A response regulator of cyanobacteria integrates diverse environmental signals and is critical for survival under extreme conditions

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ABSTRACT Microorganisms must sense their environment and rapidly tune their metabolism to ambient conditions to efficiently use available resources. We have identified a gene encoding a response regulator, NblR, that complements a cyanobacterial mutant unable to degrade its light-harvesting complex (phycobilisome), in response to nutrient deprivation. Cells of the nblR mutant (i) have more phycobilisomes than wild-type cells during nutrient-replete growth, (ii) do not degrade phycobilisomes during sulfur, nitrogen, or phosphorus limitation, (iii) cannot properly modulate the phycobilisome level during exposure to high light, and (iv) die rapidly when starved for either sulfur or nitrogen, or when exposed to high light. Apart from regulation of phycobilisome degradation, NblR modulates additional functions critical for cell survival during nutrient-limited and high-light conditions. NblR does not appear to be involved in acclimation responses that occur only during a specific nutrient limitation. In contrast, it controls at least some of the general acclimation responses; those that occur during any of a number of different stress conditions. NblR plays a pivotal role in integrating different environmental signals that link the metabolism of the cell to light harvesting capabilities and the activities of the photosynthetic apparatus; this modulation is critical for cell survival.

Microorganisms have developed diverse mechanisms for sensing and acclimating to changes in their environment (for reviews see refs. 1–4). Acclimation responses observed in photosynthetic organisms include the alteration of light harvesting complex synthesis and degradation in response to changes in light quality (5–7), light intensity (8), and nutrient availability (9–11). Such alterations help these organisms efficiently balance the absorption of excitation energy and the production of NADPH and ATP with their utilization for cell maintenance and growth. It is critical to maintain this balance because excess excitation of the photosynthetic reaction centers could result in the production of toxic oxygen species that would be damaging to many cellular processes (12).

Acclimation responses to nutrient availability can be characterized as those that are specific to the nutrient that is limiting, and those that are general and occur during any of a number of different nutrient limitation conditions. The specific acclimation responses encompass a number of metabolic changes that help the organism better scavenge the limiting nutrient from the environment. These responses include increased synthesis of high affinity transport systems (13–16) and the production of hydrolytic enzymes that allow utilization of alternate forms of the nutrient (e.g., synthesis of extracellular phosphatases and sulfatases in response to phosphorus and sulfur limitations, respectively) (17–19). The general acclimation responses that occur during nutrient starvation include the cessation of cell division and dramatic alterations in cellular structure and metabolism (10, 11, 20, 21). One of the general responses that has been studied in the cyanobacterium Synechococcus sp. strain PCC 7942 involves the degradation of the phycobilisomes (PBS), pigment-protein complexes that harvest most of the light energy for cyanobacteria (9–11, 22, 23).

The PBS give cyanobacterial cells their typical blue-green color. Imposing nitrogen or sulfur deprivation on cultures of Synechococcus sp. strain PCC 7942 triggers the rapid and complete degradation of the PBS, which causes the cells to look yellow or bleached. Therefore, mutants that fail to degrade the PBS can be identified visually. Analysis of such mutants should define the mechanism of PBS degradation and identify regulatory elements that control proteolysis or other modifications that may occur during nutrient-limited growth.

Mutants of Synechococcus sp. strain PCC 7942 that were unable to degrade their PBS during sulfur- or nitrogen-limited growth previously have been isolated (22). The mutant phenotype was complemented with the nblA (nonbleaching) gene, which encodes a polypeptide of 59 amino acids. The transcription of nblA was dramatically induced by sulfur and nitrogen starvation and, to a lesser extent, by phosphorus starvation. NblA does not show homology to a polypeptide of known function and its role in the mechanism of degradation has not been clarified.

To further study the mechanism of PBS degradation and its regulation by environmental conditions we isolated additional nbl mutants of Synechococcus sp. strain PCC 7942. Because bleaching during nutrient stress is a component of the general responses (see above), characterization of such mutants provides insights into regulatory pathways that are governed by diverse environmental factors. In this study we characterized an nbl mutant that has a lesion in a gene encoding a response regulator. This response regulator is involved in the integration of the cells responses to both nutrient deprivation and light intensity and is essential for survival during stress conditions.

MATERIALS AND METHODS

Strain, Culture Conditions, and Quantitation of Pigments. Synechococcus sp. strain PCC 7942 was cultured and starved for sulfur or nitrogen essentially as previously described (10). Briefly, cells were harvested by centrifugation (5,000 × g, 10 min) during exponential growth, resuspended in an equal volume of wash medium (BG-11 medium lacking nitrogen and sulfur), repelleted by centrifugation, and resuspended in wash...
medium to 1/10 the original culture volume. This concentrated cell culture was used to inoculate medium lacking either sulfur or nitrogen to a cell density of approximately 5 × 10^7 cells/ml. Except for high light conditions (500 μmol photons m^-2 s^-1), the cells were illuminated at an intensity of 70 μmol photons m^-2 s^-1 from incandescent bulbs. Low-sulfur agar plates were prepared as described previously (22). When appropriate, antibiotics were added to final concentrations of 2 μg/ml of ampicillin, 25 μg/ml of spectinomycin, and 25 μg/ml of kanamycin. Chlorophyll (chl) and phycocyanin (PC) were quantified as described previously (10).

Mutagenesis. The nonbleaching mutant nblR (for non-bleaching regulatory) was fortuitously isolated from a screen for genes involved in Ca^2+ homeostasis. *Synechococcus* sp. strain PCC 7942 in which pacL (a gene encoding Ca^2+ ATPase) had been inactivated by insertion of a spectinomycin cassette (24) was mutagenized with N-methyl-N′-nitro-N-nitrosoguanidine (22). The mutagenized cells were screened for a growth minus phenotype in the presence of an elevated concentration of Ca^2+ (20 mM). However, it was noted that the pacL mutant, as well as wild-type cells, appeared significantly bleached when grown on solid medium (but not liquid) containing 20 mM Ca^2+. This bleaching was shown to be a consequence of elevated expression of nblA, as determined by measuring the levels of β-glucuronidase (GUS) activity expressed from an nblA-promoter-GUS fusion in wild-type cells exposed to high levels of Ca^2+ (unpublished data). Elevated Ca^2+ levels in solid growth medium may decrease the availability of sulfate to the cells, which in turn would activate nblA and result in partial bleaching. While screening for mutants of the pacL strain that were unable to grow on 20 mM Ca^2+, we obtained the nblR mutant, which exhibited an nbl phenotype during sulfur- or nitrogen-limited growth.

Chimeric GUS Reporter Gene and Quantitation of GUS Activity. Wild-type and mutant strains were transformed with an autonomously replicating vector, pCB4’ (22), containing a chimeric gene in which the nblA promoter was fused to a promoterless GUS gene (uidA, but designated GUS throughout). This chimeric gene contained 420 bp upstream of the ATG of nblA, the first three codons of nblA (for Met, Leu, and Pro) and a nine-codon linker region (encoding for Arg, Ile, Pro, Gly, Tyr, Gly, Gin, Ser, and Leu) derived from the polylinker of pCB4’ that precedes the coding region of GUS. Expression from this chimeric gene was demonstrated to strongly depend on the nitrogen and sulfur levels of the medium. Cultures grown in nutrient-repleted medium were washed as described above and inoculated into replete, sulfur-depleted, or nitrogen-depleted medium. GUS activity was measured after 24 h of starvation, essentially as described elsewhere (25).

RNA Isolation and Northern Hybridization Analysis. RNA was isolated as previously described (22) except that the cell suspension was vortexed for 1 min in the presence of glass beads (about 100 mg, 100–300 microns, Sigma) before adding the hot phenol. This significantly improved the yield of RNA. Northern blot hybridizations and the preparation of radiolabeled RNA and DNA probes were performed as previously described (10, 22, 26).

DNA Manipulation, Complementation of Cyanobacterial Mutants, and Sequence Analysis. Molecular techniques were performed according to standard procedures (27). Genomic DNA isolated from *Synechococcus* sp. strain PCC 7942 in which a spectinomycin cassette was inserted at the BglII site of nblA (22) was used to construct a genomic library. A Sau3AI partial digest of this genomic DNA was fractionated on a 0.7% agarose gel and 7- to 10-kbp fragments were isolated and ligated into the BamHI of pUC118. This plasmid cannot replicate autonomously in *Synechococcus* sp. strain PCC 7942, but can confer ampicillin resistance to the cyanobacterium upon homologous integration into the genome. After integration, a complemented strain contains both a wild-type and a mutated copy of the complementing sequence. Complementation of the nblR mutant was achieved by transformation with the plasmid library and screening for ampicillin-resistant colonies that bleached normally when grown on low-sulfur plates (yellow colonies in a background of blue-green colonies). After several rounds of rescreening to ensure genetic homogeneity, genomic DNA was isolated from the complemented strain, digested with restriction enzymes for which there are sites on the polylinker of pUC118, and analyzed by Southern blot hybridizations using pUC118 as a probe. The EcoRI and *Pst*I digest yielded 8.0- and 5.5-kbp fragments, respectively, each containing pUC118 and flanking genomic DNA. To clone the DNA flanking the vector sequence, total genomic DNA isolated from complemented nblR was digested with EcoRI or *Pst*I, the restriction fragments were diluted and incubated with T4 DNA ligase, and the ligation products were transformed into *Escherichia coli* cells that were selected on solid medium containing 100 μg/ml of ampicillin. Each of the rescued plasmids was tested for its ability to complement nblR. The nblR strain transformed with the plasmid rescued from *Pst*I digested genomic DNA bleached on solid medium containing low levels of sulfur. The insert in this plasmid (2.5 kbp) was sequenced by using the Applied Biosystems PRISM system (Perkin–Elmer) and was used to generate subclones tested in the complementation assay. The sequence data have been submitted to the GenBank database under accession number AF049128.

Interposon Inactivation. nblR was interrupted by insertion of a 1.3-kbp kanamycin resistance gene at the XcmI site located 87 nucleotides from the beginning of the gene. The plasmids with the interrupted gene were introduced into wild-type *Synechococcus* sp. strain PCC 7942, and *in vivo* gene disruptions were verified by PCR analysis of genomic DNA isolated from the transformant.

RESULTS

Characterization of the nbl Mutant. Fig. 1A shows cultures of *Synechococcus* sp. strain PCC 7942 and the nblR mutant, nblR, after growth for 48 h in medium lacking sulfur. The wild-type culture exhibited the typical yellow color that indicates the degradation of the PBS (Fig. 1A). This observation is reflected in the absorbance spectra (Fig. 1C) as a dramatic decrease in

![Fig. 1](image-url)
the absorbance at 620 nm, a consequence of the loss of PC, the major pigment-protein of the PBS (6, 9, 10, 28). In contrast, mutant nblR exhibited a blue-green color (Fig. 1A) and retained most of the absorbance peak at 620 nm after starvation for sulfur or nitrogen (Fig. 1C). Furthermore, the nblR mutant did not exhibit (not shown) the partial loss of PBS constitutents that is observed in wild-type cells during phosphorus limitation (10).

To determine whether the accumulation of the nblA transcript was altered in the mutant, we examined the transcript levels by Northern blot hybridizations. As shown in Fig. 2A, mutant nblR had dramatically reduced levels of the nblA transcripts during sulfur and nitrogen limitation. In contrast, as shown in Fig. 2B, expression of genes such as rhdA that are activated by specific nutrient limitation conditions (26) do not appear to be altered in the nblR mutant.

To quantitate expression from the nblA promoter, we transformed both the wild-type and the nblR mutant cells with a plasmid bearing a chimeric gene consisting of the nblA promoter fused to the promoterless uidA gene encoding GUS (see Materials and Methods). GUS activity in the strains harboring the chimeric gene was measured after growth in replete medium or after growth for 24 h in medium devoid of sulfur or nitrogen. For wild-type cells, sulfur and nitrogen starvation resulted in a 20- and 30-fold increase in GUS activity, respectively, whereas mutant nblR showed only a small increase in GUS activity after starvation (Fig. 2C). Reduced expression from the chimeric gene in mutant nblR suggested that the nblA promoter was less active in the mutant than in the wild-type genetic backgrounds. Hence the nblR strain probably harbors a lesion in a gene required for transcriptional control of nblA. Furthermore, GUS activity in the nblR mutant was lower compared with that of wild-type cells even in cultures grown in nutrient-replete medium (Fig. 2D). These findings are consistent with the observation that the nblR mutant grown in complete medium exhibited a more “bluish” pigmentation (Fig. 1B) and contained 30–50% more PC/cell than wild-type cells, as calculated from whole-cell absorption spectra (Fig. 1C).

**Complementation of nblR and Analysis of the Mutated Gene.** To identify the lesion in nblR, this strain was transformed with a recombinant library of *Synechococcus* sp. strain PCC 7942 genomic DNA, and transformants were screened for bleaching during sulfur limitation. Transformation of mutant nblR with the nblA gene (integrated into the genome by homologous recombination) resulted in partial suppression of the nbl phenotype, although the nblA gene was not altered in the mutants as determined by sequence analysis. Therefore the recombinant library was constructed with genomic DNA from a strain in which nblA had been inactivated (nblAΔ1). A plasmid rescued from a transformant of nblR that was phenotypically complemented was able to reconstitute the mutant phenotype; absorbance spectra of the strain transformed with this plasmid indicated a dramatic decrease in the PC level after sulfur and nitrogen starvation (not shown). Complementation with subclones of this plasmid (not shown) defined the nblR gene, which encoded a protein with similarities to response regulators of two component signal transduction pathways. Fig. 3 shows the homology between NblR and several response regulators, including Sil0396 of *Synechocystis* sp. strain PCC 6803 (29), CopR from *Pseudomonas syringae* (30), PhoB from *E. coli* (31), SphR, the PhoB homologue in *Synechocystis* sp. strain PCC 7942 (32, 33), and OmpR from *E. coli* (34, 35). These response regulators, excluding Sil0396 that has not been studied in detail, were shown to activate transcription of genes involved in the acclimation of bacterial cells to various stress conditions. For example, PhoB activates transcription of the *pho* regulon, which is composed of several operons important for scavenging phosphorus from the environment (4). CopR is involved in the acclimation of cells to elevated levels of copper (30) and OmpR regulates transcription of the genes encoding the major outer membrane porin proteins in response to changes in osmolarity (36). The sequence of the nblR gene from the mutant strain revealed a point mutation that resulted in the replacement of threonine at position 156 with an isoleucine (T156I, Fig. 3).

Because nblR was obtained by mutagenesis of a strain in which the pacL gene (encoding Ca²⁺ ATPase), was inactivated (see Materials and Methods), it was possible that the nbl phenotype resulted from the combination of the two defective genes, pacL and nblR. Therefore, nblR of wild-type cells was disrupted by the insertion of a kanamycin-resistant gene. The resulting mutant, nblRΔ, had a pronounced nbl phenotype upon either sulfur or nitrogen starvation and contained even more PC than the nblR mutant during nutrient-replete growth (not shown). Hence, a lesion in nblR alone is sufficient to cause the nbl phenotype.

**Survival of the nblR Mutant During Nutrient Limitation and Exposure to High Light.** In addition to providing cells with a certain amount of the limiting nutrient, PBS degradation could prevent excess excitation of photosystem II reaction centers, which might be critical for surviving starvation conditions (see Discussion). To explore this question, wild-type cells and the nblRΔ1 mutant were assayed for their ability to survive nitrogen and sulfur starvation in liquid medium. The nblA-inactivated strain (nblAΔ1) was also examined to determine whether retention of the light-harvesting complex during nutrient limitation affects survival in the presence of functional NblR. After various times of starvation, aliquots of cells were removed, spotted onto nutrient-replete solid medium, and allowed to grow. Fig. 4 shows that the nblRΔ1 mutant dies rapidly during sulfur and nitrogen starvation. In contrast, the nblAΔ1 mutant survives nutrient deprivation nearly as well as wild-type cells, although there is some decreased viability. Hence, in addition to regulating nblA expression and PBS degradation, NblR functions to regulate other genes during

![Fig. 2.](image-url)
To examine whether the pleiotropic nature of the nblR mutation is the result of a polar effect on downstream genes, we inserted nblR into a neutral site in the genome of the nblR mutant. This action resulted in complementation of all aspects of the mutation, including the degradation of the PBS and altered viability during nutrient-limited and high-light growth (not shown). This observation excludes the possibility of polar effects and suggests that NblR modulates acclimation responses in addition to the degradation of the PBS.

**DISCUSSION**

NblR Integrates Diverse Environmental Signals. The cyanobacterial mutant, nblR, was isolated based on its inability to degrade PBS during sulfur and nitrogen starvation. The complementing gene encodes a response regulator of two-component signal transduction pathways (Fig. 3), which is essential for the activation of transcription of nblA during sulfur and nitrogen starvation (Fig. 2 A and C). In addition, NblR modulates phycobiliprotein levels and the activity of the nblA gene during nutrient-replete growth (Figs. 1C and 2D). Furthermore, the nblR mutant did not exhibit the partial loss of PC observed when wild-type cells are starved for phosphorus (not shown) or exposed to high light (Fig. 5A). Hence, the nblR signal transduction pathway must integrate a number of different environmental signals.

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**Fig. 3.** Alignment of NblR with various response regulators of two component signal transduction systems. (A) NblR. (B) sll0396 (putative response regulator in *Synechocystis* sp. strain PCC 6803). (C) CopR (copper-related response regulator from *Pseudomonas syringae*). (D) PhoB (phosphorus-related response regulator from E. coli). (E) SphR (PhoB homolog of *Synechococcus* sp. strain PCC 7942). (F) OmpR (osmolarity-related response regulator of E. coli). Identical and similar residues in at least two of the sequences are on a black or gray background, respectively. The threonine (T) to isoleucine (I) change at amino acid 156 and α-helical regions (a1, a2, a3) and β-sheets (β1, β2) close to the helix-turn-helix (H, T, H) of the predicted DNA binding site are indicated.

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**Fig. 4.** Whole-cell absorbance spectra of wild type (WT), nblRΩ (RΩ), and nblAΩ (AΩ) grown at medium light (ML) or high light (HL) (70 and 500 μmol photons m$^{-2}$s$^{-1}$, respectively). Cultures were adjusted to the same OD$_{750}$ before determining the spectra. The spectra were offset along the y axis. Arrows indicate the absorbance peaks of PC and chl. (B) Growth, as measured by OD$_{750}$, of WT, nblRΩ, and nblAΩ at HL (500 μmol photons m$^{-2}$s$^{-1}$).

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**Fig. 5.** (A) Whole-cell absorbance spectra of wild type (WT), nblRΩ (RΩ), and nblAΩ (AΩ) grown at medium light (ML) or high light (HL) (70 and 500 μmol photons m$^{-2}$s$^{-1}$, respectively). Cultures were adjusted to the same OD$_{750}$ before determining the spectra. The spectra were offset along the y axis. Arrows indicate the absorbance peaks of PC and chl. (B) Growth, as measured by OD$_{750}$, of WT, nblRΩ, and nblAΩ at HL (500 μmol photons m$^{-2}$s$^{-1}$).

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**Fig. 6.** Wild type (WT), nblRΩ (RΩ), and nblAΩ (AΩ) were spotted onto nutrient-replete solid medium after various times (24, 48, and 72 h) of starvation for sulfur (−S) or nitrogen (−N) in liquid medium. Each spot consisted of 10 μl from a 2-fold dilution of the original culture. Where indicated, the cultures were made 5 μM of DCMU upon initiation of starvation (t₀).
Specific regulatory proteins have been associated with the acclimation of microbes to nutrient starvation. Genes identified in cyanobacteria that encode such regulators are ntcA and ntcB (37, 38), cysR (39), sphR (32), and phoB (40). NtcA and NtcB are involved in regulation of nitrogen transport and metabolism, CysR regulates genes that respond to sulfur availability, and SphR and PhoB modulate acclimation to phosphorus limitation. In contrast to nutrient-specific responses, there are general responses, such as PBS degradation, that are triggered by changes in any of a number of different environmental parameters.

The sulfur-specific responses, mostly involved in scavenging the limiting nutrient, are not significantly affected by a mutation in nblR. The nblR mutant shows normal induction of transcription of nblA (Fig. 2B), which encodes a rhodanese-like polypeptide (26), and cysW (data not shown), which encodes one of the pore-forming proteins of the sulfate permease (41), after sulfur starvation. The nitrogen-specific responses are also likely to be normal in the nblR mutant. In *Synechococcus* sp. strain PCC 7942, most of the genes that respond to environmental levels of nitrogen-containing metabolites become active upon ammonium deprivation in the presence of nitrate (42). The nblR mutant grows normally in medium that contains nitrate as the sole nitrogen source, suggesting that elimination of NblR does not prevent the activation of genes encoding nitrate transport and assimilation functions. In sum, NblR does not appear to affect nutrient-specific responses, but is a pivotal regulatory element in a signal transduction pathway thatmodulates at least some of the general responses. Aspects of the general pathway controlled by NblR are the complete degradation of the PBS during sulfur and nitrogen starvation (Fig. 1C), the partial loss of pigmentation that accompanies phosphorus limitation (not shown) and exposure to high light (Fig. 5A), and the survival of cells starved for sulfur, nitrogen or exposed to high light (Figs. 4 and 5B, respectively). Because the nutrient-specific acclimation responses appear normal in the nblR mutant, decreased viability during starvation conditions is not a result of the inability of this strain to scavenge the limiting nutrient. Elements that regulate gene expression in response to multiple environmental factor have been found in other organisms; these include SNF1, which controls the responses of yeast to both phosphorus and glucose depletion (43).

**NblR Controls Expression of nblA.** Response regulators such as NblR often work in conjunction with a sensor protein with histidine kinase activity. Changes in the environment can trigger autophosphorylation of the sensor at a specific histidine residue. The phosphoryl group on the histidine then is transferred to an aspartate residue in the receiver domain of a response regulator. The phosphorylated response regulator then may modify the transcriptional activity of specific groups of genes (for reviews see refs. 1, 3, and 44). The strong similarity between NblR and OmpR provides some insights into the functions of specific domains of NblR. Based on the crystal structure, the C-terminal domain of OmpR consists of three α-helices packed against two antiparallel β-sheets (indicated in Fig. 3 below OmpR sequence). Helices α2 and α3 of OmpR, plus residues in the loop connecting them, constitute a variation of the helix–turn–helix (HTH) motif, with α3 being the DNA recognition helix (45). A HTH structure is predicted for the analogous region of NblR (Fig. 3, Chou-Fasman analysis). In the nblR mutant, the threonine residue located within the α1 helix, is mutated to an isoleucine (T156I, see Fig. 3). The equivalent mutation in OmpR (T162I) completely eliminates DNA binding (46). Based on the predicted three-dimensional structure of OmpR, T162 is in close proximity to helix α3 and may facilitate interactions of that helix with DNA. A similar role of T156 in NblR would explain why the T156I nblR mutant was unable to fully activate *nblA*. The isoleucine residue, unlike the threonine residue, does not have a hydroxyl group, which may function in DNA-protein interaction. Furthermore, the isoleucine residue is larger and more hydrophobic, which might result in steric retardation of NblR-DNA association. Additional evidence suggesting the functional importance of amino acid 156 in NblR comes from mutational studies of PhoB from *E. coli* (47).

**NblR Is Critical for Survival During Starvation and Exposure to High Light.** A mutant in *nblR* dies rapidly during sulfur and nitrogen starvation or upon exposure to high light. There are several reasons that could explain why the *nblR* strain is more sensitive to nutrient limitation than wild-type cells. Starvation of wild-type cells for sulfur and nitrogen triggers the disconnection of the PBS from photosystem II reaction centers, the degradation of the light-harvesting complex, and the down-regulation of photosystem II activity (22, 48). The finding that the nbl mutant strain, specifically impaired in PBS degradation, dies slightly more rapidly than wild-type cells suggests that PBS degradation has an effect on viability during acclimation to nutrient starvation. However, *nblR* dies much more rapidly than *nblA*, indicating that additional processes controlled by NblR are critical for viability during starvation. The rapid death of *nblR* during starvation makes it difficult to determine which of the general acclimation responses is impaired. However, because the photosynthetic electron transport inhibitor DCMU markedly improved survival of *nblR* during starvation (Fig. 4), it is likely that the mutant strain cannot down-regulate photosynthetic electron transport during nutrient starvation. Additional evidence for the importance of the down-regulation of photosystem II during nutrient starvation comes from studies of a mutant of *Chlamydomonas reinhardtii*, *sac1*, that is unable to acclimate to sulfur limitation (49). This strain cannot modify a number of aspects of photosystem II activity (21) and dies more rapidly in the light during sulfur deprivation than wild-type cells (49). As with *nblR*, the death of the *sac1* strain can be retarded by the addition of DCMU (49).

NblR is critical for the acclimation of *Synechococcus* sp. strain PCC 7942 to high light. The mutants *nblA* and *nblR* show much less of a decline in the levels of high-light-harvesting pigment during illumination with high light than wild-type cells (Fig. 5A). However, while *nblA* grows in high light, the *nblR* mutant does not grow (Fig. 5B) and rapidly dies (not shown). Hence, NblR modulates acclimation responses, in addition to the degradation of the PBS, that are essential for survival of cyanobacterial cells in high light. Although it is unclear what processes are controlled by NblR that allow survival in high light, they may include (i) modulation of energy transfer from the light-harvesting complex to the photosystem II reaction center, (ii) dissipation of excess absorbed light energy, (iii) repair of photosystem II reaction centers that suffer photoinhibitory damage, and (iv) detoxification of active oxygen species formed in high light. The inability of the *nblR* mutant to modulate one or more of these processes during exposure to high light could result in elevated photodamage and death.

The mechanism by which high light and nutrient deprivation control NblR activity is uncertain; however, both would create a highly reduced intracellular environment. During nutrient limitation, when the anabolism of the cell is slowed down or completely arrested, NADPH, the final electron acceptor of the photosynthetic electron transport chain, is not recycled as fast as under the analogous conditions and the electron carriers are maintained in a relatively reduced state. Retention of the PBS and light absorption during nutrient-limited and high-light conditions could further reduce the electron carriers and result in photodamage. Therefore, aside from providing the cell with some of the limiting nutrient, the degradation of the PBS might help minimize the absorption of excess excitation energy.

The redox state of the photosynthetic electron carriers is known to modulate a large number of acclimation responses...
including induction of transcription (50), regulation of translation (51), state transitions, and changes in the stoichiometry of the photosystems (52). The possibility that NbIR activity is linked to the redox state of the photosynthetic electron carriers is supported by the finding that DCMU, which prevents photosynthetic electron flow beyond QA, inhibits the accumulation of nblA mRNA in sulfur-starved cells (not shown).

Different environmental stimuli govern NbIR activity, which in turn, activates nblA transcription and PBS degradation. In addition to the environmental signals discussed above, low ambient CO2 is postulated to affect NbIR, based on the partial bleaching observed under this condition (R.S. and A. Kaplan, unpublished results). PBS degradation contributed to a small extent to cell survival during nutrient stress but did not appear to be critical for survival under the high light conditions of 2 mol photons m\(^{-2}\)s\(^{-1}\). Reduced PBS levels might become more critical when the cells experience higher light intensities or when exposed to a combination of stress conditions. Independent of its effects on nblA and PBS degradation, NbIR must alter the expression of other genes critical for survival in high light or during extended periods of nutrient deprivation. Although it is unknown what specific features of high light and nutrient limitation control NbIR activity, both conditions would lead to an increase in the intracellular redox state. Future work will help elucidate how NbIR integrates multiple environmental cues and triggers a set of responses that allow the organism to better compete under suboptimal growth conditions.

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