA interference/artificial microRNA cells, we consistently observed aberrant, prolamellar body-like structures at the origin of multiple thylakoid membrane layers, which appear to coincide with the immunofluorescent VIPP1 spots and suggest a defect in thylakoid membrane biogenesis. Accordingly, using quantitative shotgun proteomics, we found that unstressed *vipp1* mutant cells accumulate 14 to 20% less photosystems, cytochrome *b₆f* complex, and ATP synthase but 30% more light-harvesting complex II than control cells, while complex assembly, thylakoid membrane ultrastructure, and bulk lipid composition appeared unaltered. Photosystems in *vipp1* mutants are sensitive to high light, which coincides with a lowered midpoint potential of the Q_A/Q_A^- redox couple and increased thermosensitivity of photosystem II (PSII), suggesting structural defects in PSII. Moreover, swollen thylakoids, despite reduced membrane energization, in *vipp1* mutants grown on ammonium suggest defects in the supermolecular organization of thylakoid membrane complexes. Overall, our data suggest a role of VIPP1 in the biogenesis/assembly of thylakoid membrane core complexes, most likely by supplying structural lipids.


INTRODUCTION


The thylakoids of chloroplasts represent an internal membrane system that is detached from the inner envelope membrane. As thylakoid lipids are not synthesized at the thylakoid membranes but rather at the chloroplast inner and outer envelope membranes and the endoplasmic reticulum, a transport system must exist that enables a flow of lipid components from these biogenic membranes to the thylakoids (Benning, 2008, 2009). Ultrastructural studies suggested lipid transport via vesicles that bud off

from the inner envelope and fuse with the thylakoids (Carde et al., 1982). The M30 protein was identified in pea (*Pisum sativum*) as a plastid protein that is associated with both envelope and thylakoid membranes and therefore was suggested as a candidate protein for lipid transfer from envelope to thylakoids (Li et al., 1994). M30 was renamed vesicle-inducing protein in plastids1 (VIPP1) based on the phenotype observed in the *Arabidopsis thaliana hcf155* mutant, which expresses M30/VIPP1 to ~20% of wild-type levels: *hcf155* plants have much less and distorted thylakoids and reduced amounts of photosystem I (PSI), photosystem II (PSII), light-harvesting complex B, cytochrome *b₆f* complex, and ATP synthase compared with wild-type plants. Moreover, *hcf155* plants lack vesicles originating from the inner chloroplast envelope (Kroll et al., 2001; Aseeva et al., 2007). These observations led to the proposal that VIPP1 is essential for the formation of thylakoid membranes via vesicle traffic, a conclusion that was supported by the almost complete lack of

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thylakoids in a cyanobacterial *vipp1* mutant strain (Westphal et al., 2001). However, in a similarly constructed cyanobacterial *vipp1* mutant strain, Fuhrmann et al. (2009a) only found reduced, less well arranged thylakoid layers and reduced amounts of (trimeric) PSI. Because VIPP1 is an essential protein, the disruption of the *VIPP1* gene generated merodiploid cells that still accumulated >25% of wild-type VIPP1 levels. Hence, Gao and Xu (2009) generated a cyanobacterial strain expressing *VIPP1* under control of the copper-responsive *petE* promoter and under copper-depleted conditions observed that depletion of VIPP1 correlated first with a loss of photosynthetic activity (in particular of PSII) before thylakoid membranes were depleted. Therefore, Gao and Xu questioned the role of VIPP1 in thylakoid formation.

The picture becomes even more confusing when looking at the proposed function for the closest homolog of VIPP1 in prokaryotes, the phage shock protein A (PspA) (Joly et al., 2010). The phage shock response is induced by agents that potentially affect the integrity of the plasma membrane and normally lead to a loss of the proton motive force. Examples for inducing agents are filamentous phage infection, severe heat shock, depletion of the protein membrane insertase YidC, or blockage of the twin-Arg (TAT) or Sec translocons (Brissette et al., 1990; Kleerebezem and Tommassen, 1993; van der Laan et al., 2003; DeLisa et al., 2004). PspA in its oligomeric form was shown to suppress proton leakage from damaged membranes by directly interacting with membrane lipids phosphatidylserine and phosphatidylglycerol (PG; Kleerebezem et al., 1996; Kobayashi et al., 2007). Functional conservation between PspA and VIPP1 is suggested by the findings that both proteins improved protein export via the twin-Arg translocon pathway (DeLisa et al., 2004), and both proteins assemble into rotationally symmetric rings of >1 MD (Aseeva et al., 2004; Hankamer et al., 2004; Liu et al., 2007; Standar et al., 2008; Fuhrmann et al., 2009b). Still, some specificity for PspA and VIPP1 function must exist because both are present in cyanobacteria, but cyanobacterial PspA cannot substitute for the function of cyanobacterial VIPP1 (Westphal et al., 2001).

Several opinions exist not only regarding the function of VIPP1 but also concerning its localization. In chloroplasts of higher plants and algae, VIPP1 was localized to thylakoids and the inner envelope (Li et al., 1994; Kroll et al., 2001; Liu et al., 2005), but a localization only to inner envelopes was proposed by Aseeva et al. (2004). Moreover, VIPP1 was also found in stromal fractions (Li et al., 1994; Liu et al., 2005). In cyanobacteria, VIPP1 was initially reported to be localized exclusively to the plasma membrane (Westphal et al., 2001). However, a dual localization of VIPP1 to plasma membrane and thylakoids was reported (Srivastava et al., 2005), and, more recently, VIPP1 was also detected in the cytoplasm (Srivastava et al., 2006; Fuhrmann et al., 2009b). Eventually, these data suggest that VIPP1 is in an equilibrium between membrane-bound and soluble forms, as appears to be the case for bacterial PspA (Brissette et al., 1990; Kleerebezem and Tommassen, 1993).

VIPP1 activity appears to be tightly linked to plastidic chaperones, especially the HSP70 and HSP90 systems (Nordhues et al., 2010). VIPP1 interacts with a specialized J-domain protein termed CDJ2, which delivers VIPP1 to chloroplast HSP70B (Liu et al., 2005). HSP70B and CDJ2 catalyze the assembly and disassembly of VIPP1 oligomers (Liu et al., 2007). The interaction

of VIPP1 with chloroplast HSP70 was confirmed in higher plants (Aseeva et al., 2007), and VIPP1 was found to interact also with chloroplast HSP90, which might support HSP70 during VIPP1 oligomer assembly/disassembly (Heide et al., 2009). To elucidate why the chloroplast HSP90/HSP70 systems catalyze the assembly/disassembly of VIPP1 oligomers, it is essential to understand the function of VIPP1. However, as outlined above, the functional analyses done so far in higher plants and cyanobacteria did not yet provide definitive conclusions on VIPP1 function. We therefore decided to perform a functional analysis of VIPP1 in the unicellular green alga *Chlamydomonas reinhardtii*, in which the interaction between VIPP1 and the chaperones was discovered. While doing so, we found that, unlike higher plants and cyanobacteria, volvocean algae contain two VIPP paralogs. In *C. reinhardtii*, these two paralogs share 50% identical and 65% similar residues and were termed VIPP1 and VIPP2 (S. Schmollinger and M. Schroda, unpublished results). Although the presence of two VIPP paralogs complicated the analysis, our results strongly suggest that these VIPPs are redundantly involved in the biogenesis/assembly of thylakoid membrane protein complexes at distinct sites within the chloroplast and not in vesicle-mediated thylakoid membrane formation as proposed earlier.

RESULTS

C. reinhardtii VIPP1 Is Efficiently Downregulated by RNA Interference

To investigate the function of VIPP1 in *C. reinhardtii*, we first used an RNA interference (RNAi) strategy to reduce *VIPP1* function. To do this, we made a construct harboring a piece of genomic, intron-containing *VIPP1* DNA in sense orientation followed by a complementary piece of cDNA in antisense orientation (see Supplemental Figure 1 online). Expression of the RNAi construct is driven by the strong *HSP70A-RBCS2* fusion promoter (Schroda et al., 2000). The *ARG7* gene present on the same vector was used as selectable marker for the transformation of the Arg-auxotrophic cw15-325 recipient strain. This construct silenced the *VIPP1* gene efficiently, as shown by the reduction of VIPP1 protein to <5% of wild-type levels in ~10% of the Arg-prototrophic transformants (Figure 1A; see Supplemental Figure 4 online). However, as wild-type VIPP1 levels recovered within 1 to 6 months in *VIPP1*-RNAi strains the RNAi construct itself appears to be silenced, as observed earlier with other inverted repeat constructs (Yamasaki et al., 2008). Therefore, during the course of this work, we regularly had to generate fresh *VIPP1*-RNAi strains. Eventually, >30 independent lines were characterized. Control strains were generated by transforming the cw15-325 strain with a plasmid containing the *ARG7* gene alone.

VIPP1-RNAi/amiRNA Strains Are Sensitive to High Light

Despite the strong reduction of VIPP1 levels in *VIPP1*-RNAi strains (Figure 1A), we observed no obvious phenotypes in cells grown at low light intensities in Tris-acetate-phosphate (TAP)-NH₄ medium: Cells grew normally under mixotrophic conditions

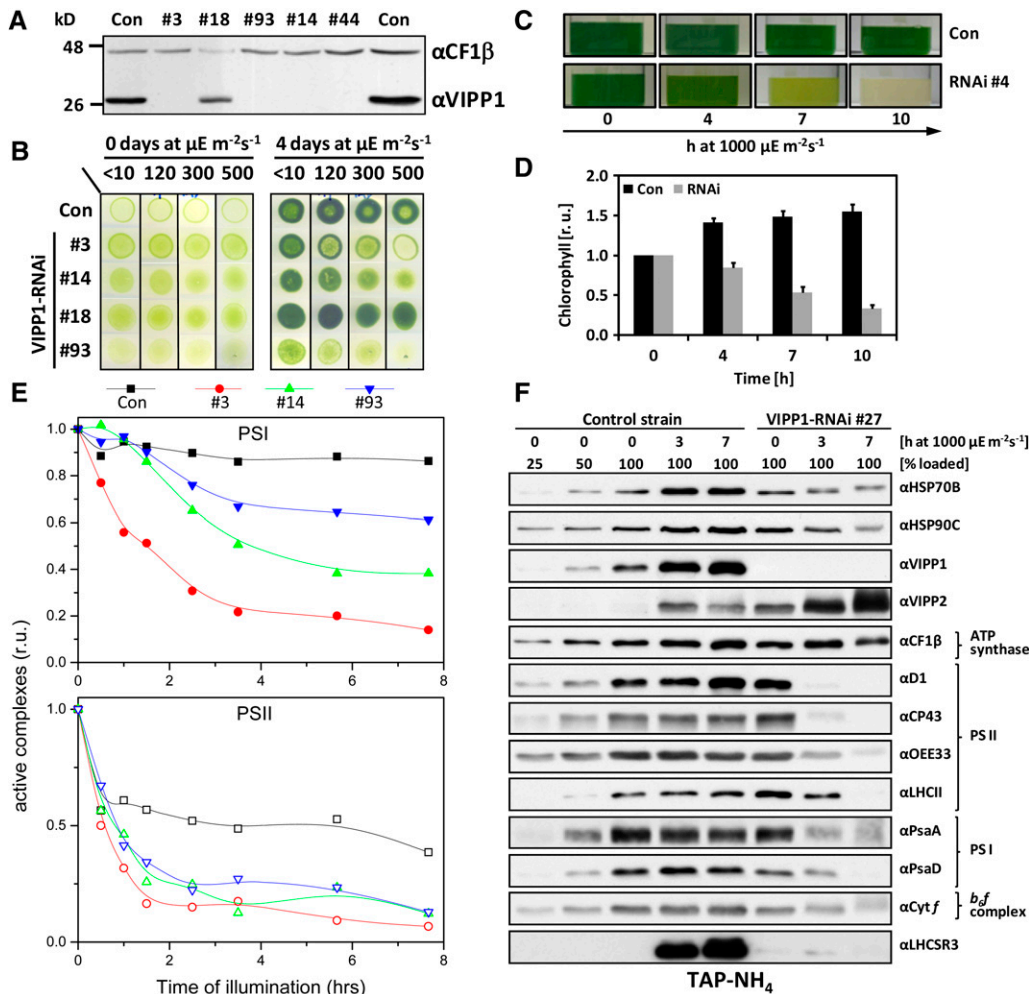


Figure 1. Photobleaching of High Light-Exposed *VIPP1*-RNAi Strains.

(A) *VIPP1* protein levels in *VIPP1*-RNAi strains are strongly reduced. Whole-cell proteins from transformants generated with *VIPP1*-RNAi construct pMS439 corresponding to 2 μg chlorophyll were separated on a 14% SDS-polyacrylamide gel and analyzed by immunoblotting. CF1 β served as loading control.

(B) High light sensitivity of *VIPP1*-RNAi strains. Ten microliters containing 10^7 cells were spotted onto TAP-NH₄ agar plates and photographed directly or after a 4-d incubation at the indicated light intensities.

(C) Bleaching of *VIPP1*-RNAi strains. A control strain and *VIPP1*-RNAi strain #4 grown in TAP-NH₄ medium were exposed to a light intensity of $\sim 1000 \mu\text{E m}^{-2} \text{s}^{-1}$ for 10 h.

(D) The chlorophyll content rapidly declines in high light-exposed *VIPP1*-RNAi strains. A control strain ($n = 3$) and five *VIPP1*-RNAi strains (#3, #26, #44, #49, and #93) were treated as in **(A)**, and the chlorophyll content was determined. Chlorophyll contents are given relative to the concentrations determined prior to the shift to high light, which were set to 1. Error bars represent SE. r.u., relative units.

(E) PSII and PSI in *VIPP1*-RNAi strains are very sensitive to high light. Equal cell densities of a control strain and *VIPP1*-RNAi strains #3, #14, and #93 were subjected to high light as in **(A)**, and PSII and PSI activities were measured by determining the DCMU-sensitive and -insensitive fractions of the electrochromic shift signal.

(F) Subunits of PSII and PSI are rapidly degraded in *VIPP1*-RNAi strains exposed to high light intensities. Whole-cell proteins from high light-exposed control and *VIPP1*-RNAi strain #27 grown in TAP-NH₄ medium were separated on 14% SDS-polyacrylamide gels and analyzed by immunoblotting.

(Figure 1B), had fully developed thylakoid membranes (see Figures 3C and 10A), and seemed to assemble thylakoid membrane protein complexes like control cells (see Supplemental Figure 2 online). The composition of bulk thylakoid membrane lipids in *VIPP1*-RNAi strains was indistinguishable from control cells (see Supplemental Figure 3 online), and we could not

observe consistent changes in the number of thylakoids per granum stack (as judged from the analysis of electron micrographs from 50 cells each of two control and five *VIPP1*-RNAi lines). This observation was unexpected, as *Arabidopsis vipp1* mutants expressing *VIPP1* at $\sim 20\%$ of wild-type levels have an albinotic phenotype and cannot grow photoautotrophically due

to a degraded thylakoid membrane system (Kroll et al., 2001; Aseeva et al., 2007).

In previous work, we observed that the *C. reinhardtii* *VIPP1* gene was strongly induced when dark-grown cells were shifted into the light (Liu et al., 2005). In *C. reinhardtii*, the thylakoid membranes are fully developed also in the dark; this suggested to us that *VIPP1* may be of particular importance in the light. We therefore exposed control and *VIPP1*-RNAi strains to increasing light intensities (up to $500 \mu\text{E m}^{-2} \text{s}^{-1}$) and observed that *VIPP1*-RNAi strains containing very low levels of *VIPP1* failed to grow and bleached at higher light intensities. By contrast, no effect of high light was observed with the control strain and with a *VIPP1*-RNAi strain expressing *VIPP1* to $\sim 50\%$ of wild-type levels (#18; Figure 1B).

To substantiate this finding, we exposed mixotrophically grown control and *VIPP1*-RNAi strains to high light intensities ($\sim 1000 \mu\text{E m}^{-2} \text{s}^{-1}$) (Figure 1C). Within 10 h, we observed severe bleaching of *VIPP1*-RNAi strains. On average, the chlorophyll content declined to $\sim 40\%$ of starting levels, while in the control strain, it increased by $\sim 60\%$ during high light treatment as a consequence of growth (Figure 1D).

Assessment of PSII and PSI activities by measurements of the electrochromic shift revealed a fast decline of PSII activity in high light-treated *VIPP1*-RNAi strains, with activities dropping to ~ 15 to 35% of the initial values already after 90 min and to $\sim 10\%$ after 7.5 h at high light (Figure 1E). In the control, PSII activity dropped only to $\sim 50\%$ of the initial value. Also, PSI activities declined in high light-treated *VIPP1*-RNAi strains but generally not as fast as PSII activities. The 7.5-h high light treatment resulted only in an $\sim 10\%$ drop of PSI activity in the control, but the loss of PSI activity in the *VIPP1*-RNAi strains ranged between ~ 40 and $\sim 90\%$ and correlated with the residual *VIPP1* levels present in these strains (Figure 1E; see Supplemental Figure 4 online).

Protein gel blot analyses revealed that the D1, CP43, and OEE33 subunits of PSII were reduced by $>75\%$ after 3 h at high light (Figure 1F). Interestingly, light-harvesting complex II (LHCII) levels in *VIPP1*-RNAi cells grown under low light conditions appeared to be higher than in control cells, and it took 7 h at high light until LHCII levels were reduced by $>50\%$. A reduction of PSI core subunits PsaA and PsaD in *VIPP1*-RNAi cells by more than half was observed only after 7 h at high light and therefore was not as fast as the loss of the PSII core subunits, which is in line with the spectroscopic data. Levels of CF1 β and cytochrome *f*, subunits of the ATP synthase and cytochrome *b₆f* complex, respectively, were barely reduced in high light-exposed *VIPP1*-RNAi strains. No decline in any of the investigated subunits was observed in high light-treated control cells.

VIPP1 and *VIPP2* were expressed constitutively in control cells grown at low light intensities, and levels of both proteins increased strongly during exposure to high light (Figure 1F). While high light treatment led to an increase in levels of plastidic chaperones HSP70B and HSP90C and of the LHCSR3 protein in control cells, none of these proteins increased in high light-treated *VIPP1*-RNAi strains. As the genes coding for these three proteins are nuclear encoded, this finding suggests that *VIPP1*-RNAi strains might be defective in a retrograde high light signaling pathway. The existence of more than one high light signaling

pathway is implied by the observation that *VIPP1*-RNAi strains are still capable of inducing *VIPP2* expression in response to high light exposure.

Interestingly, already under low light conditions, *VIPP1*-RNAi cells expressed *VIPP2* at much higher levels than control cells, and they increased *VIPP2* expression further during exposure to high light intensities (Figure 1F). Apparently, *VIPP1*-RNAi cells try to compensate for the lack of *VIPP1* by expressing additional *VIPP2*, but *VIPP2* cannot completely substitute *VIPP1*. This may have two reasons, which are not mutually exclusive: First, *VIPP2* may be functionally redundant with *VIPP1*, but its expression levels may be too low to completely compensate for the *VIPP1* loss in *VIPP1*-RNAi strains. Second, *VIPP2* overexpression in *VIPP1*-RNAi strains may replenish the total *VIPP* pool, but *VIPP2* function may be distinct from that of *VIPP1* and, therefore, *VIPP2* cannot substitute *VIPP1*. To distinguish between these possibilities, we needed to get an estimate of the relative cellular expression levels of *VIPP1* and *VIPP2*. For this, we separated total cell proteins next to dilution series of purified, recombinant *VIPP1* and *VIPP2* and immunodetected *VIPP1* and *VIPP2* with specific antisera. Exposure times were chosen such that signals from recombinant *VIPP1* and *VIPP2* were about equal, thus allowing for a direct comparison of *VIPP1* and *VIPP2* signals from total cell proteins (Figure 2). This procedure allowed us to estimate that the total *VIPP* pool size in *VIPP1*-RNAi cells of the cw15-325 background was $\sim 25\%$ of that in control cells, with a *VIPP1*:*VIPP2* ratio of $\sim 1:4$.

Although these data support the first scenario, we could not exclude the second one as the two *VIPPs* are expressed at strongly different ratios in *VIPP1*-RNAi cells compared with control cells. We therefore decided to repress *VIPP1* in a strain background (cw15-302) that we knew to be unable to increase *VIPP2* expression levels in response to *VIPP1* repression. For this, we used an artificial microRNA (amiRNA) approach to also rule out off-target effects potentially caused by the inverted

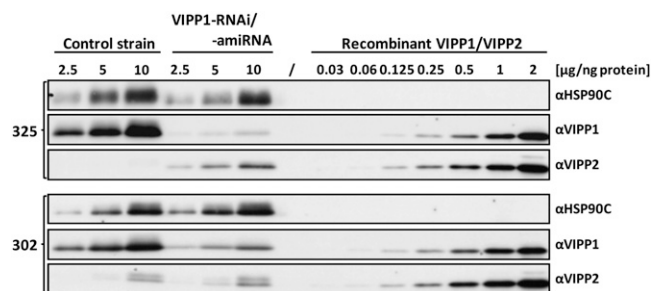


Figure 2. Examination of *VIPP1/2* Levels in *VIPP1*-RNAi and -amiRNA Strains by Immunoblotting.

Transformants generated with empty vectors (control strains) or *VIPP1*-RNAi and *VIPP1*-amiRNA constructs in the cw15-325 (#12) and cw15-302 (#18) strain backgrounds, respectively, were grown in TAP-NH₄ medium. Total protein from the transformants and purified recombinant *VIPP1* and *VIPP2* at the indicated protein concentrations were separated on 7.5 to 15% SDS-polyacrylamide gels and analyzed by immunoblotting using antisera against *VIPP1* and *VIPP2*. HSP90C served as loading control.

repeat construct (Molnar et al., 2009; see Supplemental Figure 5A online). As shown in Figure 2, VIPP2 in the cw15-302 strain background is expressed to higher levels than in the cw15-325 background and VIPP2 expression is not elevated in response to VIPP1 repression. However, VIPP1 in the cw15-302 background in several independent transformants could be repressed to only ~25% of wild-type VIPP1 levels such that the ratio of VIPP1:VIPP2 was ~2:1. Exposure of *VIPP1*-amiRNA strains to high light intensities again resulted in photobleaching (see Supplemental Figure 5B online). These results suggested that VIPP1 and VIPP2 are largely functionally redundant and that high light sensitivity is caused by a reduction in plastidic VIPP levels independent of its composition. Moreover, high light sensitivity apparently is not caused by off-target effects of the RNAi construct.

Because of the proposed link between VIPPs and chloroplast biogenesis, we investigated the effect of high light treatment on thylakoid membrane ultrastructure in *VIPP1*-depleted strains. To this end, we used transmission electron microscopy on control and *VIPP1*-amiRNA strains prior to and after a 3.5-h exposure to high light intensities. While the high light treatment hardly affected thylakoid structure in control cells, thylakoids in most cells of the *VIPP1*-amiRNA strains were extremely swollen (Figure 3).

In summary, *VIPP1* is the major VIPP in *C. reinhardtii*. Apparently, the total VIPP pool cannot be constitutively repressed to below ~25% of wild-type levels. Whereas *VIPP1*-RNAi/amiRNA strains showed no obvious phenotypes under low light conditions, they turned out to be highly sensitive to high light, as manifested by chlorophyll bleaching, loss of photosystem activities, and thylakoid swelling. PSII was most sensitive in *VIPP1*-RNAi strains, closely followed by PSI, while ATP synthase and cytochrome *b₆f* complex were only mildly affected, if at all. Finally, high light-induced accumulation of the HSP90C and HSP70B chaperones and of LHCSR3 normally observed in control cells was abolished in *VIPP1*-RNAi cells.

***VIPP1*-RNAi Cells Are More Severely Photoinhibited and Repair Photodamaged PSII More Slowly Than Control Cells**

The observed sensitivity of *VIPP1*-RNAi/amiRNA strains to high light raises two questions: First, is it a secondary effect that occurs only after prolonged exposure to high light (e.g., as a consequence of thylakoid swelling)? Second, is the fast loss of PSII activity and core subunits during high light treatment caused by the inability of *VIPP1*-RNAi strains to repair photodamaged D1 protein? To answer these questions, we exposed control and *VIPP1*-RNAi strains to ~1800 $\mu\text{E m}^{-2} \text{s}^{-1}$ for 30 min in the presence and absence of chloramphenicol, an inhibitor of organellar protein biosynthesis, and monitored PSII recovery from photoinhibition at low light intensities. As shown in Figure 4, loss of PSII maximum quantum efficiency was more severe in *VIPP1*-RNAi cells than in control cells and its recovery, both in the presence and absence of chloramphenicol, was faster in control than in *VIPP1*-RNAi strains. PSII recovery in the presence of chloramphenicol is attributable to reversibly photodamaged reaction centers, whereas the additional PSII recovery in the absence of chloramphenicol is derived from PSII centers from which irreversibly damaged D1 was exchanged by de novo synthesized D1 (Ohad et al., 1990; Schroda et al., 1999). In

VIPP1-RNAi strains, we observed that additional PSII was recovered in cells carrying out organellar protein biosynthesis compared with cells lacking organellar protein biosynthesis. We conclude that, compared with the control, *VIPP1*-RNAi strains are also more sensitive to a short application of very high intensity light. *VIPP1*-RNAi strains are generally capable of repairing photodamaged D1, but the protein biosynthesis-dependent repair rate of PSII is slower in *VIPP1*-RNAi strains than in controls.

PSII repair requires the movement of photodamaged PSII from grana stacks to stroma-exposed thylakoid regions (Goral et al., 2010). Another process requiring extensive movement of protein complexes within the thylakoid membrane is state transitions. This process is triggered by an imbalance in excitation energy between the photosystems and leads, upon overexcitation of PSII, to the phosphorylation of LHCII by the STT7 kinase and the movement of LHCII to PSI (Depège et al., 2003; Bellafiore et al., 2005). The remigration of LHCII in case of PSI overexcitation requires the dephosphorylation of LHCII. State transitions are accompanied by thylakoid membrane rearrangements, and *VIPP1* was suggested as a candidate protein that might mediate these rearrangements (Chuartzman et al., 2008). To test whether *VIPP1*-RNAi strains are impaired in state transitions, we followed state transitions in control and *VIPP1*-RNAi strains by pulse amplitude-modulated (PAM) fluorometry and 77K fluorescence emission spectroscopy. Both measurements indicated that *VIPP1*-RNAi strains were not impaired in state transitions (see Supplemental Figure 6 online), hence ruling out the possibility that the high light sensitivity of *VIPP1*-RNAi strains originated from a generally reduced capacity for lateral movement of thylakoid membrane complexes. However, we cannot rule out the possibility that membrane complex mobility is impaired in swollen thylakoids.

Thylakoid Swelling Enhances but Does Not Cause the Sensitivity of *VIPP1*-RNAi Strains to High Light

Swelling of thylakoids was observed in *C. reinhardtii* wild-type cells exposed to very high light intensities of 5000 $\mu\text{E m}^{-2} \text{s}^{-1}$ (Topf et al., 1992) and in ATP synthase mutants exposed to 70 $\mu\text{E m}^{-2} \text{s}^{-1}$ (Majeran et al., 2001). Swelling was shown to depend on the acidification of the thylakoid lumen (i.e., on the formation of a $\Delta\mu\text{H}^+$) (Majeran et al., 2001) and was suggested to depend on the presence of ammonium in the growth medium (Topf et al., 1992). The mechanism is that ammonium ions in the stroma become deprotonated to a certain extent and cross the thylakoid membrane as uncharged ammonia. In the acidified lumen, ammonia becomes reprotonated and accumulates as ammonium ions and may thereby generate an increased osmotic pressure within the lumen.

If the high light sensitivity of *VIPP1*-RNAi/amiRNA strains was caused by thylakoid swelling, which only occurs in the presence of ammonium, this phenotype should be abolished when mutant strains are grown on nitrate as nitrogen source. To test this prediction, we took advantage of the fact that the cw15-325 strain used as recipient for the *VIPP1*-RNAi constructs, unlike most *C. reinhardtii* laboratory strains, is able to use nitrate as nitrogen source. We exposed control and *VIPP1*-RNAi strains

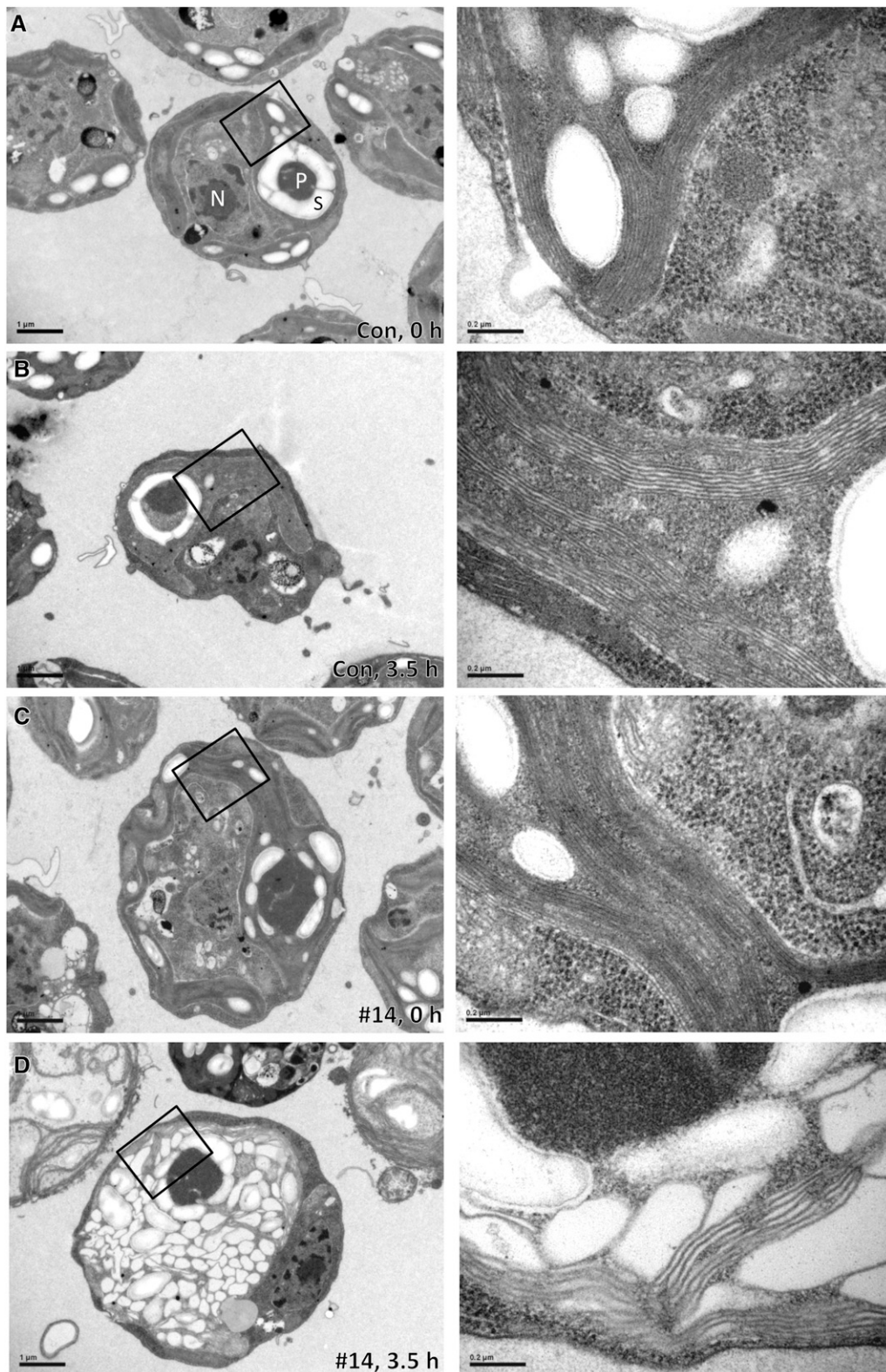


Figure 3. Thylakoids in *VIPP1*-amiRNA Strains Exposed to High Light Intensities Are Extremely Swollen.

(A) Electron microscopy image of a cell from the control strain grown at low light intensities. Cells were grown at $\sim 30 \mu\text{E m}^{-2} \text{ s}^{-1}$ in TAP-NH₄ medium. An overview image is shown on the left, and a zoom-in of the region demarcated by the black box is shown on the right. N, nucleus; P, pyrenoid; S,

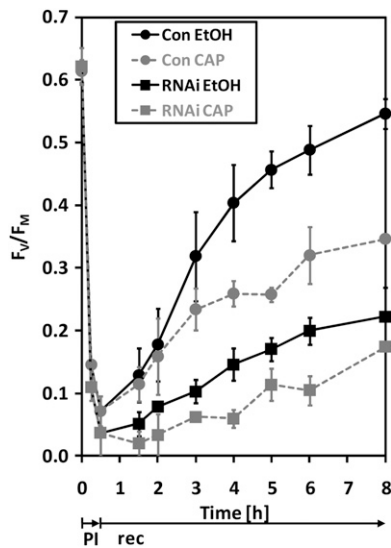


Figure 4. *VIPP1*-RNAi Cells Are More Severely Photoinhibited at Very High Light and Replace Photodamaged D1 by de Novo-Synthesized D1 Slower Than Control Cells.

Cultures of control and *VIPP1*-RNAi strain #80 grown in TAP-NH₄ medium were split into two parts. One part was supplemented with ethanol-dissolved chloramphenicol (CAP) to a final concentration of 100 μg/mL, while the other received the same volume of pure ethanol. The four cultures were exposed to ~1800 μE m⁻² s⁻¹ for 30 min (photo-inhibition [PI]) and shifted back to ~30 μE m⁻² s⁻¹ (recovery [rec]). Maximum quantum efficiency of PSII during photoinhibition and recovery was measured with a PAM fluorometer as variable fluorescence ($F_v = F_M - F_0$) normalized to F_M . Shown is the average of two independent experiments; error bars represent SE.

grown on nitrate to high light intensities (~1000 μE m⁻² s⁻¹) and observed that bleaching was retarded when compared with *VIPP1*-RNAi strains grown on ammonium (see Supplemental Figure 7 online). Also, the loss of PSII maximum quantum efficiency was slower when *VIPP1*-RNAi cells were grown on nitrate compared with ammonium but still faster and more severe than in control cells grown on either nitrogen source (Figure 5A). Similar effects were observed when cells were exposed to very high light intensities (~1800 μE m⁻² s⁻¹) for 60 min and allowed to recover at low light: The extent of PSII photodamage was highest and recovery slowest in *VIPP1*-RNAi cells grown on ammonium, followed by *VIPP1*-RNAi cells grown on nitrate, control cells grown on ammonium, and control cells grown on

nitrate (Figure 5B). Interestingly, PSII maximum quantum efficiency in control cells was also less severely affected by high light when cells were grown on nitrate as a nitrogen source compared with ammonium.

To test whether the reduced high light sensitivity of *VIPP1*-RNAi strains grown on nitrate correlated with reduced thylakoid swelling, we took 1015 electron microscopy images from control and *VIPP1*-RNAi cells, grown on nitrate or ammonium, prior to and after a 3- to 7-h exposure to high light intensities (~1000 μE m⁻² s⁻¹). The observed thylakoid phenotypes were sorted into the categories “ordered,” “disordered,” and “swollen” (examples for these categories are given in Supplemental Figure 8 online). As summarized in Table 1, cells with swollen thylakoids were only observed in *VIPP1*-RNAi strains grown on ammonium. Here, 8% of the cells displayed swollen thylakoids already at low light intensities (30 μE m⁻² s⁻¹) and the fraction of cells with swollen thylakoids increased to 68% after the 3 h of high light exposure. Even after 7 h of high light exposure, we observed no thylakoid swelling in *VIPP1*-RNAi cells grown on nitrate. However, at that time, 36% of the mutant cells were in the process of lysis, whereas this was the case for only 1 to 2% of control cells (see Supplemental Figure 9 online). These results indicate that thylakoid swelling is not causing lysis of high light-exposed *VIPP1*-RNAi cells grown on nitrate.

A similar picture emerged when we took another 800 electron microscopy images to monitor thylakoid swelling in control and *VIPP1*-RNAi cells that were exposed to very high light intensities (~1800 μE m⁻² s⁻¹) for 60 min and allowed to recover at low light. Here, 38% of ammonium-grown *VIPP1*-RNAi cells contained swollen thylakoids after 60 min at very high light, and this fraction declined to 23% after 60 min of recovery at low light intensities (Table 1). Interestingly, swollen thylakoids were also observed in 11% of ammonium-grown control cells exposed to very high light (thus corroborating the observations from Topf et al., 1992) and this fraction declined to 1% after 60 min of recovery. No thylakoid swelling was observed in nitrate-grown control and *VIPP1*-RNAi cells exposed to very high light intensities (Table 1).

Also, at the protein level, we found that exposure to high light (~1000 μE m⁻² s⁻¹) had less severe effects on *VIPP1*-RNAi cells when they were grown on nitrate as the nitrogen source compared with cells grown on ammonium (cf. Figures 1F and 5C). After 7 h of high light exposure, we found a clear decline only in levels of CP43 and perhaps PsaA. While chaperones HSP70B and HSP90C appeared to be normally induced by high light, induction of the LHCSR3 protein still was less pronounced in *VIPP1*-RNAi cells compared with control cells. Likewise, the

Figure 3. (continued).

starch. Bars in overview images correspond to 1 μm and those in zoom-ins to 0.2 μm.

(B) Electron microscopy image of a cell from the control strain exposed to high light. Cells were grown at ~30 μE m⁻² s⁻¹ in TAP-NH₄ medium and exposed to ~1000 μE m⁻² s⁻¹ for 3.5 h. Images were taken as in **(A)**.

(C) Electron microscopy image of a cell from a *VIPP1*-amiRNA strain grown at low light intensities. *VIPP1*-amiRNA strain #14 was grown at ~30 μE m⁻² s⁻¹ in TAP-NH₄ medium. Images were taken as in **(A)**.

(D) Electron microscopy image of a cell from a *VIPP1*-amiRNA strain exposed to high light. *VIPP1*-amiRNA strain #14 was grown at ~30 μE m⁻² s⁻¹ in TAP-NH₄ medium and exposed to ~1000 μE m⁻² s⁻¹ for 3.5 h. Images were taken as in **(A)**.

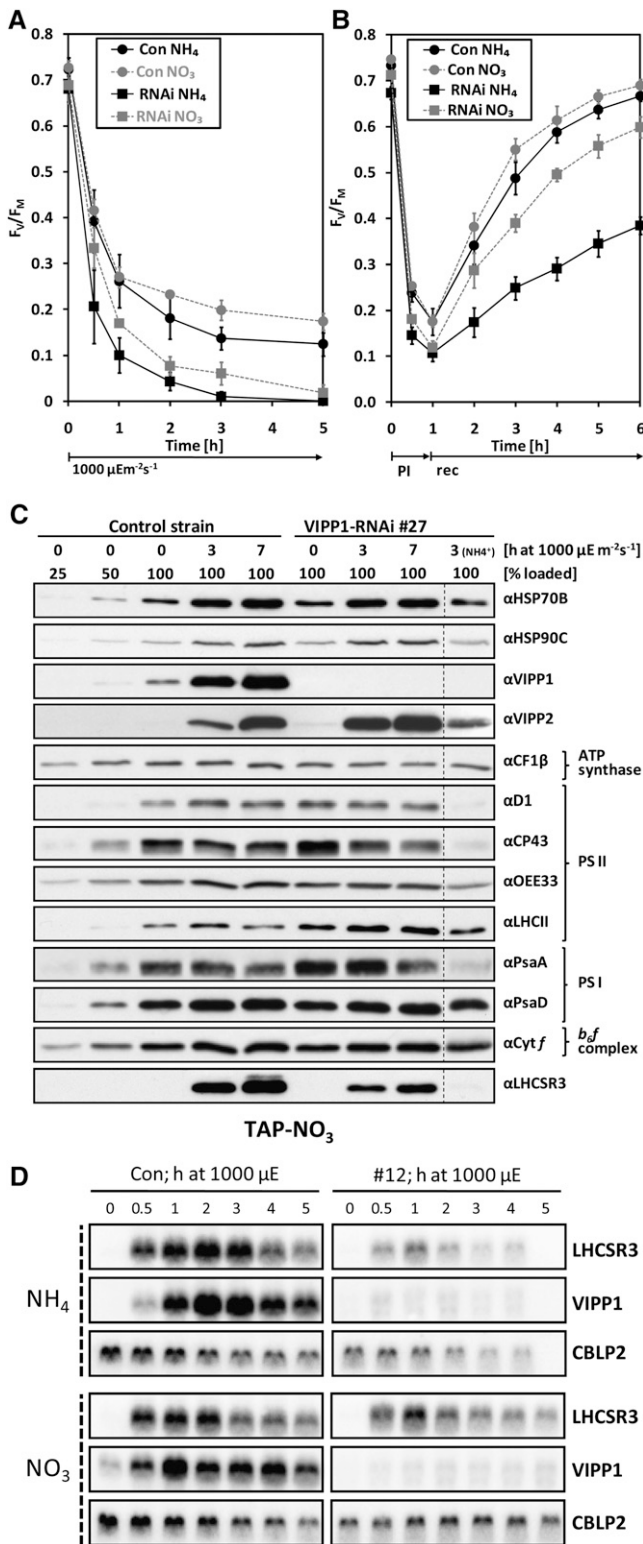


Figure 5. High Light Sensitivity Is Alleviated but Not Abolished in *VIPP1*-RNAi Strains Grown on Nitrate.

(A) PSII in *VIPP1*-RNAi strains is less high light sensitive in cells grown on nitrate compared with ammonium. Control (Con) and *VIPP1*-RNAi strains

strong decline of PSII subunits and, to a lesser extent, of PSI subunits observed after exposure of ammonium-grown *VIPP1*-RNAi cells to very high light intensities ($\sim 1800 \mu\text{E m}^{-2} \text{s}^{-1}$) was virtually absent upon exposure of nitrate-grown *VIPP1*-RNAi cells to very high light (cf. Supplemental Figures 10A and 10B online). However, similar to *VIPP1*-RNAi cells continuously exposed to $\sim 1000 \mu\text{E m}^{-2} \text{s}^{-1}$, *VIPP1*-RNAi cells exposed for 60 min to $\sim 1800 \mu\text{E m}^{-2} \text{s}^{-1}$ were still deficient in the full induction of LHCSR3 protein expression, despite growth on nitrate. And in contrast with nitrate-grown *VIPP1*-RNAi cells exposed to $\sim 1000 \mu\text{E m}^{-2} \text{s}^{-1}$, cells exposed to $\sim 1800 \mu\text{E m}^{-2} \text{s}^{-1}$ still appeared to be impaired in the full induction of the HSP70B and HSP90C chaperones and still experienced reduction of initially elevated LHCII levels (see Supplemental Figure 10B online).

Taken together, our observations show that thylakoid swelling is indeed only observed in high light-treated cells grown on ammonium as nitrogen source. Moreover, thylakoid swelling enhances, but does not cause, the sensitivity of *VIPP1*-RNAi strains to high light.

VIPP1-RNAi Strains Are Impaired in the Perception/Transmission of the High Light State

We wondered whether the reduced accumulation of the LHCSR3 protein in high light-treated *VIPP1*-RNAi strains was due to a defect in the perception/transmission of the high light conditions or due to a defect in the conversion of the *LHCSR3* transcript into a stable protein, for example, by its impaired insertion into the thylakoid membrane. In the former case, we expected a reduced induction of the *LHCSR3* transcript in response to high light, and in the latter case, we expected no effect. To distinguish between these possibilities, we exposed cells from control and *VIPP1*-RNAi strains to high light ($\sim 1000 \mu\text{E m}^{-2} \text{s}^{-1}$) and analyzed *LHCSR3* transcripts by RNA gel blot analysis. As shown in Figure

#12 and #27 were grown in TAP-NO₃ or TAP-NH₄ medium. Cells were exposed to $\sim 1000 \mu\text{E m}^{-2} \text{s}^{-1}$ and maximum quantum efficiency of PSII was measured with a PAM fluorometer as described in Figure 4. Shown is the average of two independent experiments. Error bars represent SE. **(B)** PSII in *VIPP1*-RNAi strains is less sensitive to photoinhibition in cells grown on nitrate compared with ammonium. Control and *VIPP1*-RNAi strains #5, #27, and #41 were grown in TAP-NO₃ or TAP-NH₄ medium. Cells were exposed to $\sim 1800 \mu\text{E m}^{-2} \text{s}^{-1}$ for 1 h (photoinhibition [PI]) and shifted back to $\sim 30 \mu\text{E m}^{-2} \text{s}^{-1}$ (recovery [rec]). Maximum quantum efficiency of PSII was measured with a PAM fluorometer as described in Figure 4. Shown is the average of four independent experiments. Error bars represent SE.

(C) Subunits of PSII and PSI are less prone to degradation in high light-exposed *VIPP1*-RNAi strains grown on nitrate. Whole-cell proteins from nitrate-grown control and *VIPP1*-RNAi strain #27 exposed to $\sim 1000 \mu\text{E m}^{-2} \text{s}^{-1}$ for 7 h were separated on 14% SDS-polyacrylamide gels and analyzed by immunoblotting. For comparison, whole-cell proteins from *VIPP1*-RNAi strain #27 grown on ammonium and exposed to high light for 3 h was loaded next to the other samples.

(D) RNA gel blot analysis of high light-exposed control and *VIPP1*-RNAi strains. Control and *VIPP1*-RNAi strain #12 were grown in TAP-NO₃ or TAP-NH₄ medium. Cells were exposed to $\sim 1000 \mu\text{E m}^{-2} \text{s}^{-1}$ for 5 h, and RNA was extracted from samples taken at the indicated time points and subjected to RNA gel blot analysis. *CBLP2* served as loading control.

Table 1. Categorization of Thylakoid Structure

Condition/Strain	TAP-NH ₄			TAP-NO ₃		
	Ordered	Disordered	Swollen	Ordered	Disordered	Swollen
High light						
Con (LL)	100	0	0	99	1	0
Con (180' HL)	100	0	0	98	2	0
RNAi #27 (LL)	92	0	8	92	8	0
RNAi #27 (180' HL)	27	5	68	100	0	0
Photoinhibition						
Con (60' PI)	87	0	11	98	2	0
Con (60' rec)	99	0	1	100	0	0
RNAi #27 (60' PI)	46	16	38	92	8	0
RNAi #27 (60' rec)	74	3	23	100	0	0
Heat shock						
Con (25°C)	96	0	4	n.d.	n.d.	n.d.
Con (40°C)	100	0	0	n.d.	n.d.	n.d.
RNAi #32 (25°C)	90	1	9	n.d.	n.d.	n.d.
RNAi #32 (40°C)	70	9	21	n.d.	n.d.	n.d.

Transmission electron microscopy images were taken on cells from control (Con) and *VIPP1*-RNAi strains #27 and #32 grown in TAP-NH₄ or TAP-NO₃ medium under the following conditions: LL, low light intensities of $\sim 30 \mu\text{E m}^{-2} \text{ s}^{-1}$; 180' HL, 180 min at high light intensities of $\sim 1000 \mu\text{E m}^{-2} \text{ s}^{-1}$; 60' PI, 60 min at photoinhibitory light of $\sim 1800 \mu\text{E m}^{-2} \text{ s}^{-1}$; 60' rec, 60 min at $\sim 30 \mu\text{E m}^{-2} \text{ s}^{-1}$ for recovery from photoinhibition; 25°C, 25°C and $\sim 30 \mu\text{E m}^{-2} \text{ s}^{-1}$; 40°C, 40°C for 1 h at $\sim 5 \mu\text{E m}^{-2} \text{ s}^{-1}$; n.d., not determined. On average, 101 images per strain and condition were analyzed and sorted into three categories of thylakoid structure for which examples are shown in Supplemental Figure 8 online. Values are in percentages.

5D, induction of the *LHCSR3* gene was strongly reduced in *VIPP1*-RNAi strains grown on ammonium as nitrogen source, while differences between control and *VIPP1*-RNAi strains in the high light-induced expression of *LHCSR3* were less pronounced in cells grown on nitrate. Similarly, the reduction in *LHCSR3* transcript accumulation observed in *VIPP1*-RNAi strains compared with a control strain after exposure to very high light ($\sim 1800 \mu\text{E m}^{-2} \text{ s}^{-1}$) was less pronounced in cells grown on nitrate instead of ammonium (see Supplemental Figure 10C online). In control cells grown on either nitrogen source, *VIPP1* transcripts strongly accumulated in response to high light exposure, whereas they were almost completely repressed in *VIPP1*-RNAi cells.

These data suggest that the reduced accumulation of *LHCSR3* protein observed in high light-exposed *VIPP1*-RNAi strains compared with the control strain is mainly based on the reduced expression of the *LHCSR3* gene. This in turn suggests that *VIPP1*-RNAi strains are defective in the retrograde signaling of the high light state, which is particularly true for cells grown on ammonium but to a lesser extent also for cells grown on nitrate.

Specific Photosynthesis Parameters Are Affected in *VIPP1*-RNAi Strains

Although we have shown that thylakoid swelling occurs only in high light-exposed cells grown on ammonium and not in cells grown on nitrate as nitrogen source, it is not clear why swelling is so much more severe in *VIPP1*-RNAi/amiRNA cells compared with control cells. Thylakoid swelling in high light-treated mutant cells may result from their ability to generate a higher proton-motive force (pmf) than control cells, caused, for example, by a reduced ATP synthase activity (Majeran et al., 2001). Alternatively, thylakoid swelling may have resulted from the reduced

ability of *VIPP1*-RNAi/amiRNA strains to counteract the osmotic pressure generated in the thylakoids upon high light exposure in the presence of ammonium.

To test the first hypothesis, we used electrochromic shift measurements to compare control and *VIPP1*-RNAi strains in respect of the pmf generated at saturating light intensities and with regard to thylakoid membrane conductivity (corresponding to the ATP synthase activity). These measurements were performed with cells grown under mixotrophic conditions using ammonium and nitrate as nitrogen sources prior to and after a 3-h exposure to high light intensities of $\sim 1000 \mu\text{E m}^{-2} \text{ s}^{-1}$. As shown in Figures 6A and 6B, maximum thylakoid membrane energization in saturating light was significantly lower in *VIPP1*-RNAi cells than in control cells under all conditions tested, while ATP synthase activity was never lower than that of the untreated control. Hence, thylakoid swelling in high light-exposed *VIPP1*-RNAi strains is unlikely to be caused by an increased pmf.

To identify defects in *VIPP1*-RNAi strains that might cause their increased light sensitivity and the apparently reduced ability of their thylakoids to withstand an increased osmotic pressure, we performed more biophysical measurements. We observed that PSII maximum quantum efficiency was mildly but significantly lower in *VIPP1*-RNAi compared with control cells (Figure 6C), while cytochrome *f* reduction and oxidation kinetics were unaltered, indicating similar electron transport capacities in control and *VIPP1*-RNAi strains (Figures 6E and 6F). Moreover, the chlorophyll *a/b* ratio was lower in *VIPP1*-RNAi strains than in controls (Figure 6D), which may result from the higher levels of chlorophyll *b*-rich LHCII in *VIPP1*-RNAi strains (Figures 1F and 5C; see Supplemental Figures 10A and 10B online). A larger antenna cross section in *VIPP1*-RNAi strains was also supported by the faster fluorescence induction kinetics in mutant versus

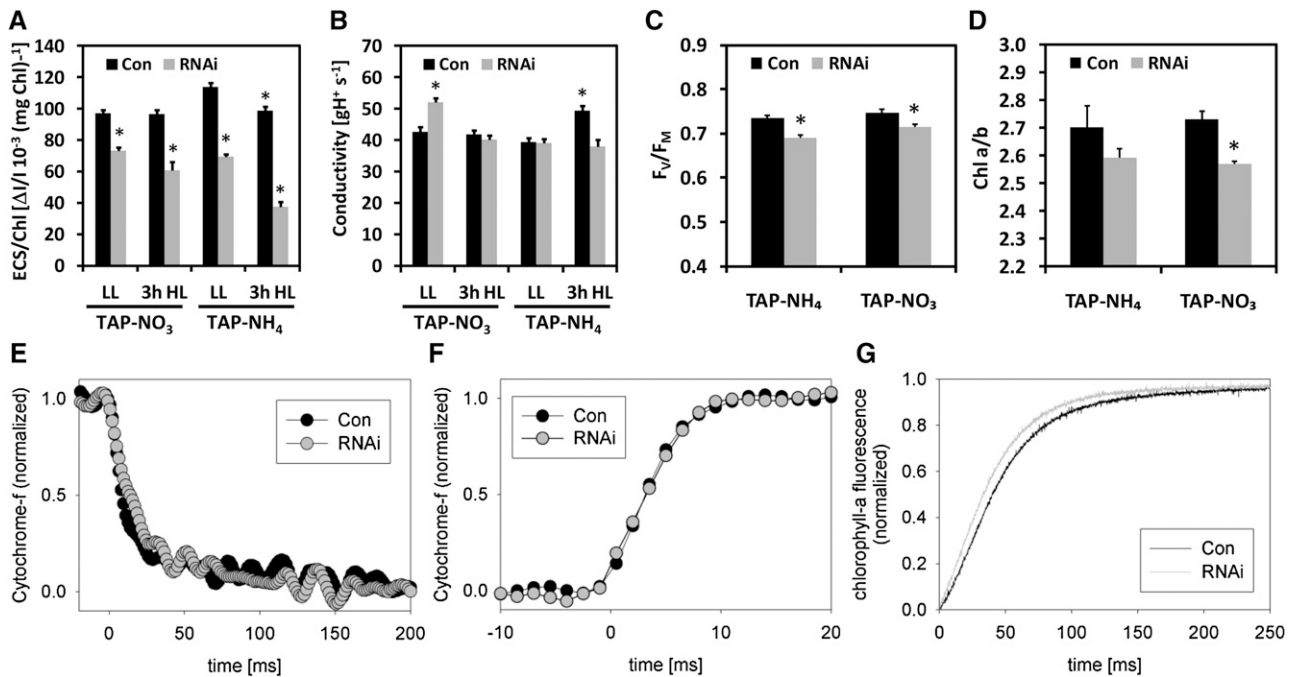


Figure 6. *VIPP1*-RNAi Strains Are Affected in Some Photosynthesis Parameters.

(A) Thylakoid membrane energization. Control (Con) and *VIPP1*-RNAi strains #2-7, #2-19, #3-30, #4-19, and #4-49 were grown in TAP-NO₃ or TAP-NH₄ medium and kept in low light (LL) of $\sim 30 \mu\text{E m}^{-2} \text{s}^{-1}$ or exposed to high light (HL) of $\sim 1000 \mu\text{E m}^{-2} \text{s}^{-1}$ for 3 h. Maximum thylakoid membrane energization was then determined by measuring the ECS in saturating light. Shown is the average of eight and six measurements on the control strain, and of 12 and 10 measurements on *VIPP1*-RNAi strains grown on NO₃ and NH₄, respectively. Error bars represent SE. Chl, chlorophyll.

(B) ATP synthase activity. The activity of the ATP synthase was inferred from the decay kinetic of the ECS signal during a short dark interval using the same cells as described in **(A)**.

(C) Maximum quantum efficiency of PSII. F_v/F_m was measured after 5 min of far-red illumination, followed by 10 min of dark adaptation of the cell suspension using control and *VIPP1*-RNAi strains #4, #9, #40, #53, #72, #111, and #129 grown in low light. Shown is the average of four and two measurements on the control strain, and of 13 and 10 measurements on *VIPP1*-RNAi strains. Error bars represent SE.

(D) Chlorophyll *a/b* ratios. Chlorophyll was extracted with 80% acetone from the same cells as described in **(C)**, and the chlorophyll concentration was determined spectrophotometrically. Asterisks indicate the significance of the difference to untreated controls grown on nitrate and ammonium as nitrogen source, respectively (*t* test, *P* value ≤ 0.05).

(E) Cytochrome *f* reduction kinetics. Control and *VIPP1*-RNAi strains #4, #9, and #72 were grown in TAP-NH₄ medium. Cytochrome *f* reduction was initiated by switching off saturating red light at time point zero. The fully reduced state of cytochrome *f* in the dark was normalized to zero and the fully oxidized state to one. Kinetics were recorded four times for each strain. Averages for the control and all three *VIPP1*-RNAi strains are shown.

(F) Cytochrome *f* oxidation kinetics. Control and *VIPP1*-RNAi strains #27, #29, and #30 were grown in TAP-NH₄ medium. Cytochrome *f* oxidation was initiated by switching on a saturating red light pulse at time point zero. Six replicates of the control strain and two each for the *VIPP1*-RNAi strains were recorded and averages plotted. Again, the fully reduced state of cytochrome *f* in the dark was normalized to zero and the fully oxidized state to one.

(G) Fluorescence induction kinetics. Control and *VIPP1*-RNAi strains #27, #29, and #30 were grown in TAP-NH₄ medium. Dark-adapted cells trapped in state 1 were supplemented with DCMU, and fluorescence was induced by switching on the light. Six replicates of the control strain and two each for the *VIPP1*-RNAi strains were recorded and averages plotted. The F_0 values were normalized to zero and the F_m values to one.

control cells (Figure 6G). The reduced sigmoidicity of the induction curve in *VIPP1*-RNAi strains also indicated that cooperativity between PSII centers was lower in *VIPP1*-RNAi cells compared with control cells. Thermoluminescence measurements revealed a reduced maximum temperature of the Q-band in *VIPP1*-RNAi cells compared with control cells, a phenotype observed in cells grown on either ammonium or nitrate as nitrogen source (Table 2). Following a saturating single turnover flash at low temperature, a thermoluminescence band, named Q-band, is detected upon heating in the region of 8 to 12°C. This luminescence arises from the recombination of the charge pairs $S_2Q_A^-$, with S_2 being an oxidation state of the Mn cluster and Q_A^- the semireduced

primary quinone acceptor in PSII (Rutherford et al., 1982). The downshift of the maximum temperature of the Q-band in *VIPP1*-RNAi cells reflects a lowering of the midpoint potential of the redox couple Q_A/Q_A^- (Krieger-Liszkay and Rutherford, 1998). A lower midpoint potential of Q_A was shown to increase the yield of $^1\text{O}_2$ generation in the light (Fufezan et al., 2007) and therefore might represent the mechanism underlying high light sensitivity of *VIPP1*-RNAi cells.

We conclude that thylakoid swelling in ammonium-grown, high light-exposed *VIPP1*-RNAi cells is not caused by a higher pmf but rather by a disturbance of thylakoid membrane organization. The defect causing this disturbance also might be

Table 2. Temperature of the Q-Band

Strain	T_{\max} (°C)	
	TAP-NO ₃ ⁻	TAP-NH ₄ ⁺
Wild type	11.5 ± 0.5	11.1 ± 0.5
VIPP1-RNAi	7.4 ± 1.2	9.0 ± 1.0

Cells were grown in TAP medium containing either ammonium or nitrate as nitrogen source at low light intensities of $\sim 30 \mu\text{E m}^{-2} \text{ s}^{-1}$. Values represent averages \pm SD from thermoluminescence measurements on two independent control and four independent VIPP1-RNAi lines (#1-29, #2-29, #4-49, and #3-30).

responsible for reduced PSII maximum quantum efficiency, lowered midpoint potential of the Q_A/Q_A^- redox couple, reduced PSII cooperativity, and increased antenna cross section in VIPP1-RNAi strains.

PSII in VIPP1-RNAi Strains Is Sensitive to Heat Shock

If VIPP1-RNAi strains indeed have a defect in thylakoid membrane organization, we can expect phenotypes to also be induced by treatments that do not involve high light intensities but similarly represent a threat to thylakoid membrane integrity. Heat shock at 40 to 42°C, combined with low light intensities that are harmless for PSII at 25°C, was demonstrated previously to result in loss of PSII activity in *C. reinhardtii* (Schuster et al., 1988). This finding was explained by cross-linking of the D1 protein with other proteins, presumably induced by the overproduction of radicals by heat-stressed, illuminated PSII.

To test whether PSII was more sensitive to heat stress in VIPP1-RNAi cells compared with control cells, we shifted mutant and control cells from 25 to 40°C at low light intensities ($\sim 5 \mu\text{E m}^{-2} \text{ s}^{-1}$) for 4 h and back to 25°C for 2 h for recovery and monitored PSII maximum quantum efficiency. This experiment was done with cells grown mixotrophically on ammonium or nitrate as nitrogen sources (Figure 7A). In control cells grown on ammonium, variable fluorescence/maximum fluorescence (F_V/F_M) declined already within the first 20 min at 40°C from 0.73 to 0.68 but recovered fast and almost completely after shifting cells back to 25°C. The loss of PSII maximum quantum efficiency in nitrate-grown control cells followed kinetics similar to those observed for ammonium-grown control cells within the first hour at 40°C, but, surprisingly, the F_V/F_M value in nitrate-grown control cells declined further to 0.58 after 4 h at 40°C and did not fully recover at 25°C. In ammonium-grown VIPP1-RNAi cells heat-shocked for 4 h, the F_V/F_M value dropped dramatically from 0.68 to 0.44 and at 25°C recovered only to 0.47. This effect was less drastic in nitrate-grown VIPP1-RNAi cells, in which the F_V/F_M value declined from 0.69 to 0.49 after 4 h at 40°C and recovered maximally to 0.57 at 25°C.

To pick up changes induced by heat shock at the ultrastructural level, we took 400 electron microscopy images of ammonium-grown control and VIPP1-RNAi strains prior to and after a 60-min heat shock at 40°C. Interestingly, the fraction of VIPP1-RNAi cells containing swollen thylakoids increased from 9% under nonstress conditions to 21% after heat shock (Table 1).

Heat stress hardly affected levels of thylakoid membrane complex subunits, no matter on which nitrogen source cells were grown (Figure 7B; see Supplemental Figure 11A online). While in control and VIPP1-RNAi strains levels of the VIPP2 protein were strongly induced by high light, they did not further increase during heat stress (cf. Figures 1F and 7B). Also, in contrast with the high light treatments, the HSP70B and HSP90C chaperones appeared to be normally induced by heat stress in VIPP1-RNAi strains grown on either nitrogen source (Figure 7B; see Supplemental Figure 11A online). In control cells grown on ammonium or nitrate, heat stress led to a slight and transient increase in levels of the LHCSR3 protein. By contrast, we observed no increase in LHCSR3 levels in VIPP1-RNAi strains after 60 min at 40°C but a strong decline of LHCSR3 after 180 min of heat stress. Also, the induction of the LHCSR3 gene by heat shock was reduced in VIPP1-RNAi strains (Figure 7C; see Supplemental Figure 11B online).

In summary, the dramatic effect of heat stress combined with very low light intensities on PSII maximum quantum efficiency in VIPP1-RNAi strains further supports the notion that VIPP1 depletion affects the structural organization of thylakoid membrane protein complexes. Again, this defect appears to have a secondary effect on retrograde signaling triggering the induction of the LHCSR3 gene by heat stress.

Quantitative Proteomics Reveals Reduced Levels of Thylakoid Membrane Core Complexes but Increased LHCII Levels in VIPP1-RNAi Cells

The conclusion that thylakoid membranes in VIPP1-RNAi/amiRNA strains must harbor structural defects prompted us to get a more quantitative picture of alterations in the abundance of thylakoid membrane protein complexes. The method of choice for this is shotgun proteomics using ¹⁵N metabolic labeling, by which a much higher quantification accuracy can be achieved than with quantitative immunoblotting (Mühlhaus et al., 2011). Using this technique, we compared the accumulation of the main protein complexes of the thylakoid membranes in control and VIPP1-RNAi strains that were grown on ¹⁵NO₃ and ¹⁴NO₃ as nitrogen sources, respectively (cells were grown on nitrate to avoid the pleiotropic effects caused by thylakoid swelling in cells grown on ammonium). Cells were kept under nonstress conditions or were exposed to 1-h high light ($2000 \mu\text{E m}^{-2} \text{ s}^{-1}$) and heat stress (40°C) treatments. For comparison, we also quantified the relative abundances of protein complexes involved in the mitochondrial respiratory chain. For each of the measured complexes, average values determined for multiple subunits are shown (see Supplemental Data Set 1 online for details). As shown in Figure 8A, thylakoid membranes from unstressed VIPP1-RNAi cells contained 30% more LHCII, but $\sim 20\%$ less PSII, PSI, and cytochrome *b₆f* complex, and 14% less ATP synthase than control cells, while levels of LHCI were unchanged. High light and heat stress led to a 5 to 15% decline in levels of LHCs and to less reduced levels of PSI (5 to 9%) and ATP synthase (8%) in VIPP1-RNAi cells compared with stressed control cells, while levels of PSII and cytochrome *b₆f* complex remained unchanged. Interestingly, levels of the mitochondrial NADH:ubiquinone oxidoreductase complex were 30% higher and those of the cytochrome oxidase 19% lower in VIPP1-RNAi cells compared

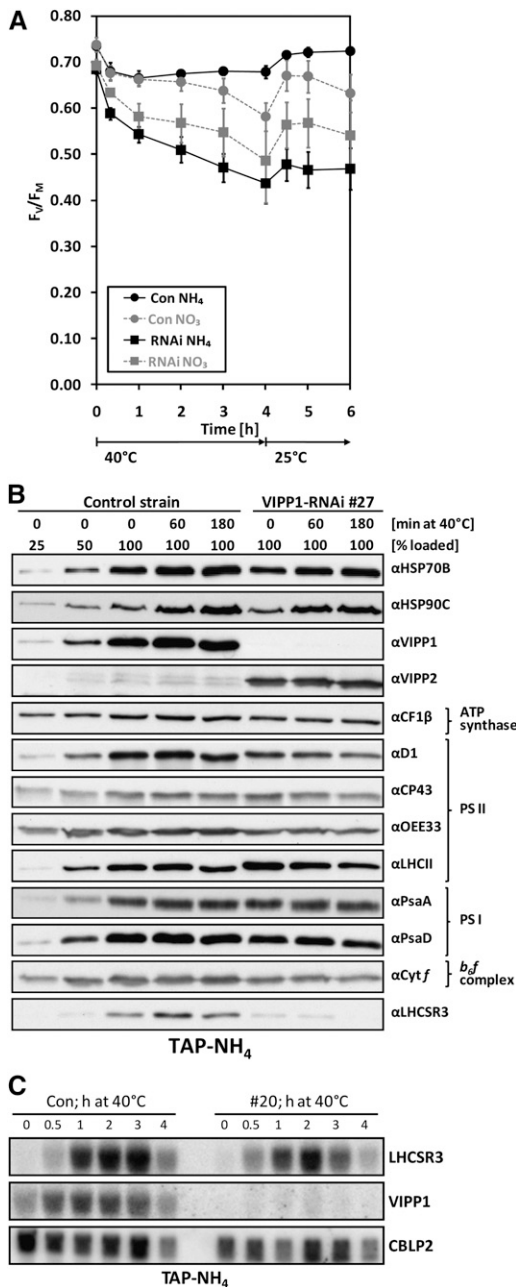


Figure 7. PSII of *VIPP1*-RNAi Strains Is Highly Susceptible to Heat Stress.

(A) PSII maximum quantum efficiency of control and *VIPP1*-RNAi strains exposed to heat stress. Control and *VIPP1*-RNAi strains #5, #20, #27, and #41 were grown in TAP- NO_3 or TAP- NH_4 medium, and cells were exposed to 40°C at $\sim 5 \mu\text{E m}^{-2} \text{s}^{-1}$. F_v/F_m was measured over time with a PAM fluorometer as described in Figure 4. Shown is the average of five and seven independent experiments for control and *VIPP1*-RNAi strains, respectively. Error bars represent SE.

(B) PSII subunits are only mildly affected by heat stress. Whole-cell proteins were extracted from control and *VIPP1*-RNAi strain #27 grown and heat-stressed as described in **(A)**. Whole-cell proteins were separated on 14% SDS-polyacrylamide gels and analyzed by immunoblotting.

(C) RNA gel blot analysis of heat-stressed control and *VIPP1*-RNAi

with control cells, while levels of the ubiquinone-cytochrome c reductase complex and the ATP synthase were unchanged (Figure 8B). High light and heat stress treatments led to increases in levels of all respiratory chain complexes in *VIPP1*-RNAi cells relative to control cells. Hence, it appears possible that *VIPP1*-RNAi cells attempt to compensate for the slightly reduced capacity of their thylakoid membrane core complexes by higher levels of the NADH:ubiquinone oxidoreductase complex.

VIPP1-RNAi/amiRNA Strains Harbor Aberrant Structures at the Origin of Thylakoid Membranes

Finally, we set out to gather information on the localization of *VIPP1* in *C. reinhardtii* by immunofluorescence microscopy using an affinity-purified antibody against *VIPP1*. As shown in Figure 9, *VIPP1* was detected as diffuse material throughout the chloroplast but also in distinct dot-like structures that sometimes also appeared to extend into rods. This localization pattern is in agreement with our previous finding that *C. reinhardtii* *VIPP1* may form rod-like structures, which by the help of molecular chaperones are kept in equilibrium with ring-like structures and smaller assembly states (Liu et al., 2007). We reasoned that rod-like structures potentially formed by *VIPP1* might be visible in electron micrographs and therefore revisited the electron micrographs taken from control and *VIPP1*-RNAi/amiRNA strains. Unfortunately, we were unable to distinguish potential *VIPP1* rods or rings from thylakoid membranes in control cells. However, we observed aberrant structures in *VIPP1*-RNAi/amiRNA cells that occurred particularly in regions from which multiple thylakoid membrane layers emerge and which may well correspond to the dot-like structures observed by immunofluorescence microscopy (Figure 10A). The aberrant structures were often found around the pyrenoid but were not restricted to that area. They were very rarely observed in control cells and occurred abundantly and consistently in *VIPP1*-RNAi and amiRNA cells (cw15-325 and cw15-302 backgrounds, respectively) grown under low light or high light on ammonium or nitrate (Table 3, Figure 10).

In summary, we could localize *VIPP1* to diffuse material and to distinct spots within the chloroplast. As judged from the distribution of these spots, they may correspond to regions from which multiple thylakoid layers emerge and which are aberrant in *VIPP1*-RNAi/amiRNA cells.

DISCUSSION

Here, we present a thorough characterization of phenotypes arising in *C. reinhardtii* as a consequence of *VIPP1* depletion mediated by inverted repeat and amiRNA constructs. *VIPP1* expression was repressed in two different strain backgrounds: In strain cw15-325, *VIPP1* levels were reduced to below 5% of wild-type levels, which, however, resulted in the upregulation of *VIPP2*

strains. Control and *VIPP1*-RNAi strain #20 were grown and heat stressed as described in **(A)**. RNA was extracted from samples taken at the indicated time points and subjected to RNA gel blot analysis. *CBLP2* served as loading control.

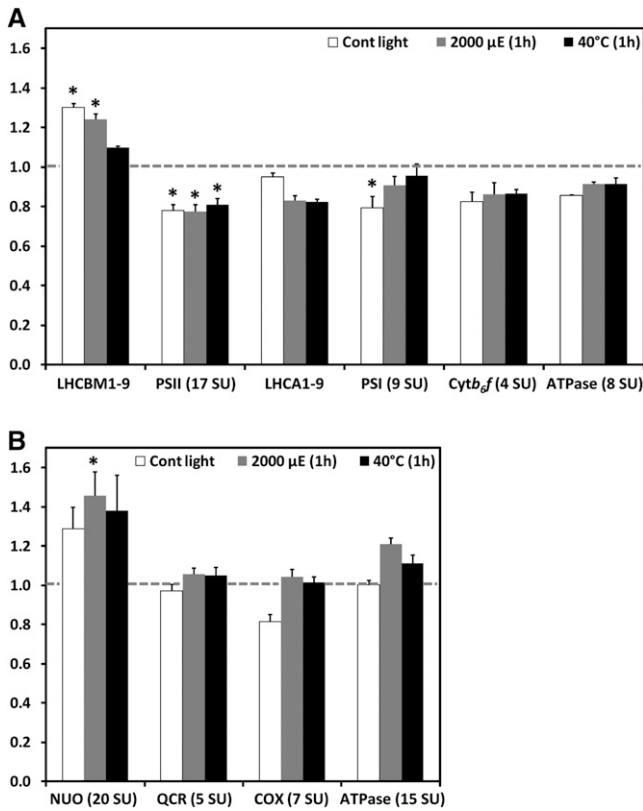


Figure 8. Levels of Thylakoid Membrane Core Complexes Are Slightly Lower in *VIPP1*-RNAi Cells Than in Control Cells.

(A) Ratios of thylakoid membrane core complexes in *VIPP1*-RNAi cells relative to control cells. A control strain and *VIPP1*-RNAi strain #111 were metabolically labeled using $^{15}\text{NO}_3$ and $^{14}\text{NO}_3$, respectively, as nitrogen source and maintained at $\sim 30 \mu\text{E m}^{-2} \text{s}^{-1}$ and 25°C for 1 h (Cont light), exposed to photoinhibitory light of $\sim 2000 \mu\text{E m}^{-2} \text{s}^{-1}$ for 1 h at 25°C , or heat-shocked at 40°C for 1 h at $\sim 5 \mu\text{E m}^{-2} \text{s}^{-1}$. After mixing control and *VIPP1*-RNAi cells from each treatment at a 1:1 ratio, proteins in membrane-enriched fractions were separated by SDS-PAGE and digested tryptically in gel. Peptides were eluted, desalted, and analyzed by liquid chromatography-MS/MS. Peptide identification and quantification was performed as described previously (Mühlhaus et al., 2011). Quantification values for single core complex subunits (SU) were computed from quantified peptides, and the average ratio of light (*VIPP1*-RNAi) to heavy (control) subunits was calculated for the different core complexes. Error bars represent SE, and asterisks indicate the significance of the difference of the ratio from one (assuming equal variance; *t* test, *P* value ≤ 0.05).

(B) Ratios of respiratory chain core complexes in *VIPP1*-RNAi relative to control cells. Ratios of respiratory chain core complexes were determined as described in **(A)**.

to $\sim 20\%$ of total VIPP levels (Figure 2). In strain cw15-302, VIPP2 was not upregulated in response to VIPP1 repression, but VIPP1 only could be reduced to $\sim 25\%$ of wild-type levels. Despite the different contributions of VIPP1 and VIPP2 to the residual VIPP pool in the two strain backgrounds, phenotypes were the same (Figures 1, 3, and 10; see Supplemental Figure 5B online). This allows us to conclude that VIPP1 and VIPP2 largely are functionally redundant and that the total VIPP pool in *C. reinhardtii*

cannot be constitutively repressed to below $\sim 25\%$ of wild-type levels, similar to the situations in *Arabidopsis* and cyanobacteria (Kroll et al., 2001; Westphal et al., 2001; Fuhrmann et al., 2009a). VIPP1 obviously is the major VIPP species in *C. reinhardtii*, a conclusion that is supported by the finding that repression of VIPP2 in strain cw15-325 did not result in high light sensitivity or other obvious phenotypes (M. Rütgers and M. Schroda, unpublished results). Very similar phenotypes obtained with *VIPP1*-RNAi and -amiRNA constructs also widely rule out off-target effects in the RNAi lines (Ossowski et al., 2008).

Aberrant Structures at the Origin of Multiple Thylakoid Membrane Layers in *VIPP1*-RNAi/amiRNA Cells Suggest Deficits in Thylakoid Biogenesis

The most striking observation we consistently made in *C. reinhardtii* *VIPP1*-RNAi/amiRNA cells under all growth conditions tested is the occurrence of aberrant structures in regions from which bundles of thylakoids emerge and often project into different directions of the chloroplast (Figure 10, Table 3). Some of these aberrant structures were reminiscent of prolamellar body (PLB)-like structures observed in dark-grown *C. reinhardtii* *y-1* (yellow-in-the-dark) mutants (Hooper and Blobel, 1969; Friedberg et al., 1971). *y-1* mutants are defective in nuclear genes that are involved in the posttranscriptional expression or accumulation of the CHLL subunit of the light-independent protochlorophyllide reductase, leading to a block in chlorophyll synthesis in the dark (Cahoon and Timko, 2000). Hence, PLB-like structures are likely made of thylakoid membrane precursors in

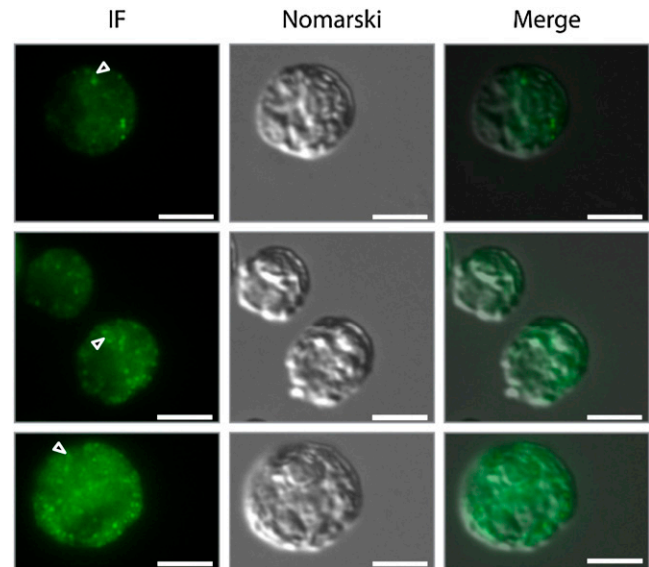


Figure 9. Immunofluorescence Microscopy Detects VIPP1 in Distinct Punctae and as Diffuse Material in the Chloroplast.

Control cells from the cw15-325 background were grown at $\sim 30 \mu\text{E m}^{-2} \text{s}^{-1}$ in TAP- NH_4 medium and fixed and processed for immunofluorescence (IF) microscopy as described in Methods. The signal recognized by the affinity-purified anti-VIPP1 antibody is shown in green. Triangles indicate potential rod-like extensions. Bars = $5 \mu\text{m}$.

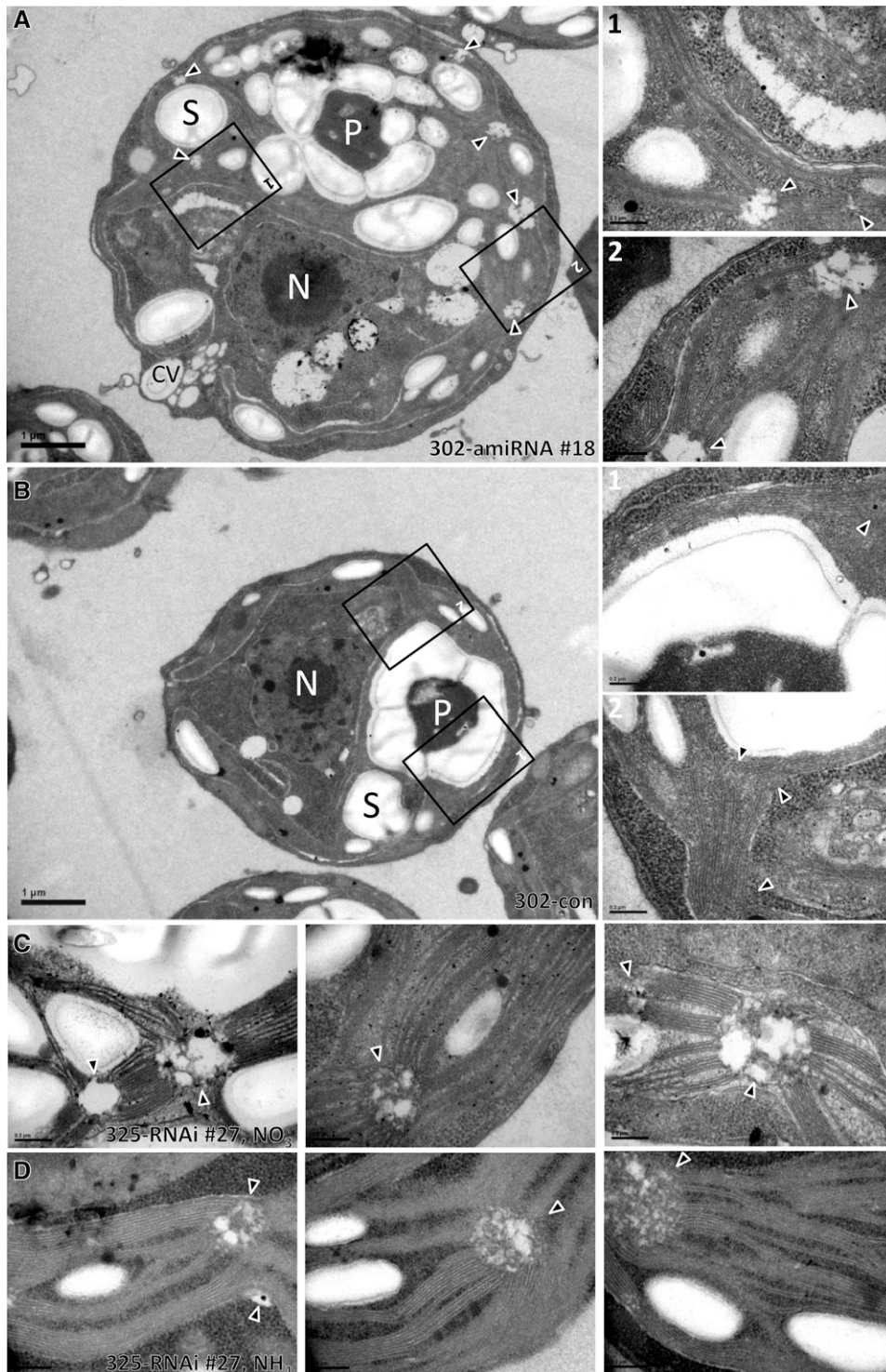


Figure 10. *VIPP1*-RNAi/amiRNA Strains Harbor Aberrant Structures at the Origin of Thylakoid Membranes.

(A) Electron microscopy image of a cell from *VIPP1*-amiRNA strain #18. Cells were grown at $\sim 30 \mu\text{E m}^{-2} \text{s}^{-1}$ in TAP- NH_4 medium. An overview image is shown on the left, and zoom-ins of the regions demarcated by black boxes are shown on the right. Triangles indicate regions at the origin of multiple thylakoid membrane ramifications. CV, contractile vacuole; N, nucleus; P, pyrenoid; S, starch. Bars in overview images = 1 μm , those in zoom-ins = 0.2 μm .
(B) Electron microscopy image of a cell from the control strain. Cells were grown at $\sim 30 \mu\text{E m}^{-2} \text{s}^{-1}$ in TAP- NH_4 medium. Symbols are as in **(A)**.
(C) Electron microscopy image of a cell from *VIPP1*-RNAi strain #27. Cells were grown at $\sim 30 \mu\text{E m}^{-2} \text{s}^{-1}$ in TAP- NO_3 medium. Abbreviations are as in **(A)**.
(D) Electron microscopy image of a cell from *VIPP1*-RNAi strain #27. Cells were grown at $\sim 30 \mu\text{E m}^{-2} \text{s}^{-1}$ in TAP- NH_4 medium. Abbreviations are as in **(A)**.

Table 3. Quantification of Aberrant PLB-Like Structures in *VIPP1*-RNAi/amiRNA Cells

Strain	Growth Conditions		
cw15-325	LL, NH ₄	3.5 h HL, NH ₄	LL, NO ₃
	Control	0	n.d.
	<i>VIPP1</i> -RNAi #27	44	n.d.
	<i>VIPP1</i> -RNAi #41	63	n.d.
cw15-302	Control	2	4
	<i>VIPP1</i> -amiRNA #14	48	69
	<i>VIPP1</i> -amiRNA #18	63	50
	<i>VIPP1</i> -amiRNA #20	51	48
			n.d.

Cells were grown in TAP medium containing either ammonium or nitrate as nitrogen source at low light (LL) intensities of $\sim 30 \mu\text{E m}^{-2} \text{ s}^{-1}$ or at high light intensities (HL) of $\sim 1000 \mu\text{E m}^{-2} \text{ s}^{-1}$. Fifty electron micrographs of each line were analyzed. n.d., not determined.

which the biogenesis of photosynthetic protein complexes is arrested or at least retarded. The aberrant structures in *VIPP1*-RNAi/amiRNA cells were frequently (but not exclusively) located close to the pyrenoid in regions previously suggested to house translation zones (T-zones; Figure 10) (Uniacke and Zerges, 2007). T-zones were postulated to represent areas where de novo PSII biogenesis and the regulated transport of newly assembled PSII complexes to thylakoid membranes occur. If indeed the aberrant, PLB-like structures in *VIPP1*-RNAi/amiRNA cells correspond to T-zones, they might be caused by a problem during photosystem biogenesis/assembly.

In cyanobacteria, groups of thylakoids converge at peripheral cytoplasmic points without any apparent connection to the plasma membrane (Kunkel, 1982). Thylakoids at these points are attached to so-called thylakoid centers, which are 30 to 50 nm wide tubular structures of up to 1 μm length that are composed of subunits generating a 14-fold rotational symmetry (Kunkel, 1982; van de Meene et al., 2006). Thylakoid centers have been postulated to be linked to a membrane fraction with a density intermediate to that of thylakoid and plasma membranes (Hinterstoisser et al., 1993). It has also been postulated that these are the sites at which protein/pigment complexes are assembled and incorporated into photosynthetic membranes and that potentially are related to T-zones in *C. reinhardtii* (Nickelsen et al., 2011). A coincidence too striking to be overlooked is that recombinant VIPP1 from *Arabidopsis*, *C. reinhardtii*, and *Synechocystis* forms ring-like structures with an outer diameter of 25 to 37 nm, a 12- to 17-fold rotational symmetry, and the capacity to assemble into rod-like structures of up to 1.4 μm (Aseeva et al., 2004; Liu et al., 2007; Fuhrmann et al., 2009b). Hence, it is tempting to speculate that thylakoid centers in fact are VIPP1 rods, which again may correspond to microtubule-like structures reported in plastids of diverse algal and plant species in various types of tissues (Liu et al., 2007, and references therein). Thylakoid centers may have moved from a peripheral position in cyanobacteria to more central ones in chloroplasts and still serve as sites of thylakoid biogenesis. This speculation is supported by the localization of VIPP1 by immunofluorescence to distinct spots within the chloroplast that sometimes appear to extend into rod-like structures (Figure 9). Support also comes from the

biochemical localization of VIPP1 to low-density membranes (Liu et al., 2005) that are membranes intermediate between envelope and thylakoids at which thylakoid protein biogenesis might occur (Zerges and Rochaix, 1998). Depletion of VIPP1 may impair the formation of proper rods, thereby leading to disordered thylakoid centers and eventually resulting in the aberrant structures observed. Alternatively, rods may be formed normally but soluble VIPP1 is depleted and not available for chaperone-mediated cycling between soluble and complexed forms (Liu et al., 2007). If this process was required for the transport of building blocks to T-zones for thylakoid biogenesis (see below), the resulting tail-back may be the cause for the aberrant structures observed, comparable to the generation of PLB-like structures by the absence of chlorophyll in the γ -1 mutant.

Is VIPP1 Involved in the Biogenesis of Thylakoid Membranes or of Thylakoid Membrane Core Complexes?

The aberrant, PLB-like structures at the origin of thylakoids point to a deficit in thylakoid biogenesis. However, is the formation of the lipid bilayer itself affected, as proposed previously (Kroll et al., 2001; Westphal et al., 2001; Aseeva et al., 2007), or is the biogenesis/assembly of the photosynthesis complexes affected, as suggested by Gao and Xu (2009)?

Similar to what has been reported for a VIPP1 depleted *Arabidopsis* mutant (Kroll et al., 2001; Aseeva et al., 2007), we find no change in lipid composition, but we do observe slightly lower levels of fully assembled photosystems, cytochrome *b₆f* complex, and ATP synthase (by 14 to 20%) in nonstressed *C. reinhardtii* *VIPP1*-RNAi strains (Figure 8A; see Supplemental Figures 2 and 3 online). In contrast with the earlier reports, however, we find neither an obvious reduction of thylakoid membranes nor changes in the number of membranes per granum (Figures 3 and 10). We rather find levels of LHCII increased by 30%, which correlates with a lower chlorophyll *a/b* ratio and faster fluorescence induction kinetics (Figures 6D, 6G, and 8A). Hence, there appears to be no limitation of thylakoid membranes in *VIPP1*-RNAi/amiRNA strains for housing protein complexes, which points to a role of VIPP1 in the biogenesis/assembly of core complexes rather than in the formation of the membranes themselves. These data are in support of those from Gao and Xu (2009), who showed in *Synechocystis* that depletion of VIPP1 first affected PSII activity, then PSI activity, and once photosystem activities were lost, thylakoids degenerated. A role for VIPP1 in core complex biogenesis/assembly is also supported by the previous finding that VIPP1 interacts with Alb3.2 (Göhre et al., 2006). Alb3.2 was suggested to play a role in PSI and PSII assembly, as it was found to directly interact with photosystem subunits. Moreover, both photosystems accumulated to reduced levels when Alb3.2 was downregulated by RNAi, while cytochrome *b₆f* and ATP synthase were barely affected. Therefore, it appears possible that VIPP1 supports Alb3.2 during photosystem assembly.

Phenotypes in Stressed *C. reinhardtii* *VIPP1*-RNAi/amiRNA Strains Point to Defects in the Structural Organization of Thylakoid Membrane Complexes

If VIPP1 is involved in the biogenesis/assembly of thylakoid membrane core complexes, which step may be affected in

VIPP1 depleted cells? We believe that the pleiotropic phenotypes associated with VIPP1 depletion under stress conditions provide some hints to answer this question.

Prolonged exposure of cells to light intensities of $\sim 1000 \mu\text{E m}^{-2} \text{s}^{-1}$ resulted in a much more drastic decline of PSII and PSI activities in *VIPP1*-RNAi strains compared with a control strain. PSII activities were more severely and more rapidly affected than PSI activities. Loss of photosystem activity in *VIPP1*-RNAi strains was accompanied by photobleaching of chlorophyll and loss of subunits of both photosystems and of LHCII, while cytochrome *b₆f* complex and ATP synthase were hardly affected (Figure 1; see Supplemental Figure 4 online). Also, a short exposure to very high light intensities ($\sim 1800 \mu\text{E m}^{-2} \text{s}^{-1}$) resulted in more severe photoinhibition of *VIPP1*-RNAi strains and slower recovery compared with a control strain (Figure 4). Accordingly, *VIPP1*-RNAi strains under these conditions also suffered from a more drastic loss of PSII and PSI subunits than control cells (see Supplemental Figure 10A online).

The sensitivity of *VIPP1*-RNAi strains to both high light treatments was drastically increased in cells grown on ammonium instead of nitrate as nitrogen source and correlated with extensive thylakoid swelling (Figures 1, 3, and 5, Table 1; see Supplemental Figure 10 online). Thylakoid swelling is likely caused by osmosis where ammonium, accumulating in the acidified lumen, acts as an osmoticum and potentially also as a denaturing agent. Interestingly, induction of LHCSR3 and HSP90C/HSP70B was more strongly impaired in *VIPP1*-RNAi cells grown on ammonium compared with cells grown on nitrate. HSP70B was shown to be involved in photoprotection and repair of PSII during and after photoinhibition (Schroda et al., 1999), and LHCSR3 was shown to be essential for nonphotochemical quenching in *C. reinhardtii* (Peers et al., 2009). It is unclear how the thylakoid defects in the *VIPP1*-RNAi strains impair the expression of LHCSR3 and the chaperones, as the signals triggering their induction are still unknown. However, similar defects in *LHCSR3* induction and increased sensitivity to high light treatments, as described here for the *VIPP1*-RNAi strains, were recently reported for knock-down mutants of the chloroplast calcium sensor protein CAS (Petroustos et al., 2011). CAS is a thylakoid membrane protein and could be involved in a Ca^{2+} -dependent step of light stress signal transduction. Therefore, one possible scenario might be that thylakoid swelling in ammonium-grown *VIPP1*-RNAi cells either impairs CAS binding to the thylakoids or precludes the correct perception of the signal triggering the induction of photoprotective mechanisms.

In *C. reinhardtii*, thylakoid swelling was observed only in severely photoinhibited wild-type cells or in ATP synthase mutants (Topf et al., 1992; Majeran et al., 2001) (Table 1) (i.e., under conditions where a large $\Delta\text{pH}/\text{pmf}$ was generated in cells grown on ammonium). It is important to stress that thylakoid swelling occurs in *VIPP1*-RNAi strains exposed to high light intensities despite the fact that they are capable of generating only a lower pmf than control cells (Figure 6A). Hence, *VIPP1*-RNAi strains appear to have a structural defect in their thylakoid membranes that impairs their ability to withstand the osmotic pressure normally generated by the diffusion of water toward the ammonium accumulating in the acidified lumen. This structural defect is most likely also the cause for the drastically increased

thermosensitivity of PSII in *VIPP1*-RNAi strains compared with control cells (Figure 7A).

In conclusion, our compiled evidence suggests that VIPP1 depletion affects the structural organization of the thylakoid membrane complexes, in particular that of the photosystems. However, at which level could this structural defect occur? In grana stacks of higher plants, interactions between protein supercomplexes, especially PSII and LHCII, result in the formation of semicrystalline arrays (Kirchhoff et al., 2007; Daum et al., 2010). Comparable semicrystalline PSII arrays have also been observed in *Synechocystis* (Folea et al., 2008), but so far, their existence has not been addressed in *C. reinhardtii* thylakoids. A disturbance in the formation of such semicrystalline PSII-LHCII complexes should not cause obvious changes in thylakoid ultrastructure or in the assembly or basic function of thylakoid membrane protein complexes. However, such arrays are expected to contribute to the rigidity of thylakoid membranes. If a depletion of VIPP1 was causing a deficit in the formation of higher order structures like the semicrystalline PSII-LHCII arrays, this would explain the thylakoid swelling phenotype observed in high light-exposed *VIPP1*-RNAi/amiRNA cells grown on ammonium as nitrogen source. It would also explain the reduced cooperativity of PSII centers suggested by the less sigmoid fluorescence induction curves in *VIPP1*-RNAi cells compared with control cells (Figure 6G).

VIPP1 Might Support Alb3.2 in Photosystem Assembly by Delivering Small Hydrophobic Polypeptides or Lipids

From the interaction of VIPP1 with Alb3.2, the occurrence of aberrant structures at the origins of thylakoids, and the reduced accumulation of core complexes in *VIPP1*-RNAi/amiRNA cells, we propose a role for VIPP1 in the biogenesis/assembly of thylakoid membrane core complexes. From the thylakoid swelling phenotype in high light and the sensitivity of PSII to heat stress, we proposed that depletion of VIPP1 may lead to deficits in the structural organization of the thylakoid membrane complexes. But how can we bring both functions together, and how can we explain the increased high light sensitivity of nitrate-grown *VIPP1*-RNAi strains, which occurs independent of thylakoid swelling?

An explanation that may account for all observations made is that VIPP1 supports Alb3.2 during photosystem assembly by delivering small hydrophobic photosystem subunits or specific structural lipids. Small hydrophobic proteins may be loaded onto VIPP1 at the TIC complex in the chloroplast inner envelope, as suggested previously as a general pathway for thylakoid-located photosynthesis proteins (Jouhet and Gray, 2009a, 2009b). Specific lipids may be loaded onto VIPP1 directly at their site of synthesis at the inner envelope (Benning, 2008). Loading of small proteins/specific lipids onto VIPP1 and their release into an Alb3.2-containing photosystem assembly complex at thylakoid origins might be achieved by the chaperone-mediated assembly/disassembly of VIPP1 oligomers (Liu et al., 2007), which is supported by the finding that levels of VIPP1, CDJ2, and HSP70B were increased twofold in *Alb3.2*-RNAi strains (Göhre et al., 2006). In this scenario, the pleiotropic phenotypes associated with VIPP1 depletion would arise from photosystems containing reduced amounts of small subunits or structural lipids and therefore are incompletely assembled or impaired in the

formation of thylakoid membrane protein supercomplexes with the consequences described above.

The idea of VIPP1 being a carrier for small hydrophobic PSI and PSII subunits appears unlikely, as there exists no evidence that VIPP1 or PspA interact with proteins other than the HSP90C/HSP70B/CDJ2/CGE1 (co-)chaperones (Heide et al., 2009) or regulatory subunits of the *psp* operon, respectively (Joly et al., 2010). Moreover, it is difficult to imagine that VIPP1 would similarly exhibit specificity for small subunits of PSI and PSII. Finally, although levels of small subunits PSBR, PSBW, PSAG, PSAH, and PSAL appear to be lower under some conditions tested (see Supplemental Data Set 1 online), this is not consistently observed under all conditions, which would be expected if their delivery to the respective photosystem was generally perturbed. By contrast, there is no doubt that VIPP1 and PspA interact with membranes and a direct interaction of PspA with PG and phosphatidylserine was demonstrated (Kobayashi et al., 2007). Furthermore, at least one of the four lipid species present in thylakoid membranes (monogalactosyl-diacylglycerol [MGDG], digalactosyl-diacylglycerol [DGDG], sulfoquinovosyl-diacylglycerol [SQDG], and PG) was found to be an integral component of thylakoid membrane complexes.

More specifically, 11 MGDGs, seven DGDGs, five SQDGs, and two PGs were identified in the crystal structure of PSII from *Thermosynechococcus elongatus* (Guskov et al., 2009), and PSII isolated from spinach (*Spinacia oleracea*) and rice (*Oryza sativa*) was found to contain approximately three MGDGs, approximately two DGDGs, approximately one SQDG, and approximately two PGs (Sakurai et al., 2006). The two PGs were localized to the cytoplasmic side close to the Q_B site (Guskov et al., 2009), and reduced electron transfer rates from Q_A to Q_B upon PG depletion suggested a role of PG in maintaining the structural integrity of the Q_B site (Gombos et al., 2002). Moreover, PG was shown to be associated with the D1 protein and to be important for PSII dimerization (Kruse and Schmid, 1995; Kruse et al., 2000). Three PGs and one MGDG were identified in the crystal structure of PSI from *Synechococcus elongatus* (Jordan et al., 2001). One MGDG was found in the crystal structure of the cytochrome *b₆f* complex (Stroebel et al., 2003). Finally, one PG and one DGDG were found in LHCII, where PG is intimately involved in trimerization (Standfuss et al., 2005). Hence, VIPP1 would need to be specific for only a single lipid species to serve in the assembly of both photosystems.

If VIPP1 is indeed acting as a lipid chaperone during photosystem assembly, which lipid may it be supplying? There are three reasons to favor PG: (1) PspA interacts specifically with phospholipids and PG is the only phospholipid in the thylakoids. (2) PG is present in both photosystems. (3) The other three lipid classes are uniformly distributed in the membranes and PSII complexes, but PG is 2 to 5 times more abundant in PSII dimers (Sakurai et al., 2006; Domonkos et al., 2008), thus suggesting the need for an active enrichment process. In several studies, the synthesis pathway for PG was disrupted in *Synechocystis*, *C. reinhardtii*, or *Arabidopsis* (Domonkos et al., 2008). Strikingly, the resulting phenotypes to a large part resemble those observed in VIPP1-depleted lines presented in this and earlier studies: To start with *Synechocystis*, the *pgsA* and *cdsA* mutants, disrupted in genes encoding phosphatidylglycerolphosphate synthase and

cytidine 5'-diphosphate diacylglycerol synthase, respectively, died unless PG was added exogenously. Depletion of exogenously supplied PG led to the loss of PSII and PSI activities (Hagio et al., 2000; Sato et al., 2000, 2004) and thus is similar to what Gao and Xu (2009) observed as a consequence of VIPP1 depletion and what we observed in high light-treated *VIPP1*-RNAi cells. PG-depleted *Synechocystis* mutants were highly susceptible to photobleaching, which was explained by an increased formation of singlet oxygen as a consequence of reduced electron transfer from Q_A to Q_B that enhances the occurrence of the dangerous triplet state of chlorophyll (Sakurai et al., 2003; Domonkos et al., 2008). Moreover, loss of PSI activity upon PG depletion was found to come along with an accumulation of PSI monomers at the expense of PSI trimers (Domonkos et al., 2004), similar to the observation of Fuhrmann et al. (2009a) in a *vipp1* merodiploid strain.

The *Arabidopsis pgp1* mutant, defective in the gene encoding phosphatidylglycerolphosphate synthase and accumulating only 12% of wild-type PG levels, is unable to grow photoautotrophically, has pale, yellow-green leaves, and its chloroplasts contain only few, swollen thylakoid membranes without granum structure (Hagio et al., 2002). This phenotype indeed resembles that of the *Arabidopsis Δvipp1* mutant (Kroll et al., 2001; Aseeva et al., 2007).

Only the phenotype of the *C. reinhardtii mf1* and *mf2* mutants, in which the deficiency of PG leads to the nearly complete absence of PSII cores due to a severely reduced translation of D1 and CP47 (Maroc et al., 1987; Pineau et al., 2004), was much more severe than that observed in the *VIPP1*-RNAi strains reported here. Also not consistent with a simple lack of PG in PSII of *VIPP1*-RNAi cells is our finding that the midpoint potential of the Q_A/Q_A^- redox couple in PSII is lowered (Table 2). This points to a problem with the integrity of the Q_A site (Fufezan et al., 2007) instead of the Q_B site, which was affected by PG depletion in cyanobacteria (Gombos et al., 2002). Certainly, a more negative midpoint potential of the Q_A/Q_A^- redox couple has been shown to increase the yield of 1O_2 (Krieger-Liszkay and Rutherford, 1998; Fufezan et al., 2007) and most likely is causing the high light sensitivity of the *VIPP1*-RNAi cells. Phenotypes that very closely resemble those of the *VIPP1*-RNAi strains were observed in the *C. reinhardtii hf-2* mutant, which is defective in the biosynthesis of SQDG. The lack of SQDG had no obvious effect on the assembly of PSI, PSII, and LHCII. While PSI activity was unaffected, PSII activity was reduced by ~40% (Sato et al., 1995). Moreover, PSII activity was more sensitive to photoinhibition and highly sensitive to heat stress (Minoda et al., 2002; Sato et al., 2003). These compiled observations indicate that at least in *C. reinhardtii*, the phenotypes in *VIPP1*-RNAi/amiRNA cells cannot be explained by a simple lack of PG in the photosystems. However, they do point to a structural problem potentially caused by an incomplete or inaccurate incorporation of PG into the photosystems, thus producing PSII centers that are similarly labile as those completely devoid of SQDG.

A New Hypothesis for a Role of VIPP1 Different from That of a Vesicle-Inducing Protein

In summary, we propose here a new hypothesis for the function of VIPP1 in the chloroplast. This hypothesis claims that VIPP1

rods are in fact the thylakoid centers found in cyanobacteria and organize sites at which thylakoid biogenesis occurs. Chaperone-mediated dynamic cycling between soluble, smaller VIPP1 assembly states and rods may serve to provide structural lipids for core complex assembly, which occurs in collaboration with Alb3.2. Depletion of VIPP1 slows down this assembly process and leads to the aberrant PLB-like structures observed at the origin of thylakoid layers. Moreover, it leads to an incomplete/inaccurate incorporation of structural lipids into thylakoid membrane core complexes, in particular the photosystems. This directly impairs their structure and function (i.e., the integrity of the Q_A site in PSII) and affects the formation of supercomplexes, like the semicrystalline PSII-LHCII arrays, thus leading to a higher susceptibility to photoinhibition and reduced ability to withstand an increased luminal turgor. The pronounced high light inducibility of VIPP1 and the reduced capacity of *VIPP1*-RNAi cells to repair photodamaged PSII by de novo D1 synthesis also indicate a role of VIPP1 during the PSII repair cycle, where a supply of structural lipids will also be needed.

There is no doubt that this hypothesis is still speculative. However, in contrast with the earlier proposal that VIPP1 acts as a vesicle inducing protein, it accounts for most phenotypes observed in VIPP1-depleted cyanobacteria, algae, and higher plants and for most of the localization studies and biochemical and structural observations made so far. For certain, this hypothesis provides a basis for new experimental approaches to eventually understand the enigmatic process of thylakoid biogenesis.

METHODS

Algal Strains and Culture Conditions

Chlamydomonas reinhardtii strains cw15-325 (cw_d, mt⁺, and *arg7*⁻) and cw15-302 (cw_d, mt⁺, *nit1*⁻, and *arg7*⁻), both kindly provided by R. Matagne (University of Liège, Belgium), were used as recipient strains for transformation. Both strains were transformed with pCB412 (containing only the wild-type *ARG7* gene) to generate control strains. Strain cw15-325 was transformed with pMS439 (containing *ARG7* and the *VIPP1*-RNAi construct) and strain cw15-302 with pMS552 (containing *ARG7* and the *VIPP1*-amiRNA construct). pCB412 was linearized with *Xho*I, pMS439 with *Spe*I, and pMS552 with *Hind*III. One microgram of DNA was used for transformation with glass beads (Kindle, 1990). Strains were grown mixotrophically in TAP medium (Harris, 2008) supplemented with 7.5 mM NH₄Cl or 4 mM KNO₃ on a rotary shaker at 25°C at ~30 μE m⁻² s⁻¹. Fifteen hours before each experiment, cultures were diluted with TAP medium containing the respective nitrogen source to obtain equal cell densities of ~4 × 10⁶ cells/mL when the experiment started. Cell numbers were determined using a Z2 Coulter Counter (Beckman Coulter).

Vector Construction and Screening for VIPP1-Underexpressing Transformants

Construction of the *VIPP1*-RNAi vector (pMS439) was described previously (Heide et al., 2009). The miRNA targeting *C. reinhardtii* *VIPP1* was designed according to the detailed instructions given by Molnar et al. (2009) using the WMD2 tool at <http://wmd2.weigelworld.org> (Ossowski et al., 2008). Resulting oligonucleotides 5'-ctagtCGCGACCTGATTAAGAAGCGAtctcgtgatcgccatccatgggggtggtgatcagcagcctaTCGCATCTTAATCAGGTCGCG-3' and 5'-ctagcCGCGACCTGATTAAGATGCGAtagcgtgatcaccaccacccatggtgc-

cgatcagcagcagcTGCCTTCTTAATCAGGTCGCGa-3' (uppercase letters indicate miRNA/miRNA sequences) were annealed by boiling and slowly cooling down in a thermocycler and ligated into *Spe*I-digested pChlamiRNA2, yielding pMS552. Screening for correct constructs was done as described by Molnar et al. (2009). VIPP1-underexpressing transformants were initially identified by analyzing ~40 transformants by immunoblotting. After discovering their sensitivity to high light, candidates for VIPP1-underexpressing strains were selected among ~60 transformants that were streaked on TAP-NH₄ agar plates and showed bleaching after exposure to ~400 μE m⁻² s⁻¹ for 4 d. Strong reduction of VIPP1 levels in high light-sensitive transformants was then verified by immunoblotting.

Protein Gel and Gel Blot Analyses

Protein extraction and quantification, SDS-PAGE, semidry blotting, and detection by enhanced chemiluminescence were performed as described previously (Schroda et al., 1999). Antisera were against HSP90C (Willmund and Schroda, 2005), HSP70B (Schroda et al., 1999), VIPP1 (Liu et al., 2005), VIPP2 (S. Schmollinger and M. Schroda, unpublished results), LHCSR3 (Naumann et al., 2007), PsaD (Naumann et al., 2005), LHCII (Vallon et al., 1986), CP43 (Vallon et al., 1985), cytochrome *f* (Pierre and Popot, 1993), PsaA (Agrisera, AS05 084), PsaA (Agrisera, AS06 172), OEE33 (Agrisera, AS06 142-33), and CF1β (Lemaire and Wollman, 1989). Depending on the abundance of the antigen in the cell and the quality of the antiserum, different amounts of whole-cell proteins were loaded per SDS-polyacrylamide minigel: 15.3 μg for the detection of HSP90C, VIPP2, CP43, OEE33, PsaA, and cytochrome *f*; 3.8 μg for HSP70B, PsaD, and LHCSR3; and 0.8 μg for VIPP1, CF1β, D1, and LHCII.

RNA Gel Blot Analyses

RNA extraction, RNA gels, blotting, and hybridization were performed as described previously (Liu et al., 2005). Probes used for hybridization were the ~1-kb full-length cDNA of *CBLP2*, a 2-kb *Nhe*I-*Aat*II cDNA fragment containing the HSP70B coding region, a 1.4-kb *Bam*HI fragment containing part of the *VIPP1* coding region, and a 0.34-kb RT-PCR product containing part of the LHCSR3 coding region (Naumann et al., 2007).

High Light, Photoinhibition, and Heat Shock Experiments

For the exposure to high light (~1000 μE m⁻² s⁻¹), 150 mL of cell cultures were transferred to 400-mL beakers and placed onto spots on a rotary shaker that were illuminated by Osram HLX 250W 64663 Xenophot bulbs. A glass plate nonpermissive to infrared and UV irradiation was placed between light sources and beakers and cooled by ventilation. Beaker positions of all simultaneously treated cultures were swapped at regular intervals during the high light treatment to ensure equal illumination. The beakers were placed on a water-cooled metal plate to maintain a temperature of 25°C in the cell cultures. The same setup was used for photoinhibition, but cultures were exposed to ~1800 μE m⁻² s⁻¹. For heat shock treatments, cell cultures were incubated and agitated in a water bath at 40°C and ~5 μE m⁻² s⁻¹. Light intensities were determined using a Luxmeter (Heinz Walz Mess- und Regeltechnik).

Quantitative Proteomics

The control strain was grown in TAP supplemented with K¹⁵NO₃ (≥98%; NLM-765; Cambridge Isotope Laboratories), and *VIPP1*-RNAi strain #111 was grown in TAP supplemented with K¹⁴NO₃. Cells were grown to a density of ~4 × 10⁶ cells/mL. The cultures were split into three parts: One part was incubated at ~30 μE m⁻² s⁻¹ and 25°C for 1 h (unstressed), the second part was treated at ~2000 μE m⁻² s⁻¹ and 25°C for 1 h (photoinhibition), and the third part was exposed to 40°C at ~5 μE m⁻²

s^{-1} for 1 h (heat shock). After mixing ^{15}N -labeled with ^{14}N -labeled cells from each treatment at a ratio of 1:1 (based on same cell numbers), membrane-enriched fractions were prepared by freeze-thawing and centrifugation (30 min at 16,000g and 4°C). Pellets were resuspended in lysis buffer (50 mM NH_4HCO_3 , 1 mM DTT, and 1 mM NaCl), sonicated, and centrifuged again as above. The pellet was resuspended with lysis buffer, and proteins corresponding to 20 μ g chlorophyll were mixed with 2 \times SDS sample buffer (125 mM Tris-HCl, pH 6.8, 20% glycerol, 4% SDS, 10% β -mercaptoethanol, and 0.005% bromophenol blue), boiled for 60 s, and separated on a 12% SDS-polyacrylamide gel. The three lanes were cut into 14 gel bands each, and proteins therein were digested with 12.5 ng/ μ L sequencing grade modified trypsin (Promega) for 16 h at 37°C. The eluted peptides from the gel slices were desalted, separated, and measured by liquid chromatography–mass spectrometry essentially as described by Mühlhaus et al. (2011) with the following alterations: For HPLC separation a gradient ramping from 0 to 35% B (A, 2% acetonitrile and 0.5% acetic acid; B, 80% acetonitrile and 0.5% acetic acid) after 45 min and then to 100% B after 50 min was employed. Data were acquired in the positive polarity mode in data-dependent fashion: one survey scan from 300 to 2000 mass-to-charge ratio in the orbitrap analyzer (resolution of 60,000 at 400 mass-to-charge ratio; target filling of 10^6 ions) was followed by tandem mass spectrometry (MS/MS) scans in the linear ion trap (target filling of 10^4 ions) of the seven most intense precursor ions detected in the survey scan. Singly charged ions were rejected for MS/MS analysis, and we allowed two MS/MS events for each ion before putting it on the exclusion list for 180 s. Identification of peptide sequences and quantification were performed by the Integration of Mass Spectrometry Identification and Quantification Software (Mühlhaus et al., 2011). The light-to-heavy ratios calculated for all unique peptides were used to determine the relative abundance of the corresponding proteins. Unequal mixing of labeled and unlabeled proteins in each sample was corrected for by normalizing the light-to-heavy ratio determined for each peptide in a sample by the median of light-to-heavy ratios of all peptides in that sample. The median of the normalized light-to-heavy ratios of all peptides from one protein was taken as the relative abundance of that protein. Quantification values of core complex subunits are listed in Supplemental Data Set 1 online. Sequences and quantification values of all identified peptides are presented in Supplemental Data Set 2 online.

Chlorophyll a Fluorescence Measurements

To monitor PSII maximum quantum efficiency during high light exposure and heat stress, the F_V/F_M ratio was determined from intact cells at 25°C using the PAM-100 fluorometer (Heinz Walz). Basal fluorescence (F_0) was detected after dark adaptation for 2 min and far-red adaptation for an additional 2 min (the measurements of Figure 6 were all done with a Dual-PAM-100 after 5 min of far-red illumination, followed by 10 min of dark adaptation). F_M was determined with a single 4000 μ E $m^{-2} s^{-1}$ light pulse. For chlorophyll a fluorescence induction curves, cells equivalent to 10 μ g/mL chlorophyll were used. Cells were first illuminated with far-red light to induce state 1, followed by the addition of 100 μ M DCMU and incubation in darkness for 15 min. To keep cells in state 1, the sample was strongly stirred (aerobic conditions). Then, chlorophyll a fluorescence induction was measured at 25 μ E light intensity with a temporal resolution of 100 μ s.

Spectroscopy

The time course of changes in PSI and PSII activities during light exposure was followed using a JTS spectrophotometer (Bio-Logic). PSI and PSII activities were estimated from changes in the amplitude of the fast phase (100 μ s) of the electrochromic absorption shift (ECS) signal (at 520 to 545 nm) upon excitation with a saturating laser flash (Bailleul et al., 2010). The ECS spectral change linearly follows the number of light-induced charge

separations within the reaction centers (Witt, 1979). Thus, PSII contribution can be calculated from the decrease in the signal amplitude upon addition of DCMU (20 μ M) and hydroxylamine (1 mM) that irreversibly block PSII charge separation once the sample has been preilluminated (Joliot and Delosme, 1974). Conversely, PSI activity was estimated as the fraction of the signal that was insensitive to these inhibitors. For the measurement of pmf and ATP synthase activity, the ECS was used as an in vivo probe of the pmf across the thylakoid membrane (Kramer et al., 2003; Takizawa et al., 2007; Bailleul et al., 2010). The difference transmission signal was measured with a KLAS-100 LED-array spectrophotometer (Heinz Walz), which allows the simultaneous measurement of light-induced difference absorption signals at eight wavelength pairs between 505 and 570 nm. The first two wavelength pairs at 505 to 520 nm and 520 to 535 nm are dominated by the electrochromic shift transmittance changes arising from the xanthophyll cycle. The other wavelength pairs (540 to 545 nm, 545 to 550 nm, 550 to 554 nm, 554 to 559 nm, 559 to 563 nm, and 563 to 570 nm) are required to completely deconvolute the ECS signal from scattering effects, the C550 pheophytin signal, and from redox changes of the cytochromes f and b_6 . The deconvoluted transmission signals ($\Delta I/I$) were normalized to the chlorophyll content of the algal suspension. The maximum amplitude of the ECS (ECS_T) was used as a measure for the light-induced pmf across the thylakoid membrane. Cells were resuspended in 20% (w/v) Ficoll and illuminated for 5 min prior to each measurement to allow photosynthesis to reach steady state. ECS_T was determined after illuminating the suspension with saturating light (1500 μ E $m^{-2} s^{-1}$), which then was interrupted by a short interval of darkness (12 s), and the dark-interval relaxation kinetics of the ECS were measured. To determine ATP synthase activity, the fast phase (between 0 and 500 ms after the end of actinic illumination) of the dark interval relaxation kinetics was fitted with a single exponential decay function. The reciprocal value of the half-time of the ECS decay, the thylakoid conductivity (gH^+), is a measure of ATP synthase activity, as fast ECS decay is exclusively attributable to proton efflux through the ATP synthase (Cruz et al., 2001; Baker et al., 2007). Cytochrome f reduction was measured in parallel to the ECS decay kinetics. For cytochrome f oxidation measurements cell numbers equivalent to 15 to 20 μ g/mL chlorophyll were used. Cells were resuspended in 20% (w/v) Ficoll. Cells were preilluminated for 5 min to fully activated the Calvin cycle, followed by the addition of 10 μ M DCMU, to inhibit PSII. Cytochrome f was oxidized by a saturating red light pulse (1660 μ E $m^{-2} s^{-1}$).

Thermoluminescence Measurements

Thermoluminescence was measured with a home-built apparatus. To measure thermoluminescence arising from the $S_2Q_A^-$ charge recombination (Q-band), *C. reinhardtii* cells were incubated in the presence of 10 μ M DCMU for 5 min in the dark at 20°C and then flashed with one saturating xenon flash at -5° C. For luminescence detection, the samples were warmed at a constant rate (0.4°C s^{-1}) from 1 to 70°C.

Indirect Immunofluorescence Microscopy

Control transformants in the cw15-325 background were grown in TAP- NH_4 media and fixed and stained as described previously (Cole et al., 1998; Pérez-Pérez et al., 2010). The primary antibody was affinity purified from polyclonal anti-VIPP1 antiserum (Liu et al., 2005; Willmund and Schroda, 2005) and used in a 1:250 dilution. A fluorescein isothiocyanate-labeled goat anti-rabbit antibody (1:500; Sigma-Aldrich) was used as secondary antibody. Preparations were photographed on a DM6000B microscope (Leica Microsystems) with an ORCA-ER camera (Hamamatsu) and the Leica Application Suite Advanced Fluorescence software package (Leica).

Transmission Electron Microscopy

Cells were fixed for 2 h (at room temperature) to 14 h (at 4°C) in 100 mM sodium cacodylate containing 2.5% glutaraldehyde at pH 7.2. After two washes with 100 mM Na-cacodylate, pH 7.2, and two washes with distilled water, the cells were osmicated in 1% OsO₄ in distilled water for 1 h at 4°C. After two washes with distilled water, cells were resuspended in 20% BSA (w/v) and incubated for 20 min at room temperature. Cells were centrifuged for 10 min at 16,000g, and the cell pellet was incubated for 30 min at room temperature with 2.5% glutaraldehyde in distilled water at pH 7.4. After two washes with distilled water, the cell pellet was cut into pieces (1 to 2 mm³), dehydrated using increasing concentrations of ethanol and embedded in glycid ether 100 (formerly Epon 812; Serva) with propylene oxide as intermediate solvent following standard procedures. Polymerization was performed for 48 h at 65°C. Ultrathin sections (~60 to 70 nm) were cut with a diamond knife (type ultra 35°; Diatome) on an EM UC6 ultramicrotome (Leica) and mounted on single-slot Pioloform-coated copper grids (Plano). The sections were stained with uranyl acetate and lead citrate (Reynolds, 1963) and viewed with a JEM-2100 (Jeol) or EM 902A (Carl Zeiss) transmission electron microscope (both operated at 80 kV). Micrographs were taken using a 4080 × 4080-pixel or 1350 × 1040-pixel charge-coupled device camera (UltraScan 4000 or Erlangshen ES500W, respectively; Gatan) and Gatan Digital Micrograph software (version 1.70.16). Image brightness and contrast were adjusted and figures assembled using Adobe Photoshop 8.0.1.

Accession Numbers

Sequence data from this article can be found in the GenBank/EMBL data libraries under the following accession numbers: *VIPP1*, AAU06582 (GenBank), Cre13.g583550 (Phytozome); and *VIPP2*, Cre11.g468050 (Phytozome).

Supplemental Data

The following materials are available in the online version of this article.

Supplemental Figure 1. Construct Used for the Suppression of *VIPP1* by RNAi.

Supplemental Figure 2. Major Thylakoid Membrane Protein Complexes Are Fully Assembled in *VIPP1*-RNAi Strains.

Supplemental Figure 3. Bulk Membrane Lipid Composition in *VIPP1*-RNAi Strains Is Indistinguishable from That of Control Strains.

Supplemental Figure 4. Subunits of PSII and PSI Are Rapidly Degraded in *VIPP1*-RNAi Strains Exposed to High Light Intensities.

Supplemental Figure 5. High Light Sensitivity Is Also Observed in *VIPP1*-amiRNA Strains Unable to Induce *VIPP2* Expression.

Supplemental Figure 6. *VIPP1*-RNAi Strains Are Not Impaired in State Transitions.

Supplemental Figure 7. Delayed Bleaching of *VIPP1*-RNAi Strains Grown on Nitrate.

Supplemental Figure 8. Categories of Thylakoid Structure.

Supplemental Figure 9. Lysis of High Light-Exposed *VIPP1*-RNAi Cells Grown on Nitrate Appears Not to Be Caused by Thylakoid Swelling.

Supplemental Figure 10. *VIPP1*-RNAi Strains Are Less Sensitive to Photoinhibition When Grown on Nitrate Compared with Ammonium.

Supplemental Figure 11. *VIPP1*-RNAi Strains Are Less Sensitive to Heat Shock When Grown on Nitrate Compared with Ammonium.

Supplemental Data Set 1. Quantification Results for Subunits of Thylakoid Membrane and Respiratory Chain Core Complexes.

Supplemental Data Set 2. Sequences and Quantification Results for All Peptides Identified by Quantitative Shotgun Proteomics.

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AUTHOR CONTRIBUTIONS

A.N. performed all growth, protein gel, and RNA gel analyses with the cw15-325 strains. M.A.S., G.F., T.R., A.K.-L., A.N., S. Schönfelder, and H.L. performed the biophysical measurements. A.-K.U., B.S., and S.G. took all electron micrographs and did the statistical evaluations. S. Schönfelder and M.R. performed all growth and protein gel analyses with the cw15-302 strains. F.S. performed the lipid analyses and the mass spectrometry measurements, and T.M. evaluated the mass spectrometry data. S. Schmollinger cloned the *VIPP1*-amiRNA construct, generated recombinant *VIPP2*, and took immunofluorescence micrographs with J.L.C. M.S. conceived and coordinated the project, prepared most of the figures, and wrote the article.

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Evidence for a Role of VIPP1 in the Structural Organization of the Photosynthetic Apparatus in *Chlamydomonas*

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