Interaction of the GntR-family transcription factor Sll1961 with thioredoxin in the cyanobacterium Synechocystis sp. PCC 6803

Junichi Kujirai1, Sato Nanba1, Taro Kadowaki1, Yoshiki Oka2, Yoshitaka Nishiyama1, Yuuki Hayashi2, Munehito Arai2 & Yukako Hihara1

Changes in the redox state of the photosynthetic electron transport chain act as a signal to trigger acclimation responses to environmental cues and thioredoxin has been suggested to work as a key factor connecting the redox change with transcriptional regulation in the cyanobacterium Synechocystis sp. PCC 6803. We screened for redox-dependent transcription factors interacting with thioredoxin M (TrxM) and isolated the GntR-type transcription factor Sll1961 previously reported to be involved in acclimation responses of the photosynthetic machinery. Biochemical analyses using recombinant Sll1961 proteins of wild type and mutants of three cysteine residues, C124, C229 and C307, revealed that an intramolecular disulfide bond is formed between C229 and C307 under oxidizing conditions and TrxM can reduce it by attacking C307. Sll1961 exists in a dimeric form of about 80 kDa both under reducing and oxidizing conditions. C124 can form an intermolecular disulfide bond but it is not essential for dimerization. Based on these observations, tertiary structure models of the Sll1961 homodimer and the Sll1961-TrxM complex were constructed.

Photosynthetic organisms perceive changes in environmental cues as the redox changes of the electron transfer components located in the photosynthetic electron transport chain and start acclimation responses to balance the photosynthetic energy supply and its consumption by various metabolic reactions1,2. DNA microarray analysis of the cyanobacterium Synechocystis sp. PCC 6803 (S.6803) with or without addition of inhibitors of photosynthetic electron transport has suggested that the redox state of components located downstream of the plastoquinone pool is critical for transcriptional regulation both under low light3 and high light conditions1.

Recently, thioredoxin (Trx), a ubiquitous small redox protein that regulates the activity of various proteins through dithiol-disulfide-exchange reactions4, has been highlighted as a key factor connecting the redox change of the photosynthetic electron transport chain with transcriptional regulation. We reported that a small LuxR-type transcription factor PedR (Ssl0564) interacts with Trx to achieve redox-dependent transcriptional regulation in S.68035,6. Elevation of photon flux density causes activation of photosynthetic electron transport and increase in the availability of reducing equivalents at the acceptor side of photosystem I. The redox change is transmitted to PedR via interaction with reduced form of Trx, leading to a transient conformational change and inactivation of PedR. As the acclimation responses take place to mitigate the over-reduction of the photosynthetic electron transport chain, PedR returns to the active form again. This mechanism seems to enable transient induction or repression of the target genes in response to sudden changes in photon flux density.

In the heterocystous cyanobacterium Anabaena sp. strain PCC 7120, an ArsR-type transcription factor RexT (Alr1867) was reported as a redox-active transcriptional repressor of the trxA2 gene (all1866)7. Binding activity of RexT to the trxA2 promoter is lost through the formation of an intramolecular disulfide bond between two cysteine residues in the presence of H2O2, whereas the binding activity can be restored by interaction with reduced TrxA2.

1Graduate School of Science and Engineering, Saitama University, 255 Shimo-Okubo, Sakura-ku, Saitama, 338-8570, Japan. 2Department of Life Sciences, Graduate School of Arts and Sciences, The University of Tokyo, 3-8-1 Komaba, Meguro-ku, Tokyo, 153-8902, Japan. Correspondence and requests for materials should be addressed to Y.Hihara (email: hihara@mail.saitama-u.ac.jp)
There have been several reports on isolation of interacting partners of Trx in S.6803 using Trx affinity chromatography\(^1\)-\(^3\). However, transcription factors have not been isolated so far, probably due to the technical difficulty identifying regulatory proteins with low abundance. Thus we established a new screening system to detect specific interaction between a transcription factor and Trx within *Escherichia coli* co-expression strain. We first focused on ten OmpR-type response regulators that are encoded in the S.6803 genome and identified three of them, RpaA (Slr0115), RpaB (Slr0947) and ManR (Slr1837), as new candidates of TrxM interacting partner\(^11\). Furthermore, we screened 29 transcription factors having cysteine residues conserved among cyanobacterial orthologs and identified two more candidates. In this study, we focused on one of them, GntR-type transcription factor Sll1961. The gene-disrupted mutant of *sll1961* has been reported to have a defect in regulation of photosystem stoichiometry under high light conditions\(^12\) and in phycobilisome degradation during nitrogen starvation\(^13\), implying contribution of Sll1961-Trx interaction to acclimation responses of the photosynthetic machinery. Biochemical analyses using recombinant Sll1961 proteins of wild-type (WT) and mutants of three cysteine residues revealed that TrxM can reduce the intramolecular disulfide bond between C229 and C307 in Sll1961. Based on these experimental results, the structure model of the Sll1961-Trx complex was constructed.

**Results**

**Interaction between Sll1961 and TrxM detected in *E. coli* co-expression strain.** We detected interaction between Sll1961 and TrxM using the previously established screening system\(^11\). In this system, a His-tagged Sll1961 (His-Sll1961) and S-tagged TrxM\(_{C35S}\) are co-expressed in *E. coli* Origami2 (DE3) cells carrying mutations in glutathione reductase and thioredoxin reductase genes to facilitate disulfide bonds formation in the cytoplasm. TrxM\(_{C35S}\) is a mutant TrxM protein whose active site C35 has been substituted with a serine residue. This substitution enables the formation of a stable complex between TrxM\(_{C35S}\) and the target protein, since thi-ol-disulfide exchange reaction is halted at the stage of the formation of the mixed-disulfide intermediate. Four different *E. coli* strains were used for this experiment: a control Origami2 strain; a Trx strain expressing only TrxM\(_{C35S}\); a Sll1961 strain expressing only Sll1961; and a Trx-Sll1961 strain expressing both TrxM\(_{C35S}\) and Sll1961. After induction of recombinant proteins by adding IPTG, soluble protein fractions of each *E. coli* strain were analyzed by non-reducing SDS-PAGE and following immunodetection. When using an anti His-tag antibody to detect His-Sll1961 protein (Fig. 1A, lanes 1–6), a 40 kDa band of the Sll1961 monomer (gray arrow) was additionally detected in that of the Trx-Sll1961 strain (lane 5) but disappeared by incubation with DTT before electrophoresis (lane 6). When using S-protein to detect the S-tagged TrxM\(_{C35S}\) (lanes 7–12), many bands were detected in the Trx strain (lane 8) but they disappeared, other than a 18 kDa Trx monomer band (white arrow), following incubation with DTT (lane 9). These bands might be complexes between Trx and endogenous *E. coli* proteins. In the Trx-Sll1961 strain protein extract, a distinct band of 58 kDa was detected (lane 11) instead of many faint bands observed in the Trx strain, but disappeared after treatment with DTT (lane 12). The 58 kDa band, which can be detected by both anti His-tag antibody and S-protein in the absence of DTT, must be the complex of the monomeric Sll1961 and the monomeric Trx.

Sll1961 possesses three cysteine residues, C124, C229 and C307, some of which can work as interacting partners of Trx. We expected that examination of the degree of conservation of cysteine residues among Sll1961 orthologs universally present in cyanobacterial species\(^14\) may give us implication for the physiological significance of three cysteine residues in Sll1961. Supplementary Fig. S1 shows the phylogenetic tree of the Sll1961 orthologs in 41 representative cyanobacterial species and the plastid (cyanelle) of *Paulinella chromatophora* in 41 representative cyanobacterial species and the plastid (cyanelle) of *Paulinella chromatophora*. In 41 representative cyanobacterial species and the plastid (cyanelle) of *Paulinella chromatophora*, some cyt-124 was conserved in 30 species out of 42, whereas C124 was only present in S.6803. It may be possible that C124 in S.6803 is used as an alternative over C134 which is widely conserved within other species (38 species out of 42). In addition, there exist some cysteine residues well-conserved in cyanobacteria other than S.6803, such as C280 present in 31 species out of 42 and C23 highly conserved in marine species.

In order to confirm the Sll1961-Trx complex formation and to examine the contribution of each cysteine residue to the complex formation, His-Sll1961 whose cysteine residue was substituted with alanine was co-expressed with S-tagged TrxM\(_{C35S}\). Reducing SDS-PAGE revealed that the amounts of accumulated His-Sll1961 and S-Trx in the soluble protein fraction were similar among co-expression strains, Trx-Sll1961\(_{C35S}\), Trx-Sll1961\(_{C229A}\) and Trx-Sll1961\(_{C307A}\). (Fig. 1B). By non-reducing SDS-PAGE followed by immunoblot analysis using an anti His-tag antibody (Fig. 1C) and S-protein (Fig. 1D,E), the 58 kDa band (black arrow) can be detected in the Trx-Sll1961\(_{C35S}\) strain (lane 11) in addition to many faint bands observed in the Trx strain, but disappeared after treatment with DTT (lane 12). The 58 kDa band, which can be detected by both anti His-tag antibody and S-protein in the absence of DTT, must be the complex of the monomeric Sll1961 and the monomeric Trx.

**Analysis of the Sll1961-Trx interaction using purified proteins.** The experiment using *E. coli* co-expression strain revealed that Sll1961 interacts with TrxM and C307 of Sll1961 is essential for the interaction. To confirm this, His-Sll1961\(_{C35S}\), His-Sll1961\(_{C229A}\), His-Sll1961\(_{C307A}\) and His-Sll1961\(_{C35S}\), His-Sll1961\(_{C229A}\), proteins were purified from the respective single-expression strains by nickel affinity chromatography. The purified Sll1961 proteins were detected as monomer by non-reducing SDS-PAGE after treatment with 100 mM DTT (Supplementary Fig. S2). Treatment with 500 μM of thiol-specific oxidizing agent diamide resulted in detection of the faint oligomer bands, but large fractions of Sll1961 protein remained as monomer.

In order to examine whether three cysteine residues in Sll1961 are redox active or not, WT and cysteine mutants of His-Sll1961 protein were incubated with DTT or diamide, treated with the thiol-modifying agent...
PEG-maleimide (average molecular mass: 5 kDa), and then subjected to non-reducing SDS-PAGE (Fig. 2A,B). When His-Sll1961 WT treated with 100 mM DTT was further incubated with PEG-maleimide, a significant mobility shift was observed (Fig. 2A, lane 2) compared to the sample without incubation with PEG-maleimide (lane 1). This indicates the attachment of PEG-maleimide to all three cysteine residues (denoted as “3PEG” in Fig. 2). On the other hand, when His-Sll1961 WT was treated with diamide and incubated with PEG-maleimide, the 3PEG band gradually decreased and a single faster migrating band appeared in a diamide concentration-dependent manner (lanes 3–6). The faster migrating band is “1PEG”, showing that two cysteine residues were oxidized. It is of note that oxidation of all three cysteines (0PEG) or only one cysteine (2PEG) was hardly observed in His-Sll1961 WT.
When His-Sll1961 C124A treated with 100 mM DTT was modified with PEG-maleimide, the attachment of PEG-maleimide to C229 and C307 resulted in band shift from 0PEG (lane 7) to 2PEG (lane 8). Oxidization with diamide resulted in decrease of the 2PEG band and increase of the 0PEG band in a diamide concentration-dependent manner (lanes 9–12). The 1PEG band was not detected, indicating oxidization of C229 and C307 proceeds simultaneously. Figure 2B shows the quite similar results on the redox change of His-Sll1961 C229A and His-Sll1961 C307A. In both cases, the 2PEG band was detected after treatment with 100 mM DTT and PEG-maleimide (lanes 2 and 8). When treated with diamide, the 2PEG and 1PEG bands with similar intensity were detected irrespective of diamide concentration (lanes 3–6, 9–12), indicating coexistence of proteins in which one of two cysteine residues became oxidized or both were insensitive to oxidation. There seems no simultaneous oxidization of C124 and C307, or C124 and C229. Based on these observations, we supposed that an intramolecular disulfide bond is formed between C229 and C307 in the oxidized His-Sll1961 WT and this redox state was detected as 1PEG.

Next, we examined the effect of addition of TrxM to the oxidized His-Sll1961 (Fig. 2C,D). In these experiments, TrxM WT but not TrxM C35S was used in order to examine the native thiol-disulfide exchange reaction. Since the electrophoretic mobility of TrxM modified with PEG-maleimide (lane 13) was similar to that of Sll1961 without modification (0PEG), addition of 5 μM TrxM resulted in detection of the band at 0PEG position (lanes 5 and 11). In His-Sll1961 WT treatment with 500 μM diamide resulted in detection of the 1PEG band (Fig. 2C, lane 2) as observed in Fig. 2A. Although incubation with 0.1 mM DTT had no effect on this band pattern (lane 3), addition of 0.5 or 5 μM TrxM concomitantly with 0.1 mM DTT resulted in decrease in the 1PEG band and increase in the 3PEG band (lanes 4 and 5). The similar effect of TrxM was observed in the case of His-Sll1961 C124A. Namely, incubation with 0.1 mM DTT had no effect on the 0PEG band formed by treatment with 500 μM diamide (lane 9), whereas addition of 0.5 or 5 μM TrxM concomitantly with 0.1 mM DTT resulted in decrease in the 0PEG band and increase in the 2PEG band (lanes 10 and 11). In the case of His-Sll1961 C229A and His-Sll1961 C307A, the intensity of 2PEG and 1PEG bands was not affected by the addition of TrxM (Fig. 2D, lanes 3–5 and 9–11). These results clearly show that TrxM can reduce the intramolecular disulfide bond between C229 and C307 in the oxidized His-Sll1961 WT and this redox state was detected as 1PEG.

Analysis of oligomerization state of Sll1961 by size exclusion chromatography. Although most GntR-family transcription factors are known to exist as a homodimer15, dimerization of Sll1961 has seldom been detected in our electrophoretic experiments. This may be due to the instability of the Sll1961 dimer in the presence of SDS. Thus we performed size exclusion chromatography to examine the oligomerization state of Sll1961. WT and cysteine mutants of His-Sll1961 protein were reduced with 100 mM DTT for 15 min at room temperature.
TrxM induces conformational changes of the N-terminal DBD, which may affect the DNA-binding ability of Sll1961. W130 is located at the N-terminal DNA-binding domain (DBD), these results suggest that reduction of Sll1961 by TrxM induces conformational changes of the protein, indicating that W130 is more exposed to solvent in the reduced form than in the oxidized form. Since W130 is involved in the overall structure of Sll1961 (Fig. 5A), the spectra showed two minima at 208 and 222 nm, indicating the presence of α-helices and β-sheets. Indeed, the secondary structure analysis of the CD spectrum of the oxidized form using six different programs indicated that the protein contains, on average, 30 ± 5% α-helices, 20 ± 4% β-sheets, and 50 ± 2% coils (Supplementary Table S1). These results suggest that Sll1961 consists of a structural domain(s) that are classified into the α/β class, in which the secondary structure is composed of α-helices and (mostly anti-parallel) β-strands that occur separately16, and/or the α/β class, in which the secondary structure is composed of alternating α-helices and parallel β-strands18. When the protein was reduced, it contained, on average, 33 ± 4% α-helices, 17 ± 3% β-sheets, and 50 ± 2% coils, indicating that reduction of a disulfide bond in Sll1961 induces a slight increase in helical structures and a concomitant decrease in sheet structures.

Fluorescence spectrum of a single tryptophan residue (W130) of Sll1961 in the oxidized form had a peak at 344 nm (Fig. 5B). The intensity at 216–219 nm is similar to that at 222 nm, suggesting the presence of α-helices. 3% ± 5% α-helices, 20 ± 4% β-sheets, and 50 ± 2% coils (Supplementary Table S1). These results suggest that Sll1961 consists of a structural domain(s) that are classified into the α/β class, in which the secondary structure is composed of α-helices and (mostly anti-parallel) β-strands that occur separately16, and/or the α/β class, in which the secondary structure is composed of alternating α-helices and parallel β-strands18. When the protein was reduced, it contained, on average, 33 ± 4% α-helices, 17 ± 3% β-sheets, and 50 ± 2% coils, indicating that reduction of a disulfide bond in Sll1961 induces a slight increase in helical structures and a concomitant decrease in sheet structures.

Fluorescence spectrum of a single tryptophan residue (W130) of Sll1961 in the oxidized form had a peak at 344 nm (Fig. 5B). The intensity at 216–219 nm is similar to that at 222 nm, suggesting the presence of α-helices. 3% ± 5% α-helices, 20 ± 4% β-sheets, and 50 ± 2% coils (Supplementary Table S1). These results suggest that Sll1961 consists of a structural domain(s) that are classified into the α/β class, in which the secondary structure is composed of α-helices and (mostly anti-parallel) β-strands that occur separately16, and/or the α/β class, in which the secondary structure is composed of alternating α-helices and parallel β-strands18. When the protein was reduced, it contained, on average, 33 ± 4% α-helices, 17 ± 3% β-sheets, and 50 ± 2% coils, indicating that reduction of a disulfide bond in Sll1961 induces a slight increase in helical structures and a concomitant decrease in sheet structures.

Fluorescence spectrum of a single tryptophan residue (W130) of Sll1961 in the oxidized form had a peak at 344 nm (Fig. 5B). The intensity at 216–219 nm is similar to that at 222 nm, suggesting the presence of α-helices. 3% ± 5% α-helices, 20 ± 4% β-sheets, and 50 ± 2% coils (Supplementary Table S1). These results suggest that Sll1961 consists of a structural domain(s) that are classified into the α/β class, in which the secondary structure is composed of α-helices and (mostly anti-parallel) β-strands that occur separately16, and/or the α/β class, in which the secondary structure is composed of alternating α-helices and parallel β-strands18. When the protein was reduced, it contained, on average, 33 ± 4% α-helices, 17 ± 3% β-sheets, and 50 ± 2% coils, indicating that reduction of a disulfide bond in Sll1961 induces a slight increase in helical structures and a concomitant decrease in sheet structures.

Fluorescence spectrum of a single tryptophan residue (W130) of Sll1961 in the oxidized form had a peak at 344 nm (Fig. 5B). The intensity at 216–219 nm is similar to that at 222 nm, suggesting the presence of α-helices. 3% ± 5% α-helices, 20 ± 4% β-sheets, and 50 ± 2% coils (Supplementary Table S1). These results suggest that Sll1961 consists of a structural domain(s) that are classified into the α/β class, in which the secondary structure is composed of α-helices and (mostly anti-parallel) β-strands that occur separately16, and/or the α/β class, in which the secondary structure is composed of alternating α-helices and parallel β-strands18. When the protein was reduced, it contained, on average, 33 ± 4% α-helices, 17 ± 3% β-sheets, and 50 ± 2% coils, indicating that reduction of a disulfide bond in Sll1961 induces a slight increase in helical structures and a concomitant decrease in sheet structures.

Fluorescence spectrum of a single tryptophan residue (W130) of Sll1961 in the oxidized form had a peak at 344 nm (Fig. 5B). The intensity at 216–219 nm is similar to that at 222 nm, suggesting the presence of α-helices. 3% ± 5% α-helices, 20 ± 4% β-sheets, and 50 ± 2% coils (Supplementary Table S1). These results suggest that Sll1961 consists of a structural domain(s) that are classified into the α/β class, in which the secondary structure is composed of α-helices and (mostly anti-parallel) β-strands that occur separately16, and/or the α/β class, in which the secondary structure is composed of alternating α-helices and parallel β-strands18. When the protein was reduced, it contained, on average, 33 ± 4% α-helices, 17 ± 3% β-sheets, and 50 ± 2% coils, indicating that reduction of a disulfide bond in Sll1961 induces a slight increase in helical structures and a concomitant decrease in sheet structures.

Fluorescence spectrum of a single tryptophan residue (W130) of Sll1961 in the oxidized form had a peak at 344 nm (Fig. 5B). The intensity at 216–219 nm is similar to that at 222 nm, suggesting the presence of α-helices. 3% ± 5% α-helices, 20 ± 4% β-sheets, and 50 ± 2% coils (Supplementary Table S1). These results suggest that Sll1961 consists of a structural domain(s) that are classified into the α/β class, in which the secondary structure is composed of α-helices and (mostly anti-parallel) β-strands that occur separately16, and/or the α/β class, in which the secondary structure is composed of alternating α-helices and parallel β-strands18. When the protein was reduced, it contained, on average, 33 ± 4% α-helices, 17 ± 3% β-sheets, and 50 ± 2% coils, indicating that reduction of a disulfide bond in Sll1961 induces a slight increase in helical structures and a concomitant decrease in sheet structures.
The protein BLAST search of Sll1961 (residues 1–343) showed that residues 13–78 corresponds to the winged helix-turn-helix (W-HTH) DNA-binding domain of the GntR family of transcriptional regulators and that residues 3–133 corresponds to a conserved domain, having high sequence similarity with a member of the GntR family, transcriptional regulator YhcF. The GntR family of transcription factors, which was named after the gluconate operon repressor from *Bacillus subtilis*, constitutes one of the most prevalent superfamilies of bacterial transcription factors. Members of the GntR family have the N-terminal DBD and the C-terminal effector-binding domain (EBD) which is also involved in dimerization. Most of GntR family regulators form a homodimer and bind to either inverted or direct repeat sequences. On the other hand, the EBD shows high heterogeneity and can be classified into several subgroups, FadR, HutC, MocR, YtrA, AraR, DevA and PlmA. Structures of EBD are diverse among subgroups of GntR. For example, FadR and YtrA consist of all-α structures; HutC consists of a core of six-stranded anti-parallel β-sheet surrounded by three α-helices; AraR is composed of two subdomains, both having a core of a six-stranded parallel β-sheet flanked on each side by two α-helices; and MocR is composed of two subdomains, each having a core of a large parallel or a small anti-parallel β-sheet surrounded by α-helices.

**Figure 4.** Mass spectrometry analysis of redox-treated His-Sll1961 WT. (A) Thiol-redox state of reduced and oxidized samples detected by modification with PEG-maleimide. The reduced (Red) and oxidized (Ox) samples were prepared by incubation with 20 mM DTT for 15 min and with 1 mM diamide for 1 h, respectively. (B) MALDI-TOF mass spectra for the tryptic digests of reduced Sll1961. (C) MALDI-TOF mass spectra for the tryptic digests of oxidized Sll1961. Peptides containing cysteine residues were marked by arrows and labeled according to the residue number of the cysteine.
Sll1961 belongs to the PlmA subfamily, which is specific to cyanobacteria and shares very limited sequence homology with the other subfamilies. Secondary structure prediction by PSIPRED showed that Sll1961 has 44% α-helices, 16% β-sheets, and 40% coils (Supplementary Fig. S5), consistent with the CD results. The prediction indicated that residues 13–78 have propensities toward three α-helices followed by two β-strands, as expected for the W-HTH motif. On the other hand, residues 97–132 have propensities to form two long α-helices, while residues 137–324 have propensities toward eight repeats of alternating β-strand and α-helix (Supplementary Fig. S4). This suggests that the C-terminal EBD of Sll1961 forms α/β structure, consistent with the CD measurement. Residues 328–343 were predicted to have random coil-like structures (Supplementary Fig. S5). These secondary structure propensities in the EBD are distinct from other GntR family members whose structures are available in the Protein Data Bank.

Table 1. Effect of redox treatment on peak areas of tryptic fragments of His-Sll1961 containing cysteine residues. *Data normalization between reduced and oxidized samples were performed using the peak area of a tryptic fragment containing no cysteine residue (m/z = 1732.84). **ND, not detected.

| Tryptic fragments containing Cys (calculated m/z) | Peak areas in the reduced sample (observed m/z) | Peak areas in the oxidized sample (observed m/z)* | Oxi/Red ratio |
|-----------------------------------------------|-----------------------------------------------|-----------------------------------------------|---------------|
| C124 (1277.46)                                | 909 (1277.65)                                 | 827 (1277.62)                                 | 0.91          |
| C229 (3030.62)                                | 149 (3029.79)                                 | 9 (3030.67)                                   | 0.06          |
| C307 (1648.91)                                | 24 (1648.76)                                  | ND**                                         | —             |
| C124 + C124 (2552.92)                        | 68 (2552.22)                                  | 451 (2552.17)                                 | 6.66          |
| C124 + C229 (4306.07)                        | ND                                            | ND**                                         | —             |
| C124 + C307 (2924.37)                        | 730 (2923.39)                                 | 210 (2923.33)                                 | 0.29          |
| C229 + C229 (6059.23)                        | ND                                            | ND**                                         | —             |
| C229 + C307 (4677.53)                        | 81 (4676.83)                                  | 674 (4676.92)                                 | 8.28          |
| C307 + C307 (3295.82)                        | ND                                            | ND**                                         | —             |

Figure 5. CD spectra and predicted structures of Sll1961. (A) Far-UV CD spectra of His-Sll1961 WT in the oxidized (red) and reduced forms (blue). (B) Predicted structure of the Sll1961 homodimer. The sulfur atoms of the cysteine residues are shown by red spheres. DBD and EBD denote the N-terminal DNA-binding domain and C-terminal effector-binding domain, respectively. (C) Overlaid structures of the Sll1961 homodimer (gray) and the Sll1961-TrxM complex. The black arrow denotes the direction of possible movements of the N-terminal DBD upon binding of TrxM (yellow and orange) to the C-terminal EBD of Sll1961. (D) Predicted structure of the Sll1961-TrxM complex, in which two TrxM molecules are bound to the Sll1961 homodimer. The figures were drawn using the PyMOL Molecular Graphics System, Schrödinger, LLC.
that the number of β-strands in the core of each subdomain is six for AraR but is four for SII1961. The model structure suggests that SII1961 is expected to have 50% α-helices, 14% β-sheets, and 36% coils, consistent with the secondary structure prediction and CD results. Moreover, W130 in the N-terminal DBD was 45% exposed to solvent (Supplementary Fig. S4), consistent with the fluorescence measurements. These coincidences support the validity of the model structure.

By docking TrxM to the SII1961 homodimer, we obtained the predicted structure of the SII1961-TrxM complex (Fig. 5D). Two TrxM molecules bind to the outer side of the homodimer and are located close to the N-terminal W-HTH domain. Comparison of the model structures of the SII1961 homodimer and the SII1961-TrxM complex indicates that upon TrxM binding, one of the two W-HTH domains of the SII1961 homodimer changes its orientation to interact with TrxM (Fig. 5C). Such structural changes may affect the mutual orientation and/or the distance between the two W-HTH domains of the homodimer and, in turn, eliminate the DNA binding ability of SII1961. These models also suggest that binding of a single TrxM molecule to the SII1961 homodimer is sufficient to modulate the DNA binding activity of SII1961. Solving crystal structures of SII1961 with and without TrxM will elucidate the mechanism of SII1961 functions.

**Discussion**

Interaction between the GntR-family transcription factor SII1961 and Trx was detected by the screening system using *E. coli* co-expression strain (Fig. 1A). So far, we have identified five transcription factors out of 39 as interacting partners of Trx in total. Despite existence of cysteine residue(s) in all of 39 transcription factors, only five showed interaction with Trx, suggesting that the screening system could detect only specific interaction between Trx and transcription factors. Alanine substitution experiments revealed that C307 in SII1961 is essential for the interaction with Trx (Fig. 1C–E). It is well established that the first cysteine (C32) in the active site of Trx forms a mixed disulfide intermediate with a cysteine of the target protein and then the second cysteine (C35) attacks the mixed disulfide bond, leading to the reduction of the target protein and oxidation of Trx itself22. Based on this information, we propose that C32 of Trx interacts with C307 of SII1961. The high degree of conservation of C307 among SII1961 orthologs (Supplementary Fig. S1) well coincides with the biochemical data showing requirement of C307 for interaction with Trx.

Both WT and cysteine mutants of SII1961 exist in a dimeric form of about 80 kDa under reducing conditions (Fig. 3). Although oligomeric forms could be hardly observed in non-reducing SDS-PAGE (Supplementary Fig. S2), SII1961 is likely to be a dimer in its native state and cysteine residues do not contribute to dimerization. Thiol modification (Fig. 2) and mass spectrometry (Fig. 4) analyses of the WT protein revealed that an intramolecular disulfide bond is formed between C229 and C307 under the oxidizing conditions. Although the 0PEG band, which indicates oxidation of all three cysteines, was not detected in the thiol modification experiments (Fig. 2A,C), it was observed in the oxidized sample for mass spectrometry analysis (Fig. 4A). The result of mass spectrometry analysis (Fig. 4C, Table 1) suggests that both a C229-C307 intramolecular disulfide bond and a C124-C124 intermolecular disulfide bond could be formed in the oxidized WT dimer, resulting in the detection of the 0PEG band. We think the C124-C124 intermolecular disulfide bond is non-specifically formed under oxidized conditions. It is not essential for dimer formation as suggested by size exclusion chromatography (Fig. 3) and low conservation among SII1961 orthologs (Supplementary Fig. S1). In the case of C124A mutant, its oxidation resulted in the detection of the 0PEG, but not the 1PEG band (Fig. 2A,C). This again indicates the formation of a C229-C307 intramolecular disulfide bond. On the other hand, oxidation of C229A and C307A mutants yielded the 2PEG and 1PEG but not the 0PEG (Fig. 2B,D), which may be due to difference in sensitivity to oxidation between C124 and other two cysteines. There seem two possibilities for the origin of the 1PEG band, one is the C124-C124 intermolecular disulfide bond and other is the oxidation of one of cysteines to oxo forms such as sulfinic acid.

The incubation with TrxM resulted in the conversion from 1PEG to 3PEG in WT and from 0PEG to 2PEG in the C124A mutant (Fig. 2C) but had no effect on the 1PEG band of C229A and C307A mutants (Fig. 2D). This indicates that TrxM can reduce the C229-C307 intramolecular disulfide bond, but not other oxidized forms of cysteines. The predicted structure of the SII1961-TrxM complex (Fig. 5D) implies that interaction with TrxM causes structural changes of the W-HTH domain of SII1961, resulting in changes in DNA binding activity.

It has been reported that binding of effector molecules to EBD affects the conformation of DBD to regulate the DNA binding activity in the GntR-family transcription factors. For example, DNA binding of the FadR repressor regulating fatty acid metabolism in *E. coli* is specifically inhibited by binding of long chain fatty acyl-CoA compounds to EBD23. Similarly, DNA binding of AraR repressor regulating arabinose metabolism in *B. subtilis* is inhibited by binding of arabinose to EBD24. To date, there have been two reports on the functional characterization of SII1961 orthologs in cyanobacterial species other than *S.6803*. In the filamentous cyanobacterium *Anabaena* sp. PCC 7120, null mutants of *plmA* (all1076) were reported to be unable to maintain the copy number of several endogenous plasmids and exogenously introduced shuttle vector25. Recently, SII1961-ortholog in *Synechococcus elongatus* PCC 7942 was isolated by yeast three-hybrid screening as an interacting factor with PII-PipX complex involved in nitrogen regulatory network26. Authors suggested that activity of the transcription factor can be regulated in response to 2-oxoglutarate and ATP/ADP signals through interaction with PII-PipX complex.

Although none of the reports on GntR-family regulator in heterotrophic bacteria provide clues to the function of EBD of PlmA subfamily, the report on interaction with PII-PipX in *Synechococcus elongatus* and our findings on interaction with Trx in *S.6803* imply the possibility that the conformational change of EBD through protein-protein interaction resulted in changes in DNA binding activity. By the characterization of *sll1961*-disrupted mutant in *S.6803*, it has been suggested that SII1961 is involved in regulation of photosystem I content under high-light conditions27 and phycobilisome content under nitrogen-limited conditions28. Under
these conditions, Sll1961 may participate in transcriptional regulation in response to photosynthetic activities through interaction with Trx and to 2-oxoglutarate level through interaction with the PH-PipX complex.

Methods

**Bacterial strains and growth conditions.** *E. coli* XL1-blue cells were used for routine cloning and Origami2 (DE3) cells (Merck Millipore, Darmstadt, Germany) for protein expression. Bacterial cultures were grown in TB or 2xYT medium at 37 °C. When necessary, antibiotics were added at the following concentrations: ampicillin (100 μg ml⁻¹), kanamycin (20 μg ml⁻¹) and spectinomycin (20 μg ml⁻¹).

**Construction of *E. coli* strains expressing His-tagged Sll1961 and/or S-tagged TrxM_C35S.** The coding region of *sll1961* was amplified for the primer set 1961-F (5′-AACATAGCTACA GTTCGCAATT-3′) and 1961-R (5′-GCGTTAGGCAAGCTCTGCGCGAC-3′). The underlining indicates the recognition sequences of Ndel and BamHI, respectively. PCR products were cloned into the pT7Blue T-vector (Merck Millipore), excised from the pT7Blue vector with Ndel and BamHI and subcloned into the same restriction sites in the pET28a vector (Merck Millipore) to express proteins with an N-terminal 6xHis-tag. C229A, C229A and C307A mutations were introduced into the *sll1961* coding sequence using a KOD-plus-mutagenesis kit (TOYOBO, Osaka, Japan) with the following primer sets: C229A-F (5′-GTTTTGAAA ACCATGATTGCC-3′) and C229A-R (5′-AAGCTTTTCTACCTGGGAAGAAGG-3′), C229A-F (5′-GCGTTAG GCATTGTAGCGTCAGC-3′) and C229A-R (5′-GGATTCTTTCGCCAGGCAGTC-3′) and C307A-F (5′-GCCCCT GAAATTACATTGGCGATA-3′) and C307A-R (5′-ATAATTTCCGGCATCCGGGA-3′). The underlining indicates the substitution from cysteine- to alanine-codon. Each expression construct was transformed into Origami2 (DE3) competent cells to yield the wild type and cysteine mutants of the Sll1961 strains.

Cloning of the coding region of *trxM* (slr6023) into the pT7Blue T-vector, introduction of C35S mutation and subcloning into the pCDFDuet-1 vector (Merck Millipore) to express TrxM_C35S with a C-terminal S-tag were performed as described in Kadowaki et al. Cloning resulted construct was transformed into Origami2 (DE3) competent cells and competent cells of the Sll1961 strains described above, to yield the Trx strain and the Trx-Sll1961 strains, respectively.

**Detection of the interaction between S-tagged TrxM_C35S and His-tagged Sll1961 in *E. coli* cells.** *E. coli* Origami2 (DE3) strains, harboring each expression construct, were pre-cultured in 2 ml of the TB medium containing kanamycin (for the pET28a vector) or spectinomycin (for the pCDFDuet-1 vector) at 37 °C overnight. The pre-culture of each strain was inoculated into 4.5 ml of the 2xYT medium to OD₅₆₅ = 0.1 and the expression of Sll1961 and TrxM_C35S was induced by addition of 100 μM isopropyl-D-thiogalactopyranoside (IPTG) at the mid-log phase. The cells were allowed to grow for an additional 3 h at 37 °C and harvested by centrifugation at 5,800 g for 2 min. Cell pellets were resuspended in the binding buffer (20 mM sodium phosphate, pH 7.4, 500 mM NaCl, 20 mM imidazole) and disrupted by sonication with Sonifier 450 (Branson, Danbury, USA) for 2 min, with two pauses of 1 min each on ice. The cell lysate was centrifuged at 16,000 g for 5 min and the supernatant was subjected to SDS-PAGE on non-reducing 12% gels, blotted onto polyvinylidene difluoride membrane (Immobilon-P, Millipore, Billerica, USA), and probed with rabbit polyclonal antibodies to His-tag (Bethyl, Montgomery, USA) or S-protein HRP conjugate (Novagen). The bound antibodies were detected using goat anti-rabbit IgG secondary antibodies conjugated to horse radish peroxidase (Bio-Rad, Hercules, USA) and the chemiluminescence detection reagent, EzWestLumi (Atto, Tokyo, Japan).

**Expression and purification of His-Sll1961 proteins from *E. coli* cells.** His-tagged Sll1961 proteins were purified from the WT and cysteine mutants of Sll1961 strains. The pre-culture of each strain was seeded into 50 ml of 2xYT medium and expression of Sll1961 was induced by addition of 100 μM IPTG at the mid-log phase. The cells were allowed to grow for an additional 3 h at 37 °C and harvested by centrifugation. All protein purification procedures were performed at 4 °C. Cells were disrupted by sonication on ice in 1.5 ml of the binding buffer and the lysate was centrifuged at 16,000 g for 5 min. The resulting supernatant was added to a 400 μl of Ni²⁺-Sepharose resin (COSMOGEL His-Accept; Nacalai tesque, Kyoto, Japan) pre-equilibrated with the binding buffer. After rotation for overnight at 4 °C, the sample was centrifuged at 1,500 g for 5 min and the resin was resuspended in 1.5 ml of the binding buffer. This washing step was repeated four times with increasing concentration of imidazole (20–40 mM). Proteins were then eluted from the resin with 200 μl of the elution buffer (20 mM sodium phosphate, pH 7.4, 500 mM NaCl, 300 mM imidazole) by incubation at 4 °C for 15 min. The supernatant was collected, and the elution step was repeated once. Purified His-Sll1961 was desalted by a PD MiniTrap G-25 column (GE Healthcare, Chicago, USA) that had been equilibrated with 20 mM sodium phosphate, pH 7.4. The eluate was added with 10% (v/v) glycerol, frozen in liquid N₂ and stored at −80 °C before use.

Protein concentration was determined using Bio-Rad Protein Assay Kit (Bio-Rad) with bovine serum albumin as the standard. The purity of the proteins was assessed by fractionating an aliquot on an SDS-PAGE gel and staining with Coomassie Brilliant Blue (CBB).

**Redox treatments of His-Sll1961 and modification of thiol group of cysteine residues.** Oxidation of His-Sll1961 (5 μM) was performed by incubation with 500 μM diamide for 60 min at room temperature. Reduction of His-Sll1961 (5 μM) was performed by incubation with 100 mM DTT or 100 μM DTT in the presence or absence of the WT TrxM protein (0.5 μM or 5 μM) for 15 min at room temperature. After the redox treatments, proteins were precipitated with 10% (w/v) trichloroacetic acid, washed with ice-cold 20% acetone and the thiols groups of cysteine residues were then modified by incubation with 4 mM methoxypolyethylene glycol (PEG) maleimide (Nihon Yushi, Tokyo, Japan) at 4 °C overnight. Modified proteins were separated by non-reducing SDS-PAGE and stained with CBB.
Mass spectrometry analysis. 25 μg of His-Sll1961 protein was oxidized with 1 mM diamide for 60 min or reduced with 20 mM DTT for 15 min at room temperature. After desalting using Zeba Spin Desalting Column (Thermo Fisher, Waltham, USA) and protein quantification using Bio–Rad Protein Assay Kit, samples were subjected to digestion by Sequencing Grade Modified Trypsin (Promega, Fitchburg, USA) at a ratio of 20:1 (w/w) for 2 h at 37°C. After desalting with ZipTip (Millipore), 1 μL of the sample was mixed with 4 μL of matrix solution of α-cyano-4-hydroxy cinnamic acid and spotted on the sample plate. MALDI-TOF mass spectra were acquired with Autoflex II (Bruker, Bremen, Germany). Calibration was performed using a Peptide Calibration Standard (Bruker) spotted on the target position next to the sample.

Size exclusion chromatography. Size exclusion chromatography was performed using high performance liquid chromatography (HPLC) system (L-20AP, Shimadzu, Kyoto, Japan). WT and cysteine mutants of His-Sll1961 protein were reduced with 100 mM DTT for 15 min at room temperature, concentrated to 60 μM using Amicon Ultra-0.5 mL (MWCO 10 kDa, Millipore), centrifuged at 20,600 g for 30 min at 4°C, and filtrated using a 0.45 μm syringe filter (Millipore) before injection. Protein samples were injected into gel filtration columns, Superdex 75 increase 3.2/300 or Superdex 200 increase 3.2/300 (GE Healthcare), using a sample loop of 20 μL and eluted with the running buffer containing 20 mM sodium phosphate (pH 7.4), 150 mM NaCl and 3 mM DTT. The flow rate of the running buffer was 0.1 mL min⁻¹. Absorbance of eluted samples was monitored at 280 nm by the UV-VIS detector (SPD-20A, Shimadzu). Alcohol dehydrogenase (160 kDa), bovine serum albumin (BSA, 66 kDa), and ovalbumin (44 kDa) were used as the standards for the Superdex 200 increase 3.2/300 column.

Determination of molecular weights was carried out on the HPLC equipped with a Viscotek TDA 305 light scattering detector (Malvern Instruments, Malvern, UK) using the Superdex 75 increase 3.2/300 column. BSA was used as a standard for calibration of the static light scattering analysis. Data analysis for estimating molecular weights was performed with the OmniSEC software (Malvern). Errors in molecular weights were estimated from duplicate or triplicate measurements.

CD and fluorescence measurements. Far-UV CD spectra of His-Sll1961,WT in the reduced and oxidized forms were measured in the buffer containing 20 mM sodium phosphate (pH 7.4) and 150 mM NaCl with and without 3 mM DTT, respectively, at 0.2 mg mL⁻¹ protein concentration adjusted using Amicon Ultra-0.5 mL (MWCO 10kDa, Millipore). CD spectra were obtained on a J-805 spectropolarimeter (Jasco, Tokyo, Japan) by scanning from 250 nm to 200 nm in a quartz cuvette with a path length of 1 mm. The temperature for measurement was maintained at 25°C by a thermostat circulating water bath. Mean residue ellipticity was calculated as described previously, assuming dimer formation of the protein. Fluorescence spectra of His-Sll1961,WT were measured as described in Supplementary Text.

Structure modeling. The procedure for modeling tertiary structures of Sll1961 and the Sll1961-TrxM complex is summarized in Supplementary Fig. S6 and described in detail in Supplementary Text and Supplementary Table S2. Three-dimensional structure predictions were performed using the I-TASSER server. Docking predictions were performed using the ZDOCK server. Foldit Standalone was used for manual refinement of predicted structures.

References
1. Muramatsu, M. & Hihara, Y. Acclimation to high-light conditions in cyanobacteria: from gene expression to physiological responses. J. Plant Res. 125, 11–39 (2012).
2. Dietz, K.-J. Efficient high light acclimation involves rapid processes at multiple mechanistic levels. J. Exp. Bot. 66, 2401–2414 (2015).
3. Hisabori, T. J. Mol. Biol. 284, 4524–4528 (1987).
4. Hosoya-Matsuda, N. et al. Anti-oxidative stress system in cyanobacteria. Significance of type II peroxiredoxin and the role of 1-Cys peroxiredoxin in Synechocystis sp. strain PCC 6803. J. Bacteriol. 185, 1719–1725 (2003).
5. Horiuchi, M. et al. Towards a functional dissection of thioredoxin networks in plant cells. Photoschem. Photobiol. 83, 145–151 (2007).
6. Fujita, Y. & Fujita, T. The gluconate operon of Bacillus subtilis encodes its own transcriptional negative regulator. Proc. Natl Acad. Sci. USA 84, 4524–4528 (1987).
7. Fujita, Y. & Fujita, T. The gluconate operon of Bacillus subtilis encodes its own transcriptional negative regulator. Proc. Natl Acad. Sci. USA 84, 4524–4528 (1987).
8. Lindahl, M. & Florencio, F. J. Thioredoxin-linked processes in cyanobacteria are as numerous as in chloroplasts, but targets are different. Proc. Natl Acad. Sci. USA 100, 16107–16112 (2003).
9. Hisabori, T. et al. The mutant of sll1961, which encodes a putative transcriptional regulator, has a defect in regulation of photosystem PipX complex. FEBS Letters 582, 1093–1096 (2008).
10. Fujita, Y. & Fujita, T. The gluconate operon of Bacillus subtilis encodes its own transcriptional negative regulator. Proc. Natl Acad. Sci. USA 84, 4524–4528 (1987).
11. Fujita, Y. & Fujita, T. The gluconate operon of Bacillus subtilis encodes its own transcriptional negative regulator. Proc. Natl Acad. Sci. USA 84, 4524–4528 (1987).
19. Rigali, S., Derouaux, A., Giannotta, F. & Dusart, J. Subdivision of the helix-turn-helix GntR family of bacterial regulators in the FadR, HutC, ModR, and YtaR subfamilies. *J. Biol. Chem.* **277**, 12507–12515 (2002).
20. Hoskisson, P. A. & Rigali, S. Variation in form and function the helix-turn-helix regulators of the GntR superfamily. *Adv. Appl. Microbiol.* **69**, 1–22 (2009).
21. McCaffery, J. M., Böhm, C., & Jones, D. T. The PSIPRED protein structure prediction server. *Bioinformatics* **16**, 404–405 (2000).
22. Holmgren, A. Thioredoxin structure and mechanism: conformational changes on oxidation of the active-site sulphhydryl to a disulfide. *Structure* **3**, 239–243 (1995).
23. DiRusso, C. C., Heimert, T. L. & Metzger, A. K. Characterization of FadR, a global transcriptional regulator of fatty acid metabolism in *Escherichia coli*. Interaction with the *fadB* promoter is prevented by long chain fatty acyl coenzyme A. *J. Biol. Chem.* **267**, 8685–8691 (1992).
24. Franco, I. S., Mota, L. J., Soares, C. M. & de Sá-Nogueira, I. Functional domains of the *Bacillus subtilis* transcription factor AraR and identification of amino acids important for nucleoprotein complex assembly and effector binding. *J. Bacteriol.* **188**, 3024–3036 (2006).
25. Lee, M. H., Scherer, M., Rigali, S. & Golden, J. W. PlmA, a new member of the GntR family, has plasmid maintenance functions in *Anabaena* sp. strain PCC 7120. *J. Bacteriol.* **185**, 4315–4325 (2003).
26. Arazi, M. & Iwakura, M. Probing the interactions between the folding elements early in the folding of *Escherichia coli* dihydrofolate reductase by systematic sequence perturbation analysis. *J. Mol. Biol.* **347**, 337–353 (2005).
27. Roy, A., Kucukural, A. & Zhang, Y. I-TASSER: a unified platform for automated protein structure and function prediction. *Nat. Protoc.* **5**, 725–738 (2010).
28. Pierce, B. G. *et al.* ZDOCK Server: Interactive docking prediction of protein–protein complexes and symmetric multimers. *Bioinformatics* **30**, 1771–1773 (2014).
29. Kleffner, R. *et al.* Foldit Standalone: a video game-derived protein structure manipulation interface using Rosetta. *Bioinformatics* **33**, 2765–2767 (2017).

**Acknowledgements**

We thank Tomoko Kunihara (The university of Tokyo) for technical assistance. This work was supported by the Cooperative Research Program of “Network Joint Research Center for Materials and Devices” (to YHi) and Grants-in-Aid for Scientific Research (C) and (A) from the Ministry of Education, Culture, Sports, Science and Technology (MEXT) (to YHi and MA, respectively).

**Author Contributions**

Y. Hi. designed the study. Y. Hi. and M.A. wrote the paper. J.K. and S.N. conducted most of the experiments and analyzed the results. T.K. isolated Sll1961 as an interaction partner of thioredoxin in the course of his screening experiment. J.K., S.N., and Y. Ha. conducted HPLC experiment. Y.O. and M.A. conducted structure modeling. Y.N. contributed to redox analysis and discussion. All authors reviewed the results and approved the final version of the manuscript.

**Additional Information**

Supplementary information accompanies this paper at https://doi.org/10.1038/s41598-018-25077-5.

**Competing Interests:** The authors declare no competing interests.

**Publisher’s note:** Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The images or other third party material in this article are included in the article’s Creative Commons license, unless indicated otherwise in a credit line to the material. If material is not included in the article’s Creative Commons license and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this license, visit http://creativecommons.org/licenses/by/4.0/.