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| 著者 Author(s) | Hasunuma, Tomohisa / Matsuda, Mami / Senga, Youhei / Aikawa, Shimpei / Toyoshima, Masakazu / Shimakawa, Ginga / Miyake, Chihiko / Kondo, Akihiko |

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Overexpression of flv3 improves photosynthesis in the cyanobacterium Synechocystis sp. PCC6803 by enhancement of alternative electron flow

Tomohisa Hasunuma1,2*, Mami Matsuda1,2, Youhei Senga1,2, Shimpei Aikawa3, Masakazu Toyoshima4, Ginga Shimakawa5, Chikahiro Miyake5,6 and Akihiko Kondo3,6,7

Abstract

Background: To ensure reliable sources of energy and raw materials, the utilization of sustainable biomass has considerable advantages over petroleum-based energy sources. Photosynthetic algae have attracted attention as a third-generation feedstock for biofuel production, because algae cultivation does not directly compete with agricultural resources, including the requirement for productive land and fresh water. In particular, cyanobacteria are a promising biomass feedstock because of their high photosynthetic capability.

Results: In the present study, the expression of the flv3 gene, which encodes a flavodiiron protein involved in alternative electron flow (AEF) associated with NADPH-coupled O2 photoreduction in photosystem I, was enhanced in Synechocystis sp. PCC6803. Overexpression of flv3 improved cell growth with corresponding increases in O2 evolution, intracellular ATP level, and turnover of the Calvin cycle. The combination of in vivo 13C-labeling of metabolites and metabolomic analysis confirmed that the photosynthetic carbon flow was enhanced in the flv3-overexpressing strain.

Conclusions: Overexpression of flv3 improved cell growth and glycogen production in the recombinant Synechocystis sp. PCC6803. Direct measurement of metabolic turnover provided conclusive evidence that CO2 incorporation is enhanced by the flv3 overexpression. Increase in O2 evolution and ATP accumulation indicates enhancement of the AEF. Overexpression of flv3 improves photosynthesis in the Synechocystis sp. PCC6803 by enhancement of the AEF.

Keywords: Synechocystis sp. PCC6803, flv3, Alternative electron flow, Carbon metabolism, Photosynthesis, Metabolite profiling

Background

Environmental concerns associated with the continued depletion of oil reserves and greenhouse gas emissions have prompted governments to promote research on environmentally benign and sustainable fuels towards realizing energy independence. In particular, utilization of biomass as the starting material for fuel production has recently attracted considerable attention [1,2]. It is anticipated that the use of biofuels as an alternative to fossil fuels will help reduce CO2 accumulation through reduced emissions and recycling of the CO2 that is released when biofuel is combusted as a transport fuel.

Photosynthetic algae are of increasing interest as a renewable source of biomass for the sustainable production of biofuels [3-5], because algal feedstocks do not directly compete with agriculture, as they require no productive land. Cyanobacteria are particularly attractive, because their photosynthetic and biomass production rates are higher than those of terrestrial plants [3,6]. Several cyanobacterial species have the ability to store large amounts of energy-rich glycogen that can be utilized for the production of biofuels, including bio-ethanol [7,8]. In addition, because cyanobacteria are prokaryotes, their metabolic processes are more readily amenable to genetic modifications designed to enhance photosynthetic activity than are those of eukaryotic algae.
In the phototrophic metabolism, carbon assimilation through the Calvin cycle is coupled with consumption of ATP and NADPH. Cyanobacterial cell growth rate is dependent on the intensity of light with the exception of photoinduction and photooxidative stress, as it drives photosynthetic electron transport and the production of ATP and NADPH. ATP is required not only for turnover of the Calvin cycle, but also for the CO₂ concentrating mechanism, which serves to increase the intracellular CO₂ concentration [9-11]. Furthermore, ATP is required for glycogen biosynthesis in photosynthetic cells [12], meaning that a continuous supply of ATP is necessary for a higher rate of photosynthesis in cyanobacteria.

ATP is produced by ATP synthase via a proton gradient (ΔpH) across the thylakoid membrane. ΔpH is mainly generated by the photosynthetic linear electron flow (LEF) pathway, although phototrophs, including cyanobacteria, have also developed a large arsenal of alternative electron flow (AEF) pathways, which assist in modulating the ATP/NADPH ratio in response to metabolic demand [13,14]. O₂-dependent electron flow, termed the Mehler pathway or water-water cycle (WWC), is one AEF pathway that helps generate a pH gradient [15-17]. In the Mehler pathway, electrons extracted from water by photosystem II (PSII) are used to generate water by the photoreduction of O₂ coupled with the oxidation of NADPH in photosystem I (PSI).

In cyanobacteria, flavodiiron (Flv) proteins possess an NAD(P)H:flavin oxidoreductase module [18]. The cyanobacteria Synechocystis sp. PCC6803 has four genes encoding Flv proteins (Flv1, Flv2, Flv3, and Flv4). The results of an in vitro study with an flv3 mutant provided evidence that Flv3 functions as an NAD(P)H:oxygen oxidoreductase [18]. A subsequent in vivo study with Δflv1 and Δflv3 mutants of Synechocystis sp. PCC6803 confirmed that Flv1 and Flv3 are involved in the photoreduction of O₂ to H₂O in the Mehler reaction [19]. Under fluctuating light conditions, the growth and photosynthesis of Δflv1 and Δflv3 mutants of Synechocystis sp. PCC6803 are arrested [20].

In the present study, a recombinant flv3-overexpressing Synechocystis strain (Flv3ox) was constructed to examine the effects of flv3 overexpression on the photosynthetic ability of Synechocystis sp. PCC6803. Enhancement in the AEF pathway through the regeneration of NADP⁺ improved ATP accumulation in the Flv3ox cell. Recently, we developed an analytical method to directly measure the turnover of metabolic intermediates in cyanobacteria [21]. The combination of in vivo ¹³C-labeling of metabolites and metabolome analysis using mass spectrometry (MS) enables kinetic visualization of carbon assimilation in central metabolic pathways, such as the Calvin cycle, glycolysis, and glycogen biosynthesis. Here, the dynamic metabolic profiling approach was used to reveal the underlying mechanisms of the enhancement of the photosynthetic carbon flow in the Flv3ox strain.

**Results**

**Construction of an flv3-overexpressing recombinant strain**

To enhance flv3 expression in Synechocystis sp. PCC6803, we constructed the transformation vector pTCP2031V-flv3, which contained flv3 linked to the psbA2 promoter between the shr2030 and shr2031 genes, which acted as anchoring regions for site-specific integration into the Synechocystis genome through homologous recombination (Figure 1a). A glucose-tolerant (GT) strain of Synechocystis sp. PCC6803 was transformed with pTCP2031V-flv3 to yield strain Flv3ox. The chromosomal integration of flv3 was confirmed by genomic PCR (Figure 1b). A vector control (VC) strain, in which the chloramphenicol resistance cassette was inserted into the genome of GT, was constructed with an empty vector pTCP2031V. Immunoblot analysis showed higher levels of Flv3 protein in the Flv3ox strain compared to the parental GT and vector control strains (Figure 1c).

**Cell growth and O₂ evolution by Flv3ox**

Figure 2 shows the growth of strain GT, Flv3ox, and VC cells cultivated under continuous irradiation with 120 μmol photons m⁻² s⁻¹ light intensity and 1% (v/v) CO₂ at 30°C. Under these conditions, Flv3ox showed a higher growth rate than the GT and VC strains. After a 7-day cultivation, the cell concentration of Flv3ox reached 3.20 g dry cell weight (DCW) L⁻¹ and that of GT was 2.48 g-DCW L⁻¹. According to statistical analysis, the biomass concentration of Flv3ox was significantly different from that of GT and VC (Additional file 1). VC demonstrated a similar cell growth ability as GT. Measurements of glycogen showed that Flv3ox had a higher glycogen content (0.117 g-glycogen g-DCW⁻¹) than that of GT (0.091 g-glycogen g-DCW⁻¹) (Figure 3).

The light dependence of the O₂ evolution rate by the two strains of Synechocystis cells was evaluated with an O₂ electrode system (Figure 4). Flv3ox exhibited a higher O₂ evolution rate than that of GT.

**Metabolic analysis of Synechocystis cells**

Photosynthetic electron flow generates a proton gradient across the cyanobacterial thylakoid membrane, driving the ATP synthesis necessary for carbon assimilation. Here, the effect of flv3 overexpression on intracellular carbon metabolism was investigated using a dynamic profiling technique [21] that measures the turnover of metabolic intermediates in cyanobacterial cells. The kinetic measurements were performed by the combination of an in-vivo ¹³C-labeling technique and mass isotopomer detection with capillary electrophoresis coupled to MS (CE/MS) for the efficient separation of highly polar molecules. To
assess metabolic turnover, extracted intracellular metabolites were analyzed after labeling for between 1 and 30 min (Figure 5). The $^{13}\text{C}$ fraction, which was defined as the ratio of $^{13}\text{C}$ to total carbon in each metabolite, was calculated from mass isotopomer distributions.

Ribulose-1,5-bisphosphate (RuBP) carboxylase/oxygenase (Rubisco) catalyzes the first major step of carbon fixation in the Calvin cycle to produce 3-phosphoglycerate (3PGA) from CO$_2$ and RuBP. For Flv3ox, the $^{13}\text{C}$ fraction of 3PGA reached a maximum of 92% after 20 min of labeling, whereas a maximum of 85% was observed after 30 min of labeling for GT (Figure 5). The overexpression of flv3 resulted in an increase in $^{13}\text{C}$-labeling
rate of metabolites involved in the Calvin cycle, including 3PGA, fructose-6-phosphate (F6P), and sedoheptulose-7-phosphate (S7P). Flv3ox also displayed a higher turnover rate of metabolites involved in glycolysis and glycogen biosynthesis, such as phosphoenolpyruvate (PEP), 2-phosphoglycerate (2PGA), acetyl-CoA (AcCoA), glucose-6-phosphate (G6P), glucose-1-phosphate (G1P), and ADP-glucose (ADP-Glc). These results clearly showed that the enhancement of flv3 expression accelerates the photosynthetic carbon assimilation rate. In addition, flv3 overexpression resulted in an increase in the turnover rate of citrate, while the $^{13}$C fraction of other metabolites involved in the citrate cycle, including cis-aconitate, isocitrate, and malate, was almost the same between Flv3ox and GT (Figure 6).

Table 1 shows the pool sizes of intracellular metabolites in the two strains of cyanobacterial cells. Flv3ox accumulated 0.207 μmol g-DCW$^{-1}$ ATP, while ATP level in GT was lower than the quantification limit (0.01 μmol g-DCW$^{-1}$) of the CE/MS analysis. There was no significant difference in NADPH level between Flv3ox and GT. NADP$^+$ was slightly (1.6-fold) increased by the flv3 overexpression. The levels of F6P, G6P, and S7P in Flv3ox were similar to those in GT, while an increase in 3PGA and a decrease in RuBP were caused by the flv3 overexpression. Flv3ox showed higher amounts of ADP-Glc and lower G1P than GT. 2PGA and PEP were also increased by the flv3 overexpression.

Discussion
The overexpression of flv3 improves the cell growth of Synechocystis sp. PCC6803, as observed by the increase

![Figure 4](image-url) Light response curves for O$_2$ evolution rate in GT and Flv3ox cells. When the OD$_{570}$ reached approximately 4.5, the cells were applied for the photosynthesis analysis. The values are the mean ± SD of three measurements.

![Figure 5](image-url) Time course of the $^{13}$C fraction for various metabolites in GT and Flv3ox cells cultivated under 120 μmol photons m$^{-2}$ s$^{-1}$ light intensity and 1% CO$_2$ conditions. The values are the mean ± SD of at least five different experiments.
in the carbon assimilation rate as well as O$_2$ evolution activity in Flv3ox cells. In Figure 5, an increase in the flux of photosynthetic carbon metabolism in Flv3ox was clearly demonstrated using a dynamic metabolic profiling approach developed by our group. So far, there has been no report that the improvement of photosynthetic CO$_2$ incorporation was directly observed in recombinant cyanobacterial strains. When metabolism is in a dynamic steady state in vivo, metabolites are replaced with newly synthesized compounds at a constant rate, and the total amount of each metabolite remains unchanged. Thus, in order to understand metabolic flux, the turnover of metabolic intermediates should be examined directly. The measurement of metabolic turnover using this in vivo $^{13}$C-labeling assay has provided conclusive evidence that CO$_2$ incorporation is enhanced by $flv3$ overexpression in Synechocystis cells.

In the photosynthetic electron transport chain, electrons extracted from H$_2$O in PSII are transferred to NADP$^+$ to produce NADPH in PSI (Figure 7). Figure 4 demonstrated that the photosynthetic O$_2$ evolution was improved by the enhancement of $flv3$ expression. Because Flv3 regenerates NADP$^+$ with the photoreduction

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Table 1 Metabolite pools in GT and Flv3ox cells cultivated under 120 μmol photons m$^{-2}$ s$^{-1}$ light intensity and 1% CO$_2$ conditions

| Metabolite | Concentration (μmol g-DCW$^{-1}$) | GT          | Flv3ox       |
|-----------|----------------------------------|-------------|--------------|
| 3PGA      | 2.267 ± 0.061                    | 3.157 ± 0.191$^{**}$ |
| F6P       | 0.071 ± 0.007                    | 0.069 ± 0.011 |
| S7P       | 0.096 ± 0.003                    | 0.094 ± 0.006 |
| RuBP      | 0.089 ± 0.006                    | 0.026 ± 0.005$^{**}$ |
| G6P       | 0.116 ± 0.015                    | 0.138 ± 0.023 |
| G1P       | 0.101 ± 0.016                    | 0.020 ± 0.003$^{**}$ |
| ADP-Glc   | 0.080 ± 0.010                    | 0.115 ± 0.004$^{**}$ |
| PEP       | 1.134 ± 0.039                    | 1.562 ± 0.100$^{**}$ |
| 2PGA      | 0.282 ± 0.008                    | 0.401 ± 0.023$^{**}$ |
| AcCoA     | 0.075 ± 0.006                    | 0.075 ± 0.015 |
| ATP       | < 0.01                           | 0.207 ± 0.021 |
| NADPH     | 0.001 ± 0.000                    | 0.003 ± 0.002 |
| NADP$^+$  | 0.495 ± 0.012                    | 0.549 ± 0.021$^{*}$ |

Values are the mean ± SEM of nine different measurements. Statistical significance was determined using the Student’s t-test ($^*$P < 0.05, $^{**}$P < 0.01).
of O\textsubscript{2}, overexpression of \textit{flv3} in \textit{Synechocystis} cells would increase photosynthetic electron transport.

In general, the energy budgets of phototrophs are dependent on the ratio of protons pumped across the thylakoid membrane to electrons passed through photosynthetic electron transfer complexes, which produce ATP and NADPH used in cellular biochemical pathways [22]. CO\textsubscript{2} fixation through the Calvin cycle is the main ATP and NADPH sink in autotrophic metabolism, requiring an ATP/NADPH ratio of 1.5, whereas the ratio generated by the LEF pathway is 1.28 [13]. AEF pathways contribute to the balancing of ATP/NADPH by supplying additional ATP [14]. For example, the Mehler reaction generates a proton gradient across the thylakoid membranes, leading to ATP generation, but the net production of reduced NADPH is not expected [16]. Although the physiological function of the Mehler reaction in cyanobacteria is not fully understood, here, increased expression of the \textit{flv3} gene in \textit{Synechocystis} promoted NADPH\textsuperscript{+} regeneration coupled with O\textsubscript{2} photoreduction in the Mehler reaction, leading to a higher photosynthetic electron transport rate and increased ATP supply, as shown in Figure 4 and Table 1. Enhancement in the intracellular ATP level led to a higher ATP/NADPH ratio in the recombinant strain. The increased ATP supply would be coupled with the cell growth and improved carbon metabolic flux through the Calvin cycle, sugar phosphate pathway, and glycogen biosynthesis.

The overexpression of \textit{flv3} increased the glycogen content in \textit{Synechocystis} cells (Figure 3). Glycogen, an α-1,6-branched α-1,4-glucan, is the primary storage polysaccharide in cyanobacteria and is biosynthesized from F6P via G6P, G1P, and ADP-Glc as intermediates [12]. As ADP-Glc is produced from G1P and ATP by G1P adenyltransferase, the enhanced supply of ATP by \textit{flv3} overexpression would lead to increased glycogen production. Increase in ADP-Glc and decrease in G1P by the \textit{flv3} overexpression, which are demonstrated in Table 1, would indicate enhanced conversion of G1P to ADP-Glc.

Flv1 and Flv3 are essential for photoreduction of O\textsubscript{2} in \textit{Synechocystis} sp. PCC6803 [19]. Allahverdiyeva et al. reported that the growth and photosynthesis of \textit{Δflv1} and \textit{Δflv3} mutants was arrested in fluctuating light, although Flv1 and Flv3 proteins were dispensable under constant-light conditions [20]. Flv1 and Flv3 were proposed to form a heterodimer [23], while Flv3 alone is sufficient for the O\textsubscript{2} photoreduction reaction \textit{in vitro} [19]. The observed difference in the phenotypes of \textit{Δflv1} and \textit{Δflv3} mutants suggests that a homodimer of Flv3 exhibits NADPH-dependent O\textsubscript{2} reduction \textit{in vivo} under excess-light stress conditions [24]. Zhang et al. reported that \textit{flv3} transcripts were markedly higher than those of \textit{flv1} in wild-type \textit{Synechocystis} under both 3% CO\textsubscript{2} and ambient CO\textsubscript{2} condition [25]. Our present results in an \textit{flv3}-overexpressing strain of \textit{Synechocystis} raise the possibility that the Flv3 protein can function alone, since overexpression of an \textit{flv3} increased the level of ATP and flux of carbon assimilation pathways.

\textit{Synechocystis} sp. PCC6803 is one of the most widely used species in studies of photosynthetic bacteria because of its unicellular morphology and fully sequenced genome [26]. In addition, the natural transformability of
this strain has allowed for the establishment of reliable genome manipulation techniques, which enables the easy modification of metabolic pathways. This has been followed by systems biology approaches, such as metabolomics, dynamic metabolic profiling, and flux balance analyses, for gaining insight into cellular metabolic processes [5,27]. A systems biology approach is needed to exploit cyanobacteria as cell factories for the production of fuels and commodity chemicals through metabolic engineering. Our present findings suggest that the ATP required for the synthetic pathway could potentially be supplied through an enhanced AEF pathway to further increase the efficiency of cyanobacterial cell factories.

Conclusions
Overexpression of the flv3 gene, encoding an NAD(P)H: oxygen oxidoreductase, enhanced photosynthetic electron transport and ATP supply through the regeneration of NADP⁺ in the recombinant Synechocystis sp. PCC6803. We also observed improvement of CO₂ fixation and carbon assimilation in central metabolic pathways such as the Calvin cycle and glycogen biosynthesis in the flv3-overexpressing strain by the combination of in vivo ¹³C-labeling of metabolites and metabolome analysis, which would be due to the enhancement in the intracellular ATP level. The flv3 overexpression improved not only the cell growth but also glycogen production in the Synechocystis, which would be a promising approach for the development of cyanobacterial biofuel production.

Methods
Strains and culture conditions
A GT strain of Synechocystis sp. PCC6803 [28], an flv3-overexpressing strain (Flv3ox), and its vector control (VC) strain were grown in BG11 liquid medium [29]. Cells were pre-cultivated in 500-mL flasks containing BG11 medium in an NC350-HC plant chamber (Nippon Medical and Chemical Instruments Co., Ltd., Osaka, Japan) under continuous irradiation at 50 μmol white light photons m⁻² s⁻¹ and 0.04% (v/v) CO₂ condition with 100-rpm agitation at 30°C. Cultivation was performed in a closed double-deck flask, which had a first stage containing 50 mL of 2 M NaHCO₃/Na₂CO₃ buffer with the appropriate pH to obtain the desired CO₂ concentration of 1% (v/v), and a second stage containing 70 mL BG11 culture medium [30,31]. Pre-cultured cells were inoculated into the fresh BG11 medium at a biomass concentration of 1.26 mg DCW L⁻¹ (the optical density at 750 nm [OD₇₅₀] was 0.04) and cultivated under continuous irradiation at 50 or 120 μmol white light photons m⁻² s⁻¹ and 1% (v/v) CO₂ with 100-rpm agitation at 30°C. Light intensity was measured in the middle of the medium using an LI-250A light meter equipped with an LI-190SA quantum sensor (LI-COR, Lincoln, NE). Cell density in the medium was determined as DCW, as a linear correlation was observed between DCW and optical density measured at OD₇₅₀ using a UV mini spectrophotometer (Shimadzu, Kyoto, Japan). For plate cultures, BG-11 was solidified using 1.5% (w/v) agar (BD Biosciences, San Jose, CA) and incubated in air at 30°C under continuous white light irradiation (50 to 70 μmol photons m⁻² s⁻¹).

Construction of recombinant strains
The flv3 (sll0550) coding region was amplified from Synechocystis sp. PCC6803 genomic DNA by PCR using the primer set 5′-GGAATTTAACCATAATGTTCACT ACCCCCCCTCCC-3′ and 5′-GGCATGGAGGACATATA TAGTAATAATTGCGACTT-3′. The resulting 1.7-kb fragment was integrated into NdeI-digested pTCP2031V [32] using an In-Fusion HD Cloning Kit (Takara Bio, Shiga, Japan) to yield pTCP2031-flv3. The GT strain was transformed with pTCP2031V-flv3 using a previously described method [33] to yield strain Flv3ox. Also, the plasmid pTCP2031V was introduced into the GT strain to yield the VC strain. Colonies resistant to 34 μg mL⁻¹ chloramphenicol were selected, and isolation of a single colony was repeated three times. The chromosomal integration of flv3 and plasmid-derived sequence was confirmed by PCR using the specific primers, Pr1 (5′-GATGGGCAATGCGTGAAGAACCGTGGTTGAC-3′) and Pr2 (5′-CGGTAAATTGCGACTTTGACCGTCAAAATAAC-3′), pTCP2031V was a kind gift from Professor M. Ikeuchi (University of Tokyo, Japan).

Immunoblot analysis
Cellular proteins were extracted from 1.6 mg DCW Synechocystis cells with 50 μL of 5% lithium lauryl sulfate containing 75 mM dithiothreitol by sonication in a Bioruptor UCD-200 bath sonicator (Cosmo Bio, Tokyo, Japan). After centrifugation of the mixture at 15,000 g for 5 min at 4°C, 10 μL of the extracts were subjected to sodium dodecyl sulfate-polyacrylamide gel electrophoresis on 12% and 1.5% (w/v) agar (BD Biosciences, San Jose, CA) and visualized with 5-bromo-4-chloro-3-indolyl-phosphate/nitroblue tetrazolium after incubation with alkaline phosphatase-conjugated Goat Anti-Rabbit IgG antibody (Promega, Madison, WI). Proteins separated on the 12% gel were blotted onto a polyvinylidene fluoride membrane, reacted with rabbit anti-Flv3 antibody, which was kindly provided by Dr. H. Yamamoto (Kyoto University, Japan), and visualized with 5-bromo-4-chloro-3-indolyl-phosphate/nitroblue tetrazolium after incubation with alkaline phosphatase-conjugated Goat Anti-Rabbit IgG antibody (Promega, Madison, WI). Proteins separated on the 12% gel were stained with Coomassie brilliant blue R-250 (Nacalai Tesque, Inc., Kyoto, Japan). Protein concentrations were determined by using the QuantiPro BCA assay kit (Sigma-Aldrich, St. Louis, MO), with bovine serum albumin as the standard.
Metabolite analysis
Cyanobacterial cells equivalent to 5 mg DCW were collected from the cultivation flasks and immediately filtered using 1-μm pore size polytetrafluoroethylene (PTFE) filter disks (Omnipore; Millipore, Billerica, MA). After washing the filter disks with 20 mM ammonium carbonate pre-chilled to 4°C, the cells retained on the filters were immediately placed into 2 mL pre-cooled (-30°C) methanol containing 37.38 μM L-methionine sulfone and 37.38 μM piperazine-1,4-bis(2-ethanesulfonic acid) as internal standards for mass analysis. Intracellular metabolites were extracted according to a previously reported method [21], with minor modification. The cells were suspended by vortexing, and then 0.5 mL of the cell suspension was mixed with 0.2 mL of pre-cooled (4°C) water and 0.5 mL of chloroform. After vortexing for 30 s, phase separation of aqueous and organic layers was performed by centrifugation at 14,000 g for 5 min at 4°C. 500 μL of the aqueous layer was filtered with a Millipore 5 kDa cut-off membrane for the removal of solubilized proteins. After evaporation of the aqueous-layer extracts under vacuum using a FreeZone 2.5 Plus freeze dry system (Labconco, Kansas City, MO), the dried metabolites were dissolved in 20 μL of Milli-Q water. The metabolites were analyzed using a CE/MS system (CE, Agilent G7100; MS, Agilent G6224AA LC/MSD TOF; Agilent Technologies, Palo Alto, CA) controlled by MassHunter Workstation Data Acquisition software (Agilent Technologies), as described previously [21].

13C-labeling experiment
To analyze metabolic turnover in cyanobacteria, in vivo 13C-labeling was performed according to a previously reported method [21]. Briefly, Synechocystis cells were cultivated in BG11 medium under 120 μmol photons m-2 s-1 light intensity and 1% (v/v) CO2 conditions at 30°C. When the OD750 reached approximately 4.5, cells were harvested using PTFE filter disks, as described above. After washing the filters with 20 mM ammonium carbonate, the cells on the filter were immediately frozen in liquid nitrogen, and then freeze-dried using a Labconco freeze dryer. After glycogen was extracted from the cells, the glycogen content was determined by high performance liquid chromatography (HPLC) (Shimadzu) using a size-exclusion HPLC column (OHpak SB-806 M HQ; Shimadzu) and reflective index detector (RID-10A; Shimadzu), as described previously [36].

Glycogen measurement
When the OD750 of Synechocystis cultures reached approximately 4.5, cells were harvested using PTFE filter disks, as described above. After washing the filters with 20 mM ammonium carbonate, the cells on the filter were immediately frozen in liquid nitrogen, and then freeze-dried using a Labconco freeze dryer. After glyco- gen was extracted from the cells, the glycogen content was determined by high performance liquid chromatography (HPLC) (Shimadzu) using a size-exclusion HPLC column (OHpak SB-806 M HQ; Shimadzu) and reflective index detector (RID-10A; Shimadzu), as described previously [36].

Photosynthesis analysis
Oxygen exchange was monitored with an oxygen electrode (Hansatech, King’s Lynn, UK). For the measurement, a reaction mixture (2 mL) containing 50 mM HEPES-KOH (pH 7.6), 10 mM NaHCO3 and an arbitrary amount of cells (10 μg chlorophyll mL-1) was first incubated in the dark for 10 min under air-equilibrated conditions and was then illuminated with actinic light (red light of wavelength >640 nm) at the indicated light intensity at 25°C. During the measurements, the reaction mixture was mixed with a magnetically controlled micro-stirrer.

Additional file

Additional file 1: Concentration of cellular biomass cultivated under 120 μmol photons m-2 s-1 light intensity and 1% CO2 conditions. Difference in biomass concentration was statistically analyzed by one-way ANOVA. A post hoc Tukey’s Honestly Significant Difference test was carried out on the grouped means. Values are the averages from measurements of ten different cell cultures, ± SD. Values followed by the same letter are not significantly different (P > 0.01).

Abbreviations
AcCoA: acetyl-CoA; ADP-Glc: ADP-glucose; AEF: alternative electron flow; CE: capillary electrophoresis; DCW: dry cell weight; Flv: flavodiiron; F6P: fructose-6-phosphate; G1P: glucose-1-phosphate; G6P: glucose-6-phosphate; GT: glucose-tolerant; LEF: linear electron flow; MS: mass spectrometry; PEP: phosphoenolpyruvate; 2PGA: 2-phosphoglycerate; 3PGA: 3-phosphoglycerate; PTFE: polytetrafluoroethylene; RuBP: ribulose-1,5-bisphosphate; S7P: sedoheptulose-7-phosphate; VC: vector control; WWC: water-water cycle.

Competing interests
The authors declare that they have no competing interests.
Authors’ contributions
TH supervised the research and drafted the manuscript. MM carried out the cultivation of recombinant strains, the metabolite analysis, the 13C-labeling experiment, and the glycerogen measurement. YS participated in the strain cultivation and glycerogen measurement. SA carried out the immunoblot analysis. MT constructed the recombinant strains. GS carried out the photosynthesis analysis. CM assisted in drafting the manuscript. AK supervised the research. All authors have read and approved the manuscript.

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Author details
1 Organization of Advanced Science and Technology, Kobe University, 1-1 Rokkodai, Nada, Kobe 657-8501, Japan. 2 Precursory Research for Embryonic Science and Technology (PRESTO), Japan Science and Technology Agency, 3-5 Sanbancho, Chiyoda-ku, Tokyo 102-0075, Japan. 3 Department of Chemical Science and Engineering, Graduate School of Engineering, Kobe University, 1-1 Rokkodai, Nada, Kobe 657-8501, Japan. 4 Kobe University Center for Inland Sea, 1-1 Rokkodai, Nada, Kobe 657-8501, Japan. 5 Department of Biological and Environmental Science, Faculty of Agriculture, Graduate School of Agricultural Science, Kobe University, 1-1 Rokkodai, Nada, Kobe 657-8501, Japan. 6 Core Research for Evolutional Science and Technology (CREST), Japan Science and Technology Agency, 3-5 Sanbancho, Chiyoda, Tokyo 102-0075, Japan. 7 Biomass Engineering Program, RIKEN, 1-7-22 Suehiro, Tsurumi-ku, Yokohama 230-0045, Japan.

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