Chlorophyll biosynthesis under the control of arginine metabolism

Graphical abstract

Highlights

- ArgD of arginine pathway interacts with the Gun4 protein of chlorophyll biosynthesis
- Ornithine stimulates the ArgD-Gun4 interaction and inhibits tetrapyrrole synthesis
- The inhibition by ornithine is released in a strain lacking the ArgD-Gun4 complex

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In brief
Kiss et al. demonstrate that the synthesis of the main photosynthetic pigment, chlorophyll, and the metabolism of arginine (managing N-stockpile) are functionally coupled via protein-protein interactions in cyanobacteria.
Chlorophyll biosynthesis under the control of arginine metabolism

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SUMMARY

In natural environments, photosynthetic organisms adjust their metabolism to cope with the fluctuating availability of combined nitrogen sources, a growth-limiting factor. For acclimation, the dynamic degradation/synthesis of tetrapyrrole pigments, as well as of the amino acid arginine, is pivotal; however, there has been no evidence that these processes could be functionally coupled. Using co-immunopurification and spectral shift assays, we found that in the cyanobacterium Synechocystis sp. PCC 6803, the arginine metabolism-related ArgD and CphB enzymes form protein complexes with Gun4, an essential protein for chlorophyll biosynthesis. Gun4 binds ArgD with high affinity, and the Gun4-ArgD complex accumulates in cells supplemented with ornithine, a key intermediate of the arginine pathway. Elevated ornithine levels restricted de novo synthesis of tetrapyrroles, which arrested the recovery from nitrogen deficiency. Our data reveal a direct crosstalk between tetrapyrrole biosynthesis and arginine metabolism that highlights the importance of balancing photosynthetic pigment synthesis with nitrogen homeostasis.

INTRODUCTION

In natural environments, the fluctuating combined nitrogen (N) source is a growth-limiting factor. To cope with this constraint, photosynthetic organisms have to adjust their metabolism to the availability of N. For acclimation, the dynamic degradation/synthesis of tetrapyrrole pigments (chlorophyll [Chl], bilins, hemes) as well as of the amino acid arginine (Arg) are evidently pivotal. Arg metabolism is in aid of balancing N availability for anabolic processes in a fluctuating environment. The surplus N is primarily used for the synthesis of Arg that can be stockpiled in storage material, such as cyanophycin, which can be metabolized during N deficiency. At the same time, N fluctuation causes extensive changes in the metabolism of tetrapyrroles. At low N, the accumulation of Chl and bilins is predominantly downregulated in a process called chlorosis or bleaching. Since these tetrapyrroles are essential co-factors for photosynthetic apparatus, bleached cyanobacterial cells stop their photosynthetic activity. When N becomes available, pigment synthesis is reactivated (regreening) for the biogenesis of photosynthetic machinery to allow photoautotrophic growth.

Oxygenic phototrophs synthesize the main photosynthetic pigment, Chl, together with other tetrapyrroles via a common, branched pathway. The tetrapyrrole pathway must be tightly regulated in virtually any type of organism since protoporphyrins and other pyrrolic intermediates are highly phototoxic. In oxygenic phototrophs, the regulation of the tetrapyrrole pathway appears to be a particularly complicated task, as the pathway is branched and high quantities of end products (Chl, heme, bilins) are required for photosynthesis. A sophisticated control mechanism(s) must evolve to counterbalance the pathway with other processes in the cell. In this way, the amounts of tetrapyrrole end products can vary depending on developmental stage and/or environmental conditions, and the accumulation of the phototoxic biosynthetic intermediates can be tightly coordinated. How this regulation is accomplished remains mostly unclear. However, the main regulatory mechanisms appear linked to the formation of the rate-limiting precursor 5-aminolevulinic acid (5-ALA) and the branching between Chl and heme pathways. At this branchpoint, the magnesium chelatase (MgCh) and ferrochelatase enzymes compete for the same protoporphyrin IX (PPIX) substrate.

The Gun4 protein has been recognized as a critical factor for the synthesis of Chl in cyanobacteria, as well as in algae and plants, and has been extensively studied by various biochemical and physiological approaches. It is functionally linked to MgCh; however, in vivo characterization of Gun4 mutants revealed complex changes in tetrapyrrole metabolism far beyond
what is expected from the altered MgCh activity. These results imply that Gun4 can regulate tetrapyrrole biosynthesis in multiple ways.

Gun4 binds tightly to the ChlH subunit of MgCh, forming a membrane-localized complex, which is expected to be the site of Mg-PIX (MgP) synthesis. However, a fraction of Gun4 is localized in the cytosol/stroma. Here, we show that in the cyanobacterium *Synechocystis* sp. PCC 6803 (hereafter *Synechocystis*), the soluble N-acetylornithine aminotransferase (ArgD) and cyanophycinase (CphB) enzymes interact with Gun4. CphB is a peptidase that breaks down the cyanophycin biopolymer into β-Asp-Arg dipeptides. It is categorically found in cyanophycin-containing bacteria and is apparently dispensable in *Synechocystis*. On the other hand, ArgD is an essential enzyme in Arg biosynthesis from bacteria to plants. Our study shows that ArgD binds Gun4 with high affinity and that the accumulation of the Gun4-ArgD complex is modulated by the cellular level of ornithine (Orn), a central intermediate of Arg biosynthesis. High levels of Orn inhibit Chl biosynthesis via an ArgD-dependent mechanism, supporting the role of the Gun4-ArgD complex in a cross-talk between the Arg and tetrapyrrole pathways.

**RESULTS**

**The Gun4 protein interacts with enzymes involved in Arg metabolism**

To better understand the metabolism of N in *Synechocystis*, we focused on the Arg metabolic pathway playing a central role in N homeostasis. Specifically, we used anti-FLAG pull-downs to identify protein interactors of enzymes involved in Arg biosynthesis. We constructed a strain expressing 3×FLAG-tagged ArgD (f.ArgD) from the constitutive *psbAII* promoter while lacking the native enzyme (*f.argD/DargD strain; see Table S1*). Unlike the *DargD* mutant, which is an Orn auxotroph, the *f.argD/DargD* mutant proliferated without Orn supplement, showing growth and pigmentation comparable with the wild-type (WT) strain (Figure 1A).

The f.ArgD enzyme was isolated from the soluble fraction according to Koskela et al., and the obtained eluate was separated on a 2D blue native (BN)/SDS-PAGE together with a WT control. The resulting gel was stained, and the visible protein spots were identified by mass spectrometry (MS) (Figure 1B; Table S2). Tryptophanyl t-RNA synthetase (TrpS) and fructose-1,6-biphosphatase (Fbp-1) were evident contaminants, eluted from the FLAG resin also when using the WT control. On the other hand, despite their detachment from f.ArgD during the separation by BN, Slr0983 (hypothetical glucose-1-phosphate cytidylyltransferase homolog), CphB, and Gun4 were identified as specific co-eluates of f.ArgD (Figure 1B). CphB (29.4 kDa) migrated like a large oligomer (>200 kDa), which was more intensely stained than the spot of the apparently monomeric Gun4 (26.6 kDa). The mass of f.ArgD (29.4 kDa) on the BN gel appeared at about 100 kDa, indicating a dimer consistent with other studies. Given the well-established and critical role of Gun4 in Chl biosynthesis, we focused on the interaction of Gun4 with enzymes involved in Arg metabolism (ArgD and CphB).
To assess the *in vitro* affinities of *Synechocystis* ArgD to CphB and Gun4, we expressed these proteins in *Escherichia coli* (*E. coli*) as 6×His-tagged variants. After purification on a nickel column, recombinant proteins were subjected to isothermal spectral shift assays, in which a standard concentration of labeled ArgD was titrated with CphB or with Gun4. Fitting of the resultant binding isotherms revealed $K_d$ values of $29.8 \pm 0.96 \mu M$ and $55.8 \pm 11.5 \mu M$ for the titration of ArgD with CphB and Gun4, respectively (*Figure 1C*). These results indicate that even though CphB and ArgD both participate in Arg metabolism (see *Figure 1D*), the *in vitro* binding affinity of ArgD to Gun4 is much stronger than to CphB.

Although Gun4 was recognized in a membrane-localized complex, herein, it was shown to interact with the strictly soluble ArgD protein. To confirm the presence of Gun4 in a cytosolic enzyme complex, the co-purification assay was repeated using a cytosolic extract of a *Synechocystis* strain expressing 3× FLAG-tagged Gun4 (f.Gun4). *In vivo* activity of Gun4 has been shown previously to be unaffected by the addition of FLAG tag. The obtained f.Gun4 and control WT eluates were separated by SDS-PAGE and stained, and the most intense protein bands were analyzed by proteomic MS. Pyruvate kinase 1 (Pyk-1), ArgD, and CphB were the most abundant, specific co-eluates of f.Gun4 (*Figure 2A; Table S3*). The interaction of Gun4 with CphB was analyzed by *spectral shift* assays, revealing a $K_d$ of 2.16 ± 0.64 μM (*Figure 2B*). This value indicates a ~40 times weaker binding affinity of Gun4 to CphB compared with ArgD (*Figure 1C*).

ArgD strongly binds Gun4 *in vitro* (*Figure 1C*). Nevertheless, the relatively high level of CphB in the f.ArgD and f.Gun4 pull-downs (*Figures 1B, 2A, 2C, and 2D*) raises the question of whether CphB is important for the Gun4-ArgD interaction *in vivo*. To address this, we purified f.ArgD from the cphB-deletion background (∆cphB) that shows no phenotypic changes during phototrophic growth (see also *Figure 1A*). Since co-purification of Gun4 with f.ArgD was apparently not affected by the absence of CphB (*Figures 2C and 2D*), we concluded that CphB is not required for the accumulation of the Gun4-ArgD complex *in vivo*.

While Gun4 is specific for oxygenic phototrophs, ArgD is a housekeeping enzyme in virtually any organism. To investigate the role of the Gun4-ArgD complex in *Synechocystis*, we replaced the ArgD enzyme with its FLAG-tagged homolog (AstC) from *E. coli*. The resulting strain (f.astC∗/∆argD) expressed the f.AstC enzyme to a level comparable to f.ArgD (*Figure S1A*) while showing no alteration in the accumulation of amino acids including Orn and Arg (*Figures S1B and S1C*). The purified f.AstC, however, did not co-elute with either Gun4 or CphB (*Figures 2C and 2D*), implying that the *E. coli* enzyme does not bind these proteins *in vivo*. The f.astC∗/∆argD strain showed phototrophic growth similar to f.argD∗/∆argD (*Figure 1A*) and exhibited no apparent phenotype under the various physiological conditions tested (*Figures S2A–S2E*).

**Orn triggers the formation of the Gun4-ArgD complex and reduces the steady states of Chl precursors**

Although the interaction of Gun4 with enzymes participating in Arg metabolism was not detectable in f.astC∗/∆argD, no apparent growth defect was caused by the absence of these protein assemblages (*Figures 1A and S2A–S2E*). However, it is possible that Gun4-ArgD is important for regulation during severe or prolonged shifts in Arg metabolism, and such conditions are difficult to mimic in laboratory. As an alternative, we monitored the phenotype of studied strains after feeding with biosynthetic intermediates of Arg. Using metabolic profiling, we checked first the ability of *Synechocystis* to uptake N-acetylornithine (AcOrn), the enzymatic product of ArgD. The treatment of WT with 100 μM AcOrn for 40 min did not, however, increase the concentration of this compound in cells above the detection limit, suggesting an inefficient uptake (*Figure S3A; Table S4*). On the other hand, the same treatment with 100 μM Orn, which is the center intermediate metabolite for both Arg biosynthesis and degradation (see *Figure 1D*), elevated the concentration of intracellular Orn by three magnitudes. Orn also caused severe changes in the accumulation of other metabolites, including an increase of AcOrn (*Figure S3A; Table S4*). In line with previous studies, Orn effectively redirected metabolic fluxes in the Arg pathway and likely blocked the initial steps of Arg synthesis.

Given the well-established role of Gun4 in Chl biosynthesis, we then compared the levels of biosynthetic intermediates of Chl in cells grown with NaNO3 supplement with or without 1 mM Orn. We employed as a negative control the f.astC∗/+argD strain, in which the ArgD-Gun4 complex was undetectable (*Figures 2C and 2D*). While Orn apparently did not affect the pool of Chl precursors in f.astC∗/+argD, the f.argD∗/+argD and WT strains accumulated significantly (p < 0.05) lower relative amounts of the monitored tetrapyrroles (*Figures 3A and S3B*). Further cultivation in the Orn-containing BG-11 media resulted in decreased amount of bilins and Chl (*Figure S3C*). Consequently, all the strains used in the study had impaired photoautotrophy in the presence of Orn except for those that did not contain the ArgD enzyme (ΔargD, f.astC∗/+argD; *Figure 3B*). Orn was converted also to Arg (*Figure S3A*); however, the direct addition of Arg had a milder or no effect on the photoautotrophic growth (*Figure S3D*).

To test whether the lower accumulation of Chl precursors in the Orn-fed f.argD∗/+argD cells can be related to Gun4-ArgD, we compared the *in vivo* abundance of the complex in cells that were fed with nitrate supplement only or also with Orn before FLAG-affinity purification. Analysis of the obtained f.ArgD pull-downs revealed six times more Gun4 co-purified with f.ArgD from cells pre-treated with 1 mM Orn for 24 h (*Figure 3C*). It is notable that CphB was also more enriched in the eluate after Orn feeding. In our standard purification protocol, FLAG-tagged proteins are released from the resin under native conditions using FLAG-tag peptide. To exclude the possibility that a fraction of the specifically bound proteins remained attached to the resin after elution, we performed another preparation in which proteins were released from the FLAG resin using SDS instead of FLAG peptide. Using this approach and subsequent immunodetection, we detected an even stronger Orn-stimulated enrichment of Gun4 (>12×) in the pull-down (*Figure 3D*). In this experiment, we monitored the effect of short-term (1 h) feeding with 100 μM Orn that also substantially increased the amount of f.ArgD-co-purified Gun4 (~6 times; *Figure 3D*).

To see the extent of Gun4 sequestered from the membranes to the soluble ArgD-Gun4 complex, we performed western blot...
analysis of the soluble (cytoplasm [C]) and insoluble (membrane [M]) fractions of cells with or without Orn treatment. In fact, a higher portion of Gun4 could be detected from the soluble fraction of the Orn-treated compared with the untreated f.argD+/−DargD and WT but not of the f.astC+/DargD cells (Figure 3E).

On the other hand, Orn had the opposite effect on the relative distribution of CphB, as it could be detected from the insoluble fraction dependent on Orn treatment.

Orn inhibits de novo biosynthesis of tetrapyroles and postpones recovery from nitrogen deprivation

To clarify whether the lower accumulation of Chl precursors in the presence of additional Orn was due to inhibition of de novo tetrapyrole synthesis, we tested the effect of Orn on regreening cells, in which de novo pigment synthesis is cardinal.37 Bleaching of the cultures was achieved by 20 h of N starvation, and the recovery was induced by the addition of 1 mM NaNO3 alternatively combined with 100 μM Orn. When Orn was omitted from the N supply of the N-starved cultures, Chl precursors were synthesized within 40 min, and after an hour of N upshift, their relative amounts substantially increased in all strains studied (Figures 4A and S4A). However, unlike in the f.astC+/DargD strains, in which Orn did not affect tetrapyrole synthesis, in the control WT as well as in the f.argD+/DargD strains, the accumulation of Chl precursors (including PIX) was arrested for at least 3 h (Figures 4A and S4A). Consequently, in the Orn-fed WT and f.argD+/DargD, the overall accumulation of Chl was inhibited (Figures 4B and S4B), and in the first 14 h of N upshift, the cell absorption remained comparable to that of the N-starved cultures (Figures 4C and S4C). On the other hand, 14 h of N upshift increased the tetrapyrole pigment-binding complexes in f.astC+/DargD regardless of the presence of Orn (Figure 4C), while in the Orn-fed WT as well as f.argD+/DargD, the cell absorption peaks at 625 and 682 nm started to increase only after 30 h (Figures 4C and S4C), indicating accumulation of the phycobilin-(625 nm) and Chl-binding (682 nm) photosynthetic proteins that eventually allowed photoautotrophic growth (Figures 4D and S4E). The recovery of the N-starved, bleached WT cultures in the presence of Orn was thus much slower compared with the strain possessing the E. coli AstC enzyme (Figure 4E), while it was comparable with the Orn-fed f.argD+/DargD cultures (Figure S4).

DISCUSSION

The Gun4 protein is specific and ubiquitous to oxygenic phototrophs, where it is essential for efficient Chl production. Consistent with the model of active, membrane-bound MgCh,19 Gun4 was co-isolated with ChlH from thylakoids34 or total cell extracts29 but not from the cytosolic fraction (Figure 2A). Prior to this work, the only known protein partner of Gun4 was the ChlH subunit of MgCh. Herein, we identified Synechocystis Gun4 in a soluble protein complex(es) consisting of the Arg metabolism-related ArgD and CphB, as well as the Pyk-1

Figure 2. Gun4 is co-isolated with ArgD and CphB but not with the heterologously expressed E. coli AstC

(A) SDS-PAGE separation of the f.Gun4 and control (WT) pull-downs. The indicated proteins were identified by MS (Table S3). The molar ratios of co-eluted proteins, calculated from raw band intensities normalized to the molecular weights, are shown in brackets. (B) Recombinant Gun4 and CphB proteins were subjected to a spectral shift assay, in which 20 nM labeled Gun4 was titrated with CphB. Symbols and error bars represent the average data of three independent experiments and their standard deviation; the obtained Kd value is shown. (C) Protein pull-downs prepared from indicated strains were separated on SDS-PAGE and stained. Asterisk indicates the native ArgD that was present in the ΔcphB background. The molar ratios of co-eluted proteins are also indicated (see A). (D) The SDS-PAGE gel shown in (C) was blotted to a polyvinylidene fluoride (PVDF) membrane and probed with specific antibodies against Gun4 and CphB.
enzymes (Figure 2A). Apart from Pyk-1, this protein set can be isolated by using both f.ArgD (Figures 1B and 2C) and f.Gun4 (Figure 2A) as bait. The putative interaction of Gun4 with Pyk-1 needs further confirmation and studies. Herein, we confirmed and evaluated the interactions between Gun4-ArgD and ArgD-CphB, as well as Gun4-CphB; however, it is not definite whether we isolate these separate protein-protein interactions or one ternary complex (ArgD-Gun4-CphB). The absence of CphB does not cause obvious phenotypic changes under the photoautotrophic conditions used in our and in previous studies.27 However, it was previously noted that the ΔcphB strain accumulates slightly more bilin-containing light-harvesting complexes.27 The herein identified interaction of CphB with Gun4 implies that CphB can directly modulate the distribution of Chl and bilin pigments via its interaction with Gun4.

The interaction of ArgD with CphB is likely connected to the regulation of the Arg levels in the cell since both these enzymes contribute to the accumulation of Arg. Additional Orn intensified the interaction of these soluble enzymes; however, at the same time, approximately half of the total amount of CphB could be isolated from the insoluble fraction of the cells. These results are difficult to interpret since the physiological importance of CphB in Synechocystis remains enigmatic.

On the other hand, the importance of ArgD as an AcOrn aminotransferase in Arg biosynthesis has been established.38 Although it was proposed to function also as Orn38 or γ-aminobutyric acid39 aminotransferases, these activities are negligible compared to its AcOrn aminotransferase activity.32 These results imply that ArgD-Gun4 participates in an Arg biosynthesis-dependent regulation of the tetrapyrrole pathway. ArgD bound Gun4 with a Kd of 55.8 ± 11.5 nM (Figure 1C), which is close to the binding constant obtained by a fluorescence quenching measurement for the Gun4-ChlH complex (Kd ≈ 10 nM).40 These data indicate that the binding affinities of Gun4 to ArgD and ChlH are comparable. We demonstrated that additional Orn, which can also boost the interaction of f.ArgD with Gun4 and CphB (Figures 3C and 3D), halted de novo Chl production in the presence, but not in the absence, of Synechocystis ArgD (Figure 4A). These results imply an Orn-dependent downregulation of tetrapyrrole biosynthesis that is likely related to the formation of the ArgD-Gun4 complex. ChlH was absent in the herein isolated soluble complexes of Gun4 (Figures 1B, 2A, and 2C). The
but also the PIX substrate of MgCh were affected (Figures 3A). Is about half that of ChlH (Figures 3A, 4A, and S4A), feedback downregulation of the entire tetrapterrole biosynthetic pathway has been repeatedly observed for MgCh and Gun4 mutants in plants. Also, a number of studies suggest that Gun4 is not merely a protein factor enhancing the activity of MgCh but that it can also directly or indirectly modulate the synthesis of 5-ALA. ArgD is an essential enzyme for the biosynthesis of Orn that is conserved from bacteria to plants. In consequence, an ArgD homolog from E. coli (AstC) functionally replaced the Synechocystis ArgD in the biosynthesis of Arg (Figures 1A and S1B), but its in vivo interaction with Gun4 and CphB was not detectable (Figures 2C and 2D). However, the absence of the Gun4-ArgD interaction did not affect the viability of cells even under severe stress conditions such as fluctuating and/or high light intensities combined with cold stress (Figure S2A), unstable availability of N supply (Figure S2B), or recovery from dormancy (Figures S2C–S2E). These results imply that the presence of this complex is not critical under the tested conditions. Alternatively, Gun4-ArgD might function in the fine regulation of tetrapyrrole biosynthesis, which is difficult to track down unless the intracellular levels of Arg metabolites are intensely altered by Orn supplement (Figure S3A; Table S4). Also, our recent data suggest that there are other sites of cross-talk besides Gun4-ArgD, where the tetrapyrrole and Arg pathways are intertwined in Synechocystis (Sobotka, R., unpublished data). Using a similar approach as described here, we discovered a stable protein complex between the bifunctional ArgJ, which catalyzes the first and fifth steps of the Arg biosynthesis (see Figure 1D), and GluTR, a rate-limiting enzyme of 5-ALA formation. Multiple regulatory checkpoints (protein complexes) between these two metabolic pathways can indeed mitigate the absence of a single point, which could explain why the fastC'ΔargD strain showed no defects in viability under the stress conditions tested (Figure S2). Nevertheless, the co-existence of multiple protein-protein interactions further supports the
importance of the co-regulation of Arg metabolism with the biosynthesis of tetrapyrroles.

Limitations of the study
Our working model postulates that at elevated Orn levels, Gun4 preferentially associates with ArgD in the cytoplasm and that its availability for membrane-localized Chl synthesis becomes limited. We observed Gun4 relocalization in Orn-treated cells using immunodetection from fractionated cells (Figure 3E). However, the weakly membrane-bound Gun4 could partially dissociate during the preparation of cellular fractions. A better approach would be to monitor the Orn-driven relocalization of Gun4 in intact cells. However, the fluorescent-tagged Gun4 was unstable and prone to degradation. The current study also lacks the identification of a physiological condition in which the discovered ArgD-Gun4 complex plays an important role. This is most likely due to the gap in our knowledge on the eco-physiology of our model strain, which was isolated 52 years ago. Since then, Synechocystis has been widely cultivated under optimal laboratory conditions in growth media containing waste amount of nutrients, including N, which particularly underlines studies related to N homeostasis. Consequently, there are no information available on Orn signaling or on the physiological significance of the CphB enzyme in our organism. It conceivably deflects the interpretation of the herein described effect of Orn on the regulation of tetrapyrrole synthesis. Similarly, without knowing the physiological relevance of CphB, it is too early to make conclusion about the importance of its interactions with other proteins in a metabolic network. Therefore, the current report inspires further studies in these directions.

STAR METHODS
Detailed methods are provided in the online version of this paper and include the following:

- KEY RESOURCES TABLE
- RESOURCE AVAILABILITY
  - Lead contact
  - Materials availability
  - Data and code availability
- EXPERIMENTAL MODEL AND SUBJECT DETAILS
- METHOD DETAILS
  - Construction of the Synechocystis model strains
  - Isolation of Synechocystis soluble and membrane proteins
  - Isolation of protein complexes from Synechocystis
  - Protein electrophoresis, immunoblotting and mass spectrometry
  - Quantification of selected metabolites
  - Purification of recombinant Gun4, CphB and ArgD proteins from Escherichia coli
  - Spectral shift assays
- QUANTIFICATION AND STATISTICAL ANALYSIS

SUPPLEMENTAL INFORMATION
Supplemental information can be found online at https://doi.org/10.1016/j.celrep.2023.113265.


## STAR METHODS

### KEY RESOURCES TABLE

<table>
<thead>
<tr>
<th>REAGENT or RESOURCE</th>
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<td><strong>Antibodies</strong></td>
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<tr>
<td>anti-FLAG</td>
<td>Sigma-Aldrich</td>
<td>Cat# F7425, RRID:AB_439687</td>
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</table>
| anti-Gun4           | Raised in rabbit against the recombinant *Synechocystis* Gun4 (Sobotka et al.
|                     | (this study). | N/A |
| anti-CphB           | Raised in rabbit against the recombinant *Synechocystis* CphB (this study). | N/A |
| anti-ArgD           | Raised in rabbit against the recombinant *Synechocystis* ArgD (this study). | N/A |
| anti-rabbit IgG-peroxidase antibody produced in goat | Sigma-Aldrich | Cat# A6154, RRID:AB_11125345 |
| goat-anti-rabbit IgG IR800 | Azure Biosystems | Cat# AC2134 |
| **Bacterial strains** |        |            |
| WT-P substrain of *Synechocystis* sp. PCC 6803 (used as a background for all other strains of this study) | Tichy et al.
| ΔargD *Synechocystis* sp. PCC 6803 | This study | N/A |
| f.argD+/ΔargD *Synechocystis* sp. PCC 6803 | This study | N/A |
| f.astC+/ΔastC *Synechocystis* sp. PCC 6803 | This study | N/A |
| ΔcphB *Synechocystis* sp. PCC 6803 | This study | N/A |
| f.argD+/ΔcphB *Synechocystis* sp. PCC 6803 | This study | N/A |
| f.gun4+/- Δgun4 *Synechocystis* sp. PCC 6803 | Sobotka et al.
| pET21a-gun4 *Escherichia coli* | This study | N/A |
| pET21a-argD *Escherichia coli* | This study | N/A |
| pET21a-cphB *Escherichia coli* | This study | N/A |
| **Chemicals, peptides, and recombinant proteins** |        |            |
| anti-FLAG M2 affinity gel | Sigma-Aldrich | Cat# A2220 |
| 3xFLAG peptide | Sigma-Aldrich | Cat# A4799 |
| Gun4-6xHis | This study | N/A |
| ArgD-6xHis | This study | N/A |
| CphB-6xHis | This study | N/A |
| **Deposited data** |        |            |
| Proteomics data (identification of protein gel bands) | MassIVE: https://massive.ucsd.edu/ProteoSAFe/static/massive.jsp | MassIVE: MSV000091280 |
| Metabolomics data | figshare: https://doi.org/10.6084/m9.figshare.24205368.v1 |
| Oligonucleotides |        |            |
| For list of primers see Supplementary Table S5 | This study | N/A |
| **Recombinant DNA** |        |            |
| pPD-NFLAG (Km) | Chidgey et al.
| | Cat# A2220 |
| **Software and algorithms** |        |            |
| Thermo Xcalibur | Thermo Fisher Scientific | RRID:SCR_014593 |
| MassHunter Quantitative Analysis software | Agilent | RRID:SCR_015040 |
| ImageJ | Schneider et al.
| | Cat# AC2134 |

## RESOURCE AVAILABILITY

### Lead contact

Further information and requests for resources and reagents should be directed to and will be fulfilled by the lead contact, Roman Sobotka (sobotka@alga.cz).
Materials availability
The E. coli and Synechocystis strains generated in this study will be sent after request from the lead contact.

Data and code availability
- Proteomics data have been deposited at MassIVE: MSV000091280 and are publicly available as of the date of publication. Accession numbers are listed in the key resources table. Metabolomics data have been deposited at figshare (https://doi.org/10.6084/m9.figshare.24205368.v1) and are publicly available as of the date of publication.
- Further data reported in this paper, and any additional information required to reanalyze the data reported in this paper will be shared by the lead contact upon request.
- This paper does not report original code.

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Synechocystis sp. PCC 6803 strain GT-P[48] was used as the wild type (WT) and as a genetic background for all prepared strains listed in Table S1. Unless stated otherwise, the Synechocystis strains were grown photoautotrophically in liquid BG-11 medium on a rotary shaker at 28°C, under continuous, moderate irradiance of 40 μmol photons m⁻² s⁻¹ given by white fluorescence tubes. The plate-drop experiments were performed by pipetting liquid cultures on a BG-11 agar plate. The drops were photographed after three days of photoautotrophic growth at 30°C, under constant illumination with 30 μmol photons m⁻² s⁻¹. Alternatively, the plates were exposed to 500 μmol photons m⁻² s⁻¹ (high light) at 28 or 22°C; or to repeated periods of 5 min illumination with high light and 5 min dark (fluctuating light). The fluctuating nitrogen (N) stress was mimicked by repeated cycles of washing the cells to nitrate-less BG-11 (N₂) for 8 h; followed by the addition of 18 mM NaNO₃ supplement to the liquid cultures for another 8 h; while agitated with 240 rpm at 28°C; illuminated by 200 μmol m⁻² s⁻¹. Dormancy and recovery were induced by 25 d of N-starvation[51] and the addition of 1 mM NaNO₃, respectively. When amino acids were included to the media the stable pH was ensured by 10 mM TES. The absorption spectra of cells were measured by a UV-3000 spectrophotometer (Shimadzu). The number and average size of cells were assessed by coulter counter (Multisizer 4, Beckman Coulter).

METHOD DETAILS

Construction of the Synechocystis model strains
The ΔargD and ΔcphB strains were constructed by replacing the slr1022 and slr2001 genes by erythromycin or spectinomycin resistance cassettes, respectively. The primers in the megaprimer PCR method used for mutagenesis are listed in Table S5. The segregation of ΔargD locus was achieved in the presence of 5 mM Orn. For the construction of strains expressing the N-terminally 3xFLAG-tagged ArgD protein (f.ArgD) we purchased a synthetic Synechocystis argD gene (GenScript, USA) with an optimized codon usage to remove the common restriction binding sites. To express the AstC enzyme from Escherichia coli (E. coli) in Synechocystis, the astC gene of E. coli was amplified by PCR from the genomic DNA. The argD and astC genes were cloned to the pPD-NFLAG plasmid[52,49] (for all primers used see Table S5). The obtained constructs were transformed into Synechocystis ΔargD and/or ΔcphB cells. All transformed Synechocystis cells were fully segregated on BG-11 plates with increasing concentrations of relevant antibiotics (see Table S1).

Isolation of Synechocystis soluble and membrane proteins
Synechocystis cells (5 x 10⁸) from exponential growth phase were pelleted, washed, and re-suspended in buffer A containing 20 mM HEPES pH 7.4, 5 mM CaCl₂, 10 mM MgCl₂, 25% glycerol, protease inhibitor (cOmplete; Roche). The cells were broken mechanically in a Precellys Evolution tissue homogenizer (Bertin Instruments) using balotina beads (100–200 μm). The breaking was performed in three cycles of shaking at 7500 rpm at 0°C; the sample was chilled for 2 min between the cycles. After removing the beads by repeated washing and sedimentation steps, the insoluble and soluble fractions were separated by three series of centrifugation at high speed (4°C, 65000 x g, 20 min).

Isolation of protein complexes from Synechocystis
Cells from 4 L of Synechocystis cultures (c. 2x10⁶ mL⁻¹ cell content) were centrifuged, washed with buffer A and broken as described in the ‘Isolation of Synechocystis soluble proteins’ section. The soluble fraction was applied to anti-FLAG M2 affinity gel chromatography (Sigma-Aldrich) essentially as described in[31]. Proteins bound to the column were washed with 15 column volumes of buffer A. FLAG-tagged protein complexes were eluted with synthetic 3xFLAG peptide (150 μg/mL) in buffer A. Alternatively, the proteins were eluted form the column with 1% SDS in buffer A.

Protein electrophoresis, immunoblotting and mass spectrometry
The isolated proteins and protein complexes were solubilized and separated on SDS-PAGE essentially as described in[53], and subsequently stained by Coomassie Blue. In case of immunoblot, the gel was stained by SYPRO Orange (Sigma-Aldrich) and
transferred onto a PVDF membrane that was subsequently incubated with specific primary antibody and then with secondary antibody for horseradish peroxidase (Sigma-Aldrich). The following primary antibodies were used in the study: anti-Gun4, anti-D154, anti-FLAG (Sigma-Aldrich Cat# F7425, RRID:AB_439867); the polyclonal anti-CphB and anti-ArgD antibodies were generated in rabbit, against the full-length Synechocystis recombinant proteins produced in E. coli (Moravia Biotech). The primary antibodies were probed with anti-rabbit IgG-peroxidase antibody produced in goat (Sigma-Aldrich Cat# A6154, RRID:AB_11125345) and visualized using Immobilon Crescendo Western HRP substrate (Millipore, Cat# WBLUR0500, RRID:AB_439867) and luminescence image analyzer (ImageQuant, LAS-4000). Alternatively, we used goat-anti-rabbit IgG IR800 antibody (Cat# AC2134) and visualize the signal using Azure Biosystems. For the identification of proteins by liquid chromatography (LC) coupled tandem mass spectrometry (MS/MS), the Coomassie Blue-stained bands were cut from the gel, digested and analyzed as described here.55 For protein identification, MS/MS spectra were searched against Synechocystis species-specific protein bases (UNIPROT Universal Protein Resource, RRID:SCR_002380) and Cyanobase (RRID:SCR_007615) using the PLGS3.0 (Waters) software package.

**Quantification of selected metabolites**

The biosynthetic precursors of Chl/heme were extracted from equal number of cells and quantified by HPLC, essentially as described in.56 The Chl content per cell was determined for three independent cultures after methanol extraction of pigments according to.57 For the quantification of selected metabolites by combined GC-MS and LC-MS analysis58 3x10^9 cells were collected to the evaporated sample extract: (1) 50 μL methanol:ACN:H2O (2:2:1 v/v/v) containing an internal standard 4-fluorophenylalanine (80 nmol). The sample was then homogenized using a Tissue Lyser II (Qiagen) at 50 Hz, 0°C for 5 min. The mixture was then centrifuged at (8000 g, 10 min, 5°C). The supernatant was removed, and the extraction step was repeated under the same conditions but without the internal standard. The supernatants were combined and the obtained sample extract stored (–80°C). A 100 μL aliquot of each extract was mixed with 13C-labeled internal standards (13C3-Serine, 13C3-Alanine, 13C6-Tyrosine, 13C6-Glutamic acid, 13C6-Arginine, 13C6-Lysine, 13C6-Phenylalanine, 13C6-Asparagine, 13C6-Methionine, 13C2-Threonine, 13C2-2-Oxoglutarate, 13C6-Glutamine, 13C6-Proline, 13C6-Ariginine) (absolute 2 nmol each) were concentrated in a vacuum concentrator (RVC 2–25 CD Plus combined with ALPHA 1–2 LD Plus, Thermo-Fischer Scientific). Each dried extract sample was then subjected to derivatization with ethanol (EtOH) - ethyl chloroformate (ECF) reaction medium under pyridine catalysis and simultaneous liquid-liquid microextraction into a lower chloroform layer as described earlier.55 Briefly, the following five sequential steps were used for the addition of a corresponding medium to the evaporated sample extract: (1) 50 μL of the mixture EtOH: water (2:1 v/v/v), (2) 50 μL of the mixture EtOH: pyridine (2:1 v/v), (3) 50 μL of the mixture ECF: chloroform (1:7 v/v), (4) 50 μL 1 M NaOH, (5) 50 μL of the mixture ECF: chloroform (1:7 v/v). The reaction mixture was stirred before addition of the particular medium. Finally, 30 μL of the lower chloroform layer was evaporated by a gentle stream of nitrogen and redissolved in 100 μL methanol: water (3:7 v/v) for LC-MS analysis. 50 μL of 1 M HCl was added to the remaining reaction mixture and stirred. The 50 μL of the lower chloroform phase was used for GC-MS analysis. An LTQ XL mass spectrometer coupled to a Accela 600 liquid chromatograph (LC) and a Accela autosampler (all Thermo Fisher Scientific) was used for quantitative analysis. Amino acids were separated on a 150 mm x 3 mm i.d., 2.6 μm, Kinetex C18 (Phenomenex) with a mobile phase flow rate of 400 μL/min, an injection volume of 5 μL, and a column temperature of 35°C. The mobile phase was A = 5 mmol/L ammonium format in methanol, B = 5 mmol/L aqueous ammonium format; gradient (A): 0.0 min, 30%; 10.0 min, 100%; 11.0 min, 30%; 14.5 min, 3%. Full scan positive ion mass spectra were acquired in a mass range of 85–850 Da. LT-Q settings were as follows: 2.5 kV spray voltage ion source parameter, 300°C capillary temperature, sheath gas at 40 au, aux gas at 10 au, spare gas at 1 au, 300°C source temperature. Data were processed using Thermo Xcalibur software (Thermo Fisher Scientific, RRID:SCR_014593), version 4.0. GC-MS analyses of amino acids were performed using a VF-17ms capillary column (30 m, 250 mm, 250 μm) and a gas chromatograph 5977B coupled to a quadrupole mass spectrometer 5977B MSD (Agilent) equipped with an electron ionization source (EI) and operated in full-scan mode (40–500 Da). The instrument settings were: Helium flow rate, 1.2 mL/min; inlet temperature, 280°C; injection mode, splitless; split flow, 40 mL/min; splitless time, 1.0 min; septum purge flow 3 mL/min; temperature program, 45°C, hold for 2 min, 16°C/min to 320°C; 2 min; transfer line temperature, 280°C; and EI source temperature, 230°C; ionization energy, 70 eV. Data were processed using Agilent MassHunter WorkStation - Qualitative Analysis for GC/MS (Agilent, RRID:SCR_016657) and Agilent Masshunter Quantitative Analysis software (Agilent, RRID:SCR_015040) version 10.

**Purification of recombinant Gun4, CphB and ArgD proteins from Escherichia coli**

C-terminal His6-tagged Gun4, CphB and ArgD proteins were over-expressed in E. coli BL21 (DE3) using a pET21a plasmid (Novagen). The expression was induced with isopropyl-β-D-thiogalactopyranoside (0.4 mM) and shaken for additional 20 h at 18°C. Cells were harvested by centrifugation (10 min, 4°C, 10000 × g), resuspended in lysis buffer (25 mM Tris/HCl pH 7.8, 150 mM NaCl, 10 mg L⁻¹ of DNase I, 10 mg L⁻¹ of lysozyme, protease inhibitor (Complete; Roche), and incubated for 30 min at 37°C. After incubation, cells were disrupted by sonication and the lysate was clarified by centrifugation (4°C, 30 min, 10000 × g). Soluble His-tagged proteins were purified by metal-affinity chromatography (Protino Ni-NTA agarose, Macherey-Nagel; buffer: 25 mM Tris/HCl pH 8.0, 150 mM NaCl); their purity and concentration were checked by SDS-PAGE.
Spectral shift assays
Recombinant Gun4 and ArgD proteins were labeled with 2nd Generation Red NHS dye using standard protocols (NanoTemper Technologies GmbH, Munich, Germany) and used for isothermal spectral shift assays.\textsuperscript{59} 5 or 20 nM labeled ArgD or Gun4 was titrated with a serial dilution of CphB or Gun4, respectively. Binding experiment were performed in triplicate, in 25 mM Tris, 0.1% Pluronic F-127, pH 8 at 25°C. Samples were loaded into Monolith Premium capillaries (NanoTemper Technologies) and loaded into a Monolith X instrument (NanoTemper Technologies) and excited at 590 nm (100% LED power), using software version Mo.Control 2.4.1. The ratio of fluorescence emission at 670 : 650 nm was collected; and the data were exported to CSV files, where custom Python scripts were used to determine the dissociation constant ($K_d$) using the equations in.\textsuperscript{59}

QUANTIFICATION AND STATISTICAL ANALYSIS

The ratios of the co-purified f.ArgD and Gun4 proteins indicated on Figures 3B and 3C were assessed from the intensity of the stained protein bands or antibody signals using the ImageJ software (RRID:SCR_003070).\textsuperscript{50} The average (arithmetic mean) and standard deviation of the data (shown on Figures 3A and 4A, 4B and 4D; S4B and S4E) were determined from measurements of $n = 3$ samples. Significance of the data (where indicated) was tested with one-tailed $t$ test, with a significance level set to $p < 0.05$. 