

The thioredoxin (Trx) redox-state sensor protein can visualize Trx activities in the light-dark response in chloroplasts

Kazunori Sugiura¹, Yuichi Yokochi^{1,2}, Nae Fu^{1,2}, Yuki Fukaya¹, Keisuke Yoshida^{1,2}, Shoko Mihara^{1,2}, and Toru Hisabori^{1,2*}

Running title: Chloroplast thioredoxin redox sensor protein CROST

¹Laboratory for Chemistry and Life Science, Institute of Innovative Science, Tokyo Institute of Technology, Nagatsuta 4259, Midori-ku, Yokohama 226-8503, Japan

²School of Life Science and Technology, Tokyo Institute of Technology, Nagatsuta 4259, Midori-ku, Yokohama 226-8503, Japan

*To whom correspondence should be addressed: Toru Hisabori, Laboratory for Chemistry and Life Science, Institute of Innovative Research, Tokyo Institute of Technology, Nagatsuta-cho 4259-R1-8, Midori-ku, Yokohama 226-8503, Japan, Tel: +81-45-924-5234, Fax: +81-45-924-5268, Email: thisabor@res.titech.ac.jp

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Thiol-based redox regulation via ferredoxin–thioredoxin (Trx) reductase/Trx controls various functions in chloroplasts in response to light/dark changes. Trx is a key factor of this regulatory system, and five Trx subtypes, including 10 isoforms, have been identified as chloroplast-localized forms in *Arabidopsis thaliana*. These subtypes display distinct target selectivity, and consequently, they form a complicated redox regulation network in chloroplasts. In this study, we developed a FRET-based sensor protein by combining CFP, YFP, and the N-terminal region of CP12, a redox-sensitive regulatory and Trx-targeted

protein in chloroplasts. This sensor protein enabled us to monitor the redox change of chloroplast thioredoxin *in vivo*, and we therefore designated this protein “CROST” (change in redox state of Trx). Using CP12 isoforms, we successfully prepared two types of CROST sensors that displayed different affinities for two major chloroplast Trx isoforms (*f*-type and *m*-type). These sensor proteins helped unravel the real-time redox dynamics of Trx molecules in chloroplasts during the light-dark transition.

To survive under fluctuating environmental conditions, green plants have evolved various

systems for regulating metabolism. The light condition is an important environmental factor for plant cell metabolism. Indeed, important biological functions such as carbon dioxide fixation in plants are adequately regulated under varying light conditions. For this purpose, the thiol-based redox regulation system in chloroplasts is a key regulation system for controlling chloroplast metabolism under fluctuating light conditions. In this regulation system, reducing power produced by water splitting in the light-driven electron transport chain is transferred to thioredoxin (Trx) *via* ferredoxin and ferredoxin–Trx reductase. Trx is a ubiquitous protein possessing a characteristic “WCGPC” motif as an active center. *Via* reduction and oxidation of two Cys residues in the motif and the dithiol-disulfide exchange reaction with target proteins, Trx regulates the activities of target proteins by reducing/oxidizing the disulfide bond on their molecular surfaces (1). Therefore, detection of Trx activity *in vivo* is important for understanding the states of the redox regulation system and various Trx regulated enzymes in chloroplasts such as ATP synthase (2), four enzymes involved the Calvin-Benson cycle (3), the malate valve (4), glucose 6-phosphate dehydrogenase for the oxidative pentose phosphate pathway (4,5), and the enzymes involved in chlorophyll biosynthesis (6).

Genome sequencing of the model plant *Arabidopsis thaliana* revealed that 10 Trx isoforms classified into five subtypes (*f*-, *m*-, *x*-, *y*-, and *z*-types) are localized in

chloroplasts. These five subtypes of Trx exhibit different target selectivities and form a complicate redox network, although these Trx isoforms maintain a common structural motif named Trx-fold (7). The target selectivity of Trx isoforms might be determined by their different molecular characteristics, such as the surface charges of the protein molecule (8) and their midpoint redox potentials (9). However, the details of the target discrimination mechanism of Trx are unclear (10). Recent studies revealed that some proteins possessing Trx-like motifs also function in the redox network in chloroplasts. For example, NADPH-Trx reductase C (NTRC), which has both NTR and Trx domains in one molecule, can reduce several Trx target proteins such as 2-Cys proxiredoxin (2-CysPrx), peroxiredoxin Q, and the Mg-chelatase I subunit using reducing power provided by NADPH (11). In addition, we recently reported the physiological significance of TrxL2, which has a “WCRKC” sequence in its active center motif, in the chloroplast redox network. TrxL2 has higher midpoint redox potential than Trx, and it can oxidize Trx target proteins using oxidation power derived from 2-CysPrx and reactive oxygen species (12). We then clarified that the TrxL2–2-CysPrx pathway is the oxidation pathway for redox-regulated enzymes in chloroplasts in the dark. This important finding reminded us of the significance of the reduction/oxidation balance of the whole redox network in chloroplasts as a metabolism regulatory system for adapting to fluctuating

environmental conditions.

To date, Trx activities *in vivo* have been assessed by visualizing the redox states of target proteins using SDS-PAGE mobility shift analysis following the chemical modification of free thiol groups (13-15). However, this method requires disruption of the desired cells or organelles. To monitor the *in vivo* redox state, roGFP, the most well-known redox sensor protein, was developed by introducing two cysteine residues at positions Ser147 and Gln204 in the wild-type GFP or the GFP S65T mutant, and these Cys residues act as switches to detect the redox state as a redox-dependent change in fluorescence (16). roGFP has been widely used for nondestructive observation in cells. However, roGFP cannot react directly with Trx or small redox molecules such as glutathione and hydrogen peroxide. Instead, roGFP was reduced and oxidized by glutaredoxin (Grx), a Trx superfamily protein containing the “CPYC” motif in its active center (17). Although several redox sensor proteins such as Oba-Q (18), rxYFP (19), rxRFP (20), and Re-Q (21) have been developed, these sensor proteins also cannot detect the redox change in Trx molecules directly. For this purpose, TrxRFP was recently developed as a genetically encoded sensor by fusing rxRFP and human Trx1 (22). This Trx1 was reduced by human Trx reductase and oxidized by human Trx peroxidases in a similar manner as native Trx1. In this system, the redox states of rxRFP and Trx1 in the fused TrxRFP molecule are in

equilibrium, and the redox state of Trx1 can be observed by measuring the change in the fluorescence intensity of rxRFP. However, TrxRFP overexpression in cells might affect the intracellular redox states because TrxRFP functions similar as Trx *in vivo*. In addition, the signal obtained from TrxRFP illustrates the redox status of Trx1 in the fused molecule but not that of other endogenous Trx molecules. Therefore, it is difficult to directly apply TrxRFP to the observation of Trx molecules in photosynthetic organisms.

In this study, we report newly designed sensor probes to detect the redox changes of Trx by combining CFP, YFP, and a partial sequence of CP12 (Fig. 1A). CP12 is a Trx target protein conserved among photosynthetic organisms that forms a super complex with glyceraldehyde-3-phosphate dehydrogenase and phosphoribulokinase and suppress their activities in the dark or under limited light or stress conditions (23). CP12 in chloroplasts has two cysteine pairs that can form disulfide bonds at the N- and C-terminal sides. Trx-*f* can reduce these disulfide bonds and induce a conformational change of CP12 (24). We therefore applied the partial sequence of CP12 for our Trx sensor. We then fused the N-terminal domain of CP12 with mTurquoise (CFP) and cp173-mVenus (YFP). When the fused protein interacted with the reduced form of Trx, the protein exhibited a remarkable change in fluorescence based on Förster resonance energy transfer (FRET). In addition, we succeeded in conferring Trx specificity to the sensor protein using CP12 from a different

origin.

Results and Discussion

Development of probes for redox changes of Trx

CP12 of *A. thaliana* (CP12-2) has two pairs of cysteines (Cys75–Cys84, Cys117–Cys126) that form disulfide bonds under oxidizing conditions, and this protein is considered the target of Trx-*f* *in vivo* (24). Although the molecular structure of CP12 was unsolved, a large conformational change of the molecule caused by changes of the redox status of these cysteine pairs was expected. We therefore applied the partial sequence of the N-terminal domain of CP12-2 from Gly58 to Ala101 including the Cys75–Cys84 pair, which was designated as AtCP12_Nd, to prepare the sensor protein (Fig. 1B, red part). As a donor protein for FRET, mTurquoiseΔ11, a variant of CFP lacking the C-terminal 11 amino acids, was adopted, and cp173-mVenus, a circular permutation mutant of YFP having the amino acid at number 173 as its N-terminus, was used as the acceptor protein (Fig. 1A). These two GFP derivatives were connected by AtCP12_Nd, and the redox changes of cysteines in the CP12 region were then confirmed using the mobility shift on SDS-PAGE following the AMS labeling of free thiols on the proteins (13) (Figure 1C). This new probe nicely displayed high FRET efficiency in the oxidized form, and a remarkable decrease of the signal was noted for the reduced form (Fig. 1D). We then designated this sensor protein CROST1

(change in redox state of thioredoxin #1).

Next, we prepared another sensor protein using CP12 from the cyanobacterium *Anabaena* sp. PCC7120, as the major Trx in cyanobacteria is Trx-*m* and they do not possess Trx-*f*. A part of CP12 from *Anabaena*, namely Lys7–Gln51 (Fig. 1B, red part), was fused to mTurquoiseΔ11 at the N-terminus and cp173-mVenus at the C-terminus. This protein also exhibited a change in fluorescence caused by the redox change of the CP12 molecule (Fig. 1D), and it was designated CROST2.

Characterization of CROST sensors in vitro

We digitalized the FRET signal changes of the sensor proteins by calculating the ratio of fluorescence intensity at 530 and 480 nm (Fig. 2). Initially, reduction of the oxidized form of CROST1 was examined in the presence of various reductants. When Trx-*f*1 from *A. thaliana* was applied together with 100 μM of the reduced form of DTT (DTT_{red}) at time zero, the signal rapidly decreased from 4.1 to 1.5 (Fig. 2A left, closed circle), indicating that the full reduction of CROST1 occurs within 20 s. By contrast, CROST1 was not reduced when the reduced form Trx-*m*2 from *A. thaliana* was applied (Fig. 2A left, open circle). In addition, CROST1 was hardly reduced by 100 μM DTT_{red}, glutaredoxin C5 (GrxC5) from *A. thaliana* (closed square), and 2 mM GSH or by NTRC and 100 μM NADPH (closed reverse triangle). These results clearly indicate that CROST1 is a specific sensor for efficiently detecting the redox status of Trx-*f*.

Similarly, reduction of CROST2 was not observed when DTT_{red}, GrxC5 or NTRC was used as a reductant (Fig. 2A right). However, CROST2 was reduced by Trx-*m2* in addition to Trx-*f1*. This result suggests that CROST2 is a redox sensor which has a lower specificity to Trx isoforms. (Fig. 2A right).

Because we recently revealed a new protein functioning in the chloroplast redox network, which inactivates thiol-regulated enzymes in the dark via oxidation (12), we examined the oxidation process of CROST proteins. As shown in Fig. 2B, oxidation of the reduced form CROST1 was observed when the protein was incubated with Trx-*f1* and the oxidized form of DTT (DTT_{ox}), whereas the newly assigned chloroplast oxidation factors TrxL2 proteins oxidized CROST1 much slower than Trx-*f1* (Fig. 2B left). DTT_{ox}, GrxC5 plus GSSG, or 2-CysPrx together with H₂O₂ did not oxidize CROST1. In contrast, CROST2 was efficiently oxidized by TrxL2.1 and TrxL2.2 (12). These results indicate that CROST proteins can reflect changes in the redox states of Trx-*f* or Trx-*m* in the solution to their FRET signals, whereas the chemical redox molecules do not affect the redox status of CROST proteins.

The Trx isoform specificities of CROST sensors were then examined using nine Trx isoforms derived from *A. thaliana*, *Physcomitrella patens*, and *Anabaena* sp. PCC7120 (*A. 7120*). Because the reaction rate of CROST1 with *At*Trx-*f1* was sufficiently fast as shown in Fig. 2A, a Trx isoform concentration of 30 nM was used together

with 100 μ M DTT_{red} in this study. Among the Trx isoforms examined, *At*Trx-*f1* and *Pp* Trx-*f* could efficiently reduce CROST1 (Fig. 3A, left), and the rate constants were obtained (Fig. 4A). We then examined the Trx isoform specificity of CROST1 in the oxidation process. When 0.1 μ M CROST1 was incubated with 1 μ M Trx isoforms together with 50 mM DTT_{ox}, slow oxidation was observed. Again, *At*Trx-*f1* and *Pp*Trx-*f* could oxidize CROST1 under these conditions (Fig. 3B, left and 4B). Although slow reduction and oxidation of CROST1 were observed when *An*Trx-*m1* was used (Fig. 3A and 3B, left), rate constants were much smaller than those of *f*-type isoforms (Fig. 4A and 4B). These results indicate that CROST1 is an *f*-type Trx-specific sensor that is reduced or oxidized via the redox equilibrium with *f*-type Trx.

To determine the specificity of CROST2, 500 nM Trx isoforms were used because the apparent reduction rates of CROST2 were much slower than those of CROST1 (Fig. 2). Based on the rate constants calculated from the reduction time course data (Fig. 3A, right), CROST2 could detect changes in the redox states of *At*Trx-*f*, *Pp*Trx-*f*, *At*Trx-*m2*, *Pp*Trx-*m*, and *An*Trx-*m1* (Fig. 4C). When the oxidation process was examined, *An*Trx-*m1* was more effective than *At*Trx-*f1* (Fig. 3B, right and 4C), and this must be attributable to the use of *A. 7120* origin CP12 to prepare CROST2. Hence, CROST2 must be applicable to indicate the redox states of *m*-type Trx in cells lacking *f*-type Trx such as cyanobacteria.

Although there was a large difference in the reduction rate constants of CROST1 ($0.40 \text{ s}^{-1} \mu\text{M}^{-1}$) and CROST2 ($0.068 \text{ s}^{-1} \mu\text{M}^{-1}$) using *AtTrx-f1*, the reduction rate constant of CROST2 for *AtTrx-f1* was equivalent to that of chloroplast F_1 -ATPase for spinach-derived *Trx-f* ($0.057 \text{ s}^{-1} \mu\text{M}^{-1}$) (25). Therefore, CROST2 must have a certain reactivity with Trx even under physiological conditions. The rate constant of the reduction process of CROST1 for *AtTrx-f1* was 21-fold larger than that for oxidized process (Fig. 4A and B), suggesting that the oxidation of *AtCP12-2* used by *Trx-f1* to reduce CROST1 does not occur efficiently *in vivo*. Conversely, the rate constant of the reduction process of CROST2 for *AnTrx-m1* was only 4.2-fold larger than that for the oxidation process (Fig. 4C and D), suggesting that the N-terminal domain of *AnCP12* used for CROST2 must be easily oxidized by *AnTrx-m1* in *A. 7120* cells.

We then measured the midpoint redox potentials of CROST1 and CROST2 to reveal the cause of the difference of reduction rate constants of *AtTrx-f1*. Consequently, a midpoint redox potential of -266 mV was obtained for CROST1 at pH 7.5 (Fig. 5A left). This value was considerably larger than that of *AtTrx-f1* (-321 mV) (9) and similar to that of *TrxL2.1*, (-258 mV) (12). By contrast, CROST2 displayed a more negative midpoint redox potential of -296 mV (Fig. 5A). Because the rate of electron transfer between two redox proteins is influenced by the difference of their midpoint redox potentials, the observed difference at approximately 30

mV between CROST1 and CROST2 must be sufficient to explain the difference of the reduction rate constants observed in Fig. 4. The lower midpoint redox potential of CROST2 may explain the difficulty of various Trx isoforms in reducing this protein.

The disadvantages of CROST sensor proteins are their remarkable pH dependencies and small fluorescent signal changes under low pH conditions (Fig. 5B). The observed pH dependencies may be attributable to the pH dependency of the YFP chromophore. Under low pH conditions, the YFP chromophore of cp173-mVenus is not ionized, and therefore, it cannot function as a FRET acceptor. However, CROST sensors fortunately had large dynamic ranges at pH of approximately 7–8 (Fig. 5B), which is equivalent to the pH in various organelles, and the F_{530}/F_{480} values under the oxidized conditions were more than double compared with those under the reduced condition. These sensors should therefore be useful for monitoring the intracellular redox states of Trx molecules.

Visualization of the redox states of Trx in chloroplasts

To observe the redox status of Trx in living cells, we expressed CROST1 and CROST2 in *A. thaliana* chloroplasts and observed the fluorescence of the sensor proteins using an LSM780 confocal fluorescence microscope (Zeiss) (Fig. 6A). Because of the unexpected appearance of bright spots, which were obviously differentiated from the fluorescence signals

from the cytosol or chloroplasts and which might be ascribed to the aggregation of sensor proteins (Fig. 6A white arrow), we have calculated the fluorescent intensities in the image by removing these bright spots. In addition, we have confirmed the expression of CROST proteins in leaves by immunoblotting analysis (Fig. 6B). Degradation of a small part of CROST sensors was observed. The emerged degraded fluorescent proteins may affect the fluorescent intensity ratios of YFP channels (518–544 nm) and CFP (465–491 nm) (F_{YFP}/F_{CFP}) in measurements, especially when free CFP appeared in cells. The auto-fluorescence of chlorophyll is also a possible disturbance factor for the in situ measurement. We therefore obtained fluorescence spectra from *A. thaliana* cells expressing CROST sensors were acquired using the lambda mode and confirmed that effects of degradation of the sensors and the auto-fluorescence of chlorophyll in chloroplasts was enough weak and did not influence the fluorescent signals of CROST sensors (Fig. 6C). The leaves of the wild type plants were used as a control. By changing the light intensity, we could observe a slight decrease of the signal intensity at 530 nm and increase of the second peak at 480 nm (Fig. 6D). F_{YFP}/F_{CFP} were then calculated from these spectra (Fig. 6C). As controls, F_{YFP}/F_{CFP} ratios were determined in the presence of 20 mM DTTred or 20 μ M 3-(3,4-dichlorophenyl)-1, 1-dimethylurea (DCMU). Finally, we observed the change in F_{YFP}/F_{CFP} during the mutual light/dark changes using CROST1 (Fig. 6E, upper panel, closed circle)

and CROST2 (Fig. 6E, lower panel, closed circle) introduced into the chloroplasts of *A. thaliana*. When the plant leaves were illuminated, both CROST sensors immediately shifted to the reduced state, implying that Trx proteins in chloroplasts, mainly Trx-*f*, were reduced. When the light source was removed, F_{YFP}/F_{CFP} gradually increased, indicating that Trx-*f* was slowly oxidized in chloroplasts. The signal decay of CROST1 was obviously slower than that of CROST2. This difference might be ascribed to the difference of the midpoint redox potentials of these two sensors. CROST1 had higher redox potential than CROST2 (Fig. 5A). In addition, the reactivity of CROST2 was not restricted to *f*-type Trx, but it was sufficient against Trx proteins and TrxL2. This property might explain the rapid oxidation of CROST2 even under the light-to-dark transition (Fig. 2, Fig. 3). In contrast, we could not detect any changes of F_{YFP}/F_{CFP} when 20 μ M DCMU was applied (Fig. 6E, open circle). Thus, we succeeded in observing the real-time redox dynamics of Trx molecules in chloroplasts caused by photosynthetic electron transport according to the light-dark transition.

Experimental Procedure

Gene construction

The cDNAs of mTurquoise Δ 11, cp173m-Venus, N-terminal domain of AtCP12-2 (At3g56690) and AnCP12 (pfam02672) were amplified by PCR and fused using hot fusion methods (26) (Fig. 1B) with the pET-23a vector to express the protein

with a C-terminal His₆-tag. cDNA fragments encoding the mature protein-coding regions, which were predicted by TargetP (27), of *AtTrx-f1* (At3g02730), *AtTrx-m2* (At4g03520), *AtTrx-x* (At1g50320), *AtTrx-y2* (At1g43560), *AtTrx-z* (At3g06730), *PpTrx-f* (PHPAT.004G014000.1), and *PpTrx-m* (PHPAT.005G085400.1) were cloned into the pET-23a vector (Novagen) to express the protein with a C-terminal His₆-tag. Other expression plasmids were prepared in our earlier studies (9,28,29).

Protein expression and purification

Escherichia coli strains BL21(DE3) and BL21(DE3) pLysS were transformed with each expression plasmid. BL21(DE3) pLysS was used to express *A. thaliana*-derived Trx, and BL21(DE3) was used for *P. patens*-derived Trx and CROST sensors. Transformed *E. coli* was then cultured at 37°C. Expression of the desired protein was induced using 0.5 mM isopropyl-β-D-thiogalactopyranoside followed by further cultivation for 16 h at 21°C. Collected cells were disrupted and ultracentrifuged (125,000 × *g* for 40 min) to obtain the supernatant containing the desired protein for subsequent purification. His-tagged *A. thaliana*-derived Trxs and *P. patens*-derived Trx and CROST sensors were purified via Ni-NTA affinity chromatography using Ni-NTA agarose (QIAGEN) resin. Ni-NTA agarose resin was washed in 25 mM Tris-HCl (pH 8.0) containing 20 mM imidazole and eluted using the same buffer containing 250 mM imidazole. Elutions including *A.*

thaliana-derived Trxs were applied to Superdex 75 Increase 10/300 GL (GE Healthcare) columns and purified. Elutions including *P. patens*-derived Trx were dialyzed and concentrated. Elutions containing CROST sensors were applied to a TOYOPEARL Butyl-650 (TOSOH, Tokyo) column and eluted using 20 mM Tris-HCl (pH 8.0) with an ammonium sulfate reverse-gradient from 30 to 0%. Peak fractions eluted from the column were collected. The elution containing CROST sensors were incubated in the presence of 0.5 mM diamide for 10 min at 25 °C, dialyzed and concentrated. Other proteins, which were partially purified by Ni-NTA agarose chromatography, were then purified according to the methods described in our earlier studies (9,28,29). Purities of the recombinant proteins used were examined by SDS-PAGE (Figure. S1).

Fluorescence spectroscopy and calculation of the reduced and oxidized ratio

Fluorescence emission spectra were measured using an FP-8500 fluorescence spectrophotometer (JASCO, Tokyo). An excitation wavelength of 430 nm was used to excite CROST sensors. Emission fluorescence intensities from CROST sensors at 480 (F₄₈₀) and 530 nm (F₅₃₀) were measured, and the fluorescence ratio (Fr) was calculated using equation 1.

$$\text{Fr} = F_{530}/F_{480} \quad \text{equation 1}$$

To establish fully reduced or oxidized

conditions, CROST sensors were incubated with 100 μM DTT_{red} or 50 mM DTT_{ox} in the presence of 1 μM *AtTrx-f1* for 10 min at 25°C, and Fr values were obtained as F_{red} and F_{ox}. The oxidation ratio, Ox/total, was then calculated using equation 2.

$$\text{Ox/total} = (\text{Fr} - \text{F}_{\text{red}}) / (\text{F}_{\text{ox}} - \text{F}_{\text{red}})$$

equation 2

Determination of rate constants

To eliminate the influence of direct reduction by DTT, we calculated the rate constant from Ox/total data in the presence of Trx isoforms after subtracting Ox/total data in the presence of only DTT at each time. Then, resulting Ox/total value at the indicated time was plotted and fitted using equation 3 to determine the rate constant k_{red} and equation 4 to determine k_{ox} .

$$\text{Ox/total} = \exp(-k_{\text{red}} [\text{Trx}] t)$$

equation 3

$$\text{Ox/total} = 1 - \exp(-k_{\text{ox}} [\text{Trx}] t)$$

equation 4

[Trx] is the concentration of Trx.

Determination of the midpoint redox potentials of CROST sensors

CROST sensor proteins (0.1 μM) were incubated in 25 mM Tris-HCl (pH 7.5), 50 mM DTT_{ox}, various concentrations of DTT_{red} (1 μM –50 mM), and 1 μM *AtTrx-f1* for 3 h at 25°C. The midpoint redox potential values of CROST sensors were calculated by fitting the titration data of the reduction level of proteins to the Nernst equation. A value of –357 mV was used as a midpoint redox

potential of DTT at pH 7.5 (9).

Expression and observation in chloroplasts of *A. thaliana*

The expression plasmids pRI201/PtCROST1 and pRI201/PtCROST2 were prepared by appending the recA DNA recombination family protein leader sequence to the 5' end of the CROST1 and CROST2 coding genes cloned in the expression vector pRI201-AN (Takara). The *A. thaliana* plants were transformed with the expression plasmids using *Agrobacterium tumefaciens* and then grown at 22°C under 16 h light/8 h dark conditions. For the observation, AxioObserverZ1 with LSM780 (Carl Zeiss) was used with an Ar 458 nm laser (25 mW). The laser power was held at less than 0.1% to avoid activating the photosystem. A Mulch channel GaAsP detector was used in the photon-counting and lambda modes. The intensity of the fluorescence image was then digitalized using ImageJ image analysis software.

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Abbreviations used are: Trx, thioredoxin;

CROST, change in redox state of Trx; NTRC, NADPH-Trx reductase C; 2-CysPrx, 2-Cys proxiredoxin; Grx, glutaredoxin; CFP, mTurquoise; YFP, cp173-mVenus; FRET, Förster resonance energy transfer; CP12-2, CP12 of *A. thaliana*; GrxC5, glutaredoxin C5; DTT_{ox}, the oxidized form of DTT; DTT_{red}, the reduced form of DTT.

Conflict of interest

The authors declare that they have no conflicts

of interest related to the contents of this article.

Author contributions

KS and TH conceived the study. KS, YY and NF performed the experiments. TH and KY supervised the research. YF and SM assisted with the experiments. KS, KY, and TH discussed the data. KS and TH wrote the paper, and SM and KY commented on the manuscript.

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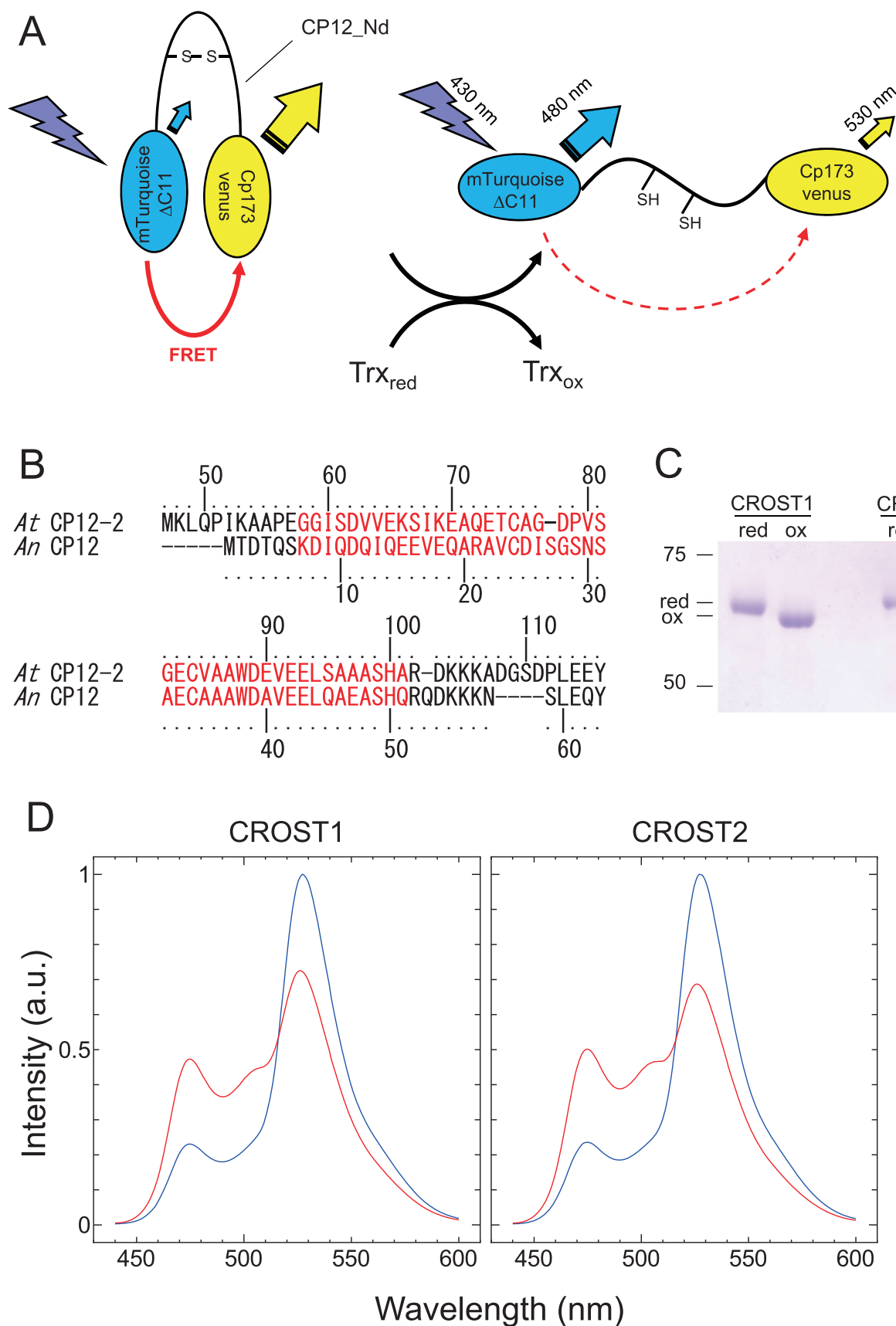


Figure 1. The molecular design of CROST sensors. *A*, schematics of CROST sensors. Mutants of CFP (mTurquoise featuring a deletion of 11 C-terminal amino acids) and YFP (cp173Venus) were connected by the N-terminal domain of CP12 derived from *A. thaliana* or *Anabaena* sp. PCC7120. *B*, amino acid sequences of CP12. N-terminal domains used for CROST sensors are highlighted in red. *C*, oxidation and reduction of CROST proteins were confirmed by non-reducing SDS-PAGE following the AMS labeling. *D*, redox-dependent fluorescence spectral changes of CROST proteins. Fluorescence emission spectra of oxidized (blue line) and reduced (red line) CROST proteins at 25° C in 50 mM Tris-HCl (pH 7.5), 50 mM NaCl, and 1 mM EDTA were measured with excitation at 430 nm. CROST proteins were oxidized during the purification process and therefore reduced in the presence of 100 mM DTT_{red} and 0.2 mM *At*Trx-f1 before the measurement.

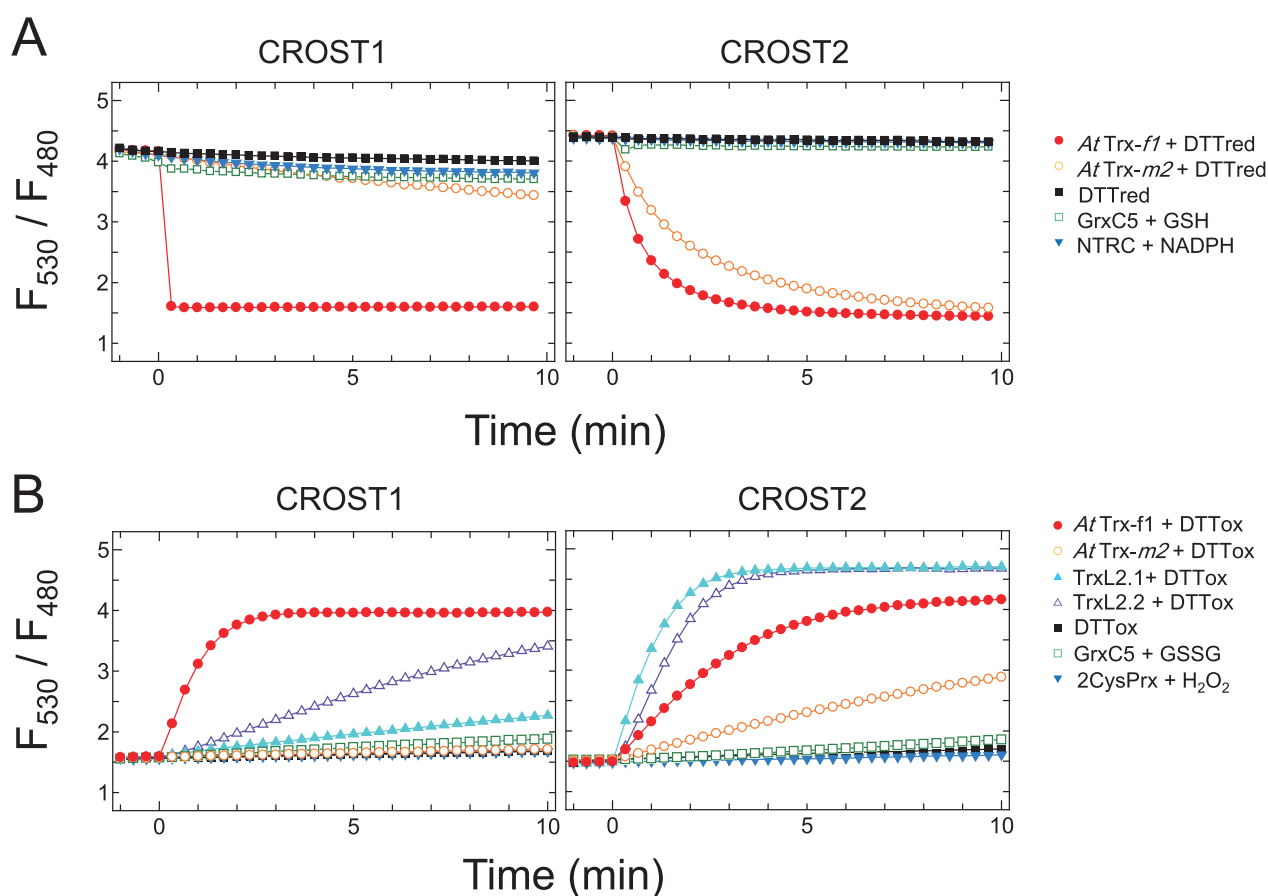


Figure 2. Reduction and oxidation of CROST proteins. *A*, reduction of CROST1 (left) and CROST2 (right) by reductants. Fluorescent intensity ratios (530/480 nm) of 0.1 mM oxidized CROST proteins at 25° C in 50 mM Tris-HCl (pH 7.5), 50 mM NaCl, and 1 mM EDTA were plotted before and after the addition of reductants at time 0 as follows: (closed circle) 0.2 mM *At*Trx-*f1* + 100 mM DTT_{red}, (open circle) 0.2 mM *At*Trx-*m2* + 100 mM DTT_{red}, (closed square) 100 mM DTT_{red}, (open square) 0.2 mM *At*GrxC5 + 2 mM GSH, and (closed inverted triangle) 0.2 mM *At*NTRC + 100 mM NADPH. *B*, oxidation of CROST1 (left) and CROST2 (right) by oxidants. Fluorescence intensity ratios of 0.1 mM reduced CROST proteins were plotted before and after the addition of reductants at time 0 as follows: (closed circle) 0.2 mM *At*Trx-*f1* + 50 mM DTT_{ox}, (closed triangle) 0.2 mM *At*Trx-*m2* + 50 mM DTT_{ox}, (closed triangle) 1 mM *At*TrxL2.1 + 50 mM DTT_{ox}, (open triangle) 1 mM *At*TrxL2.2 + 50 mM DTT_{ox}, (closed square) 50 mM DTT_{ox}, (open square) 0.2 mM *At*GrxC5 + 1 mM GSSG, and (closed inverted triangle) 0.2 mM *At*2CysPrx + 0.1 mM H₂O₂. In the presence of 0.2 mM *At*Trx-*f1* and 0.5 mM DTT_{red}, 10 mM CROST sensors were reduced for 30 min at 25° C before measurements and diluted to a using concentration.

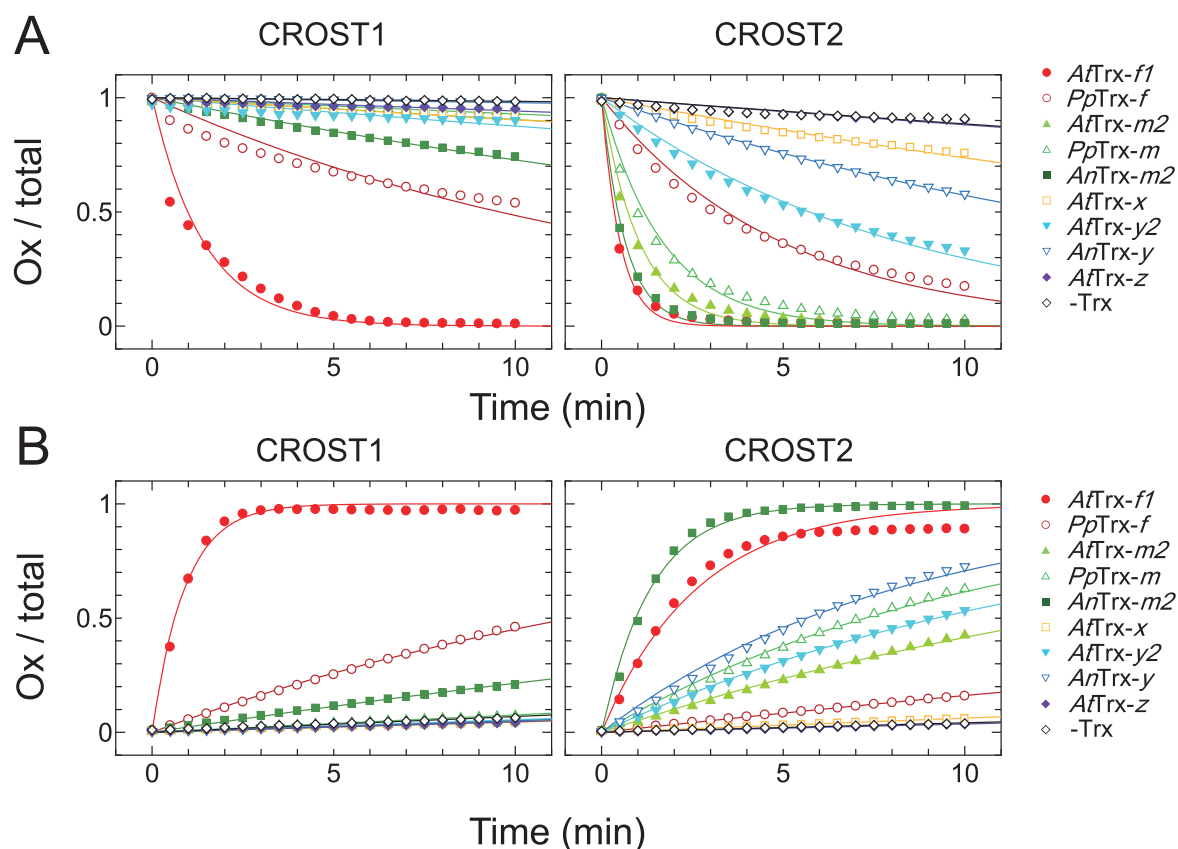


Figure 3. Reduction and oxidation of CROST proteins by five Trx subtypes. *A*, reduction levels of CROST1 (left) and CROST2 (right) at 25° C in 50 mM Tris-HCl (pH 7.5), 50 mM NaCl, and 1 mM EDTA were plotted. At time 0, each Trx subtype was added at a final concentration of 30 nM for CROST1 and 500 nM for CROST2 with 100 mM DTT_{red}: (closed circle) *AtTrx-f1*, (open circle) *PpTrx-f*, (closed triangle) *AtTrx-m2*, (open triangle) *PpTrx-m*, (closed square) *AnTrx-m1*, (open square) *AtTrx-x*, (closed inverted triangle) *AtTrx-y2*, (open inverted triangle) *AnTrx-y*, (closed diamond) *AtTrx-z*, and (open diamond) without Trx. *B*, oxidation levels of CROST1 (left) and CROST2 (right) were plotted under the conditions described for (*A*). Trx (1 mM) was used in the presence of 50 mM DTT_{ox}. The symbols follow those used in (*A*). CROST sensors (10 mM) were reduced in the presence of 0.2 mM *AtTrx-f1* and 0.5 mM DTT red for 30 min at 25° C before measurements and diluted to a using concentration.

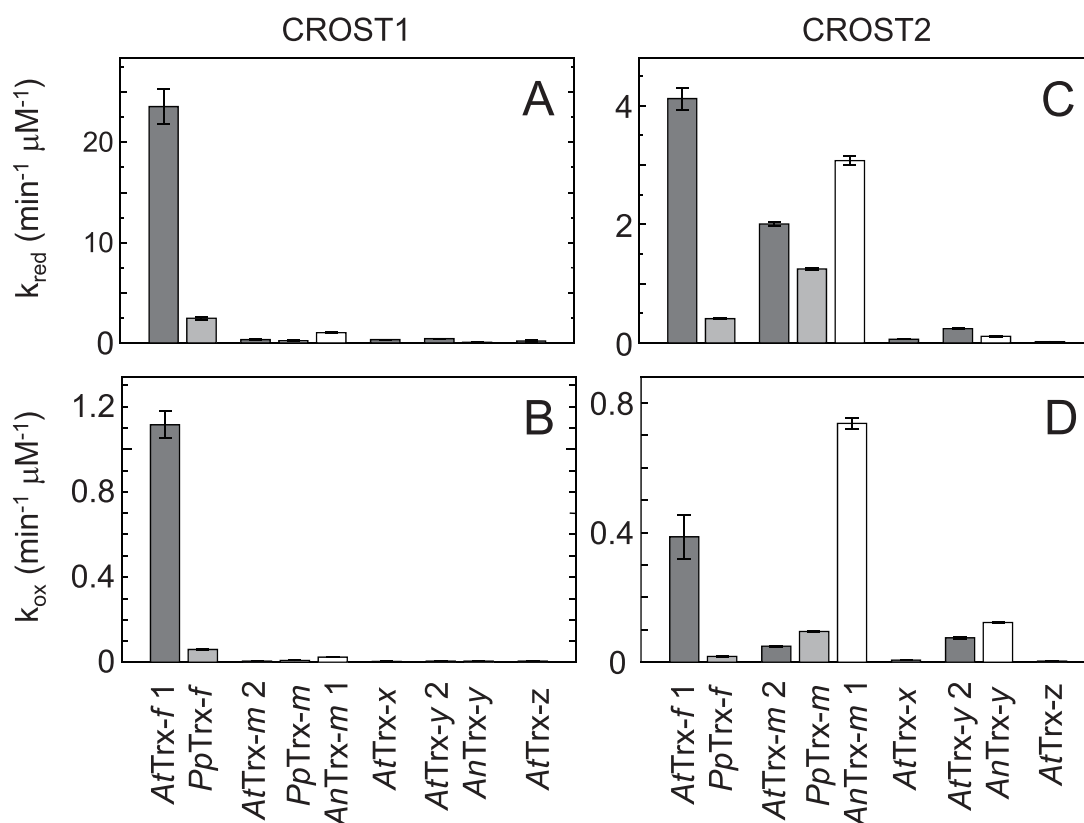


Figure 4. Rate constants of CROST reduction and oxidation

A and *B*, reaction rate constants (k_{red}) were determined via fitting with the first-order equation (see Experimental Procedures) of (*A*) the reduction process data for CROST1 shown in Figure 3A, left panel, and (*B*) the reduction process data for CROST2 shown in Figure 3A, right panel. *C* and *D*, oxidation rate constants (k_{ox}) were determined via fitting with the first-order equation (see Experimental Procedures) of (*C*) the oxidation process data for CROST1 shown in Figure 3B, left panel, and (*D*) the oxidation process data for CROST2 shown in Figure 3B, right panel.

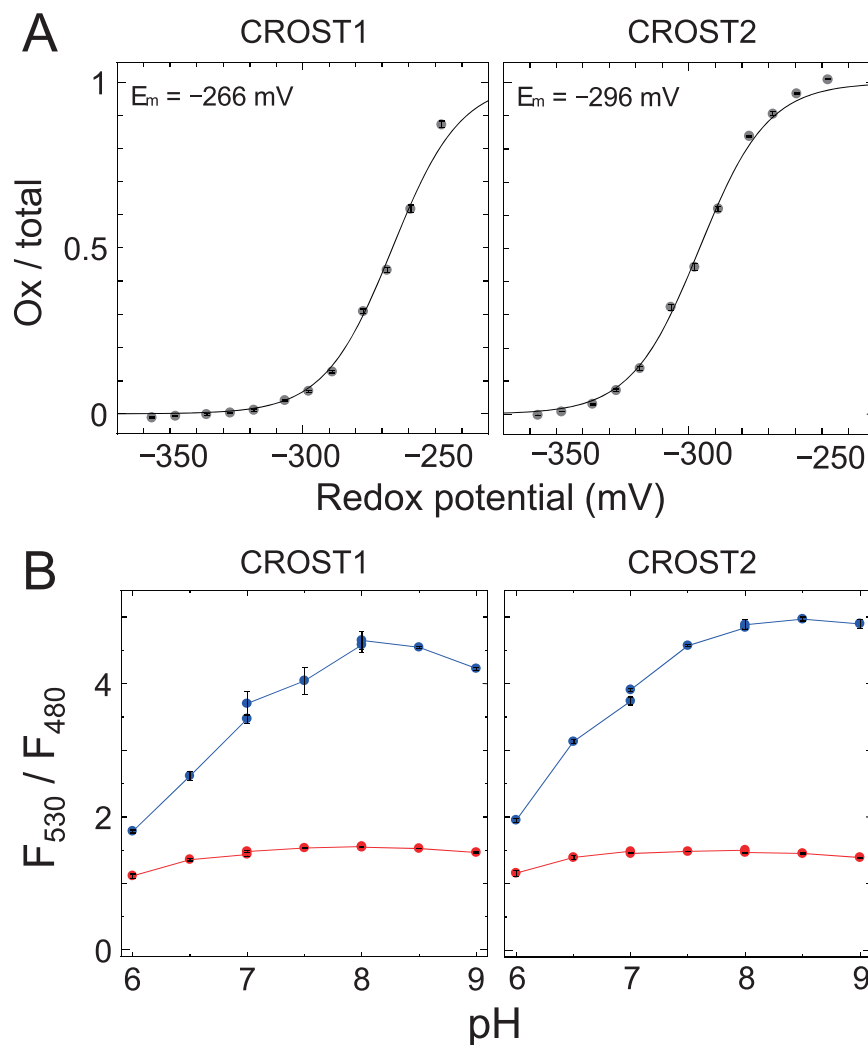
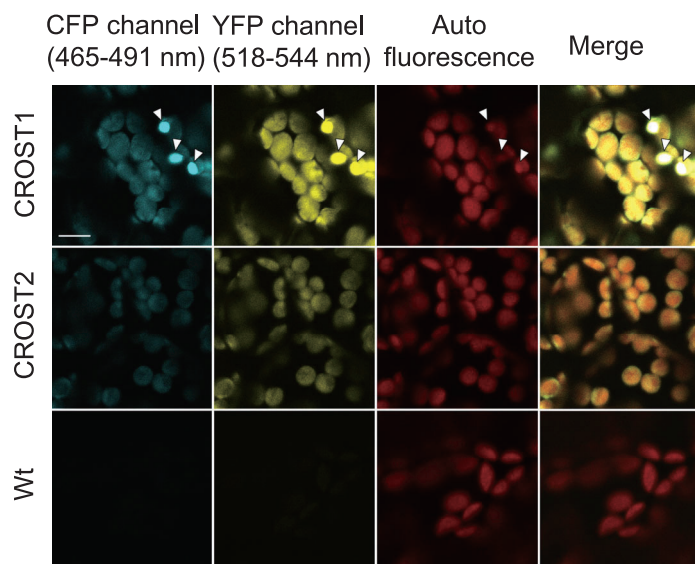
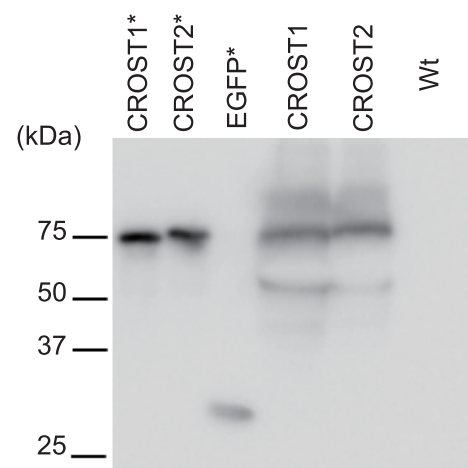


Figure 5. Redox potential and pH dependency of CROST sensors. *A*, the reduction levels of CROST1 (left) and CROST2 (right) were quantified as the ratios of the reduced forms to the total and plotted against the redox potential of DTT buffer (pH 7.5). Data were fitted to the Nernst equation for two electrons exchanged ($y = 1/(1 + \exp(0.078(x - E_m)))$). The midpoint redox potential value shown in the figure is the mean $\pm$ SD (four independent experiments using two different protein preparations). *B*, fluorescence ratio (F_{530}/F_{480}) of recombinant CROST1 (left) and CROST2 (right) in different pHs were measured for their oxidized (blue line) and reduced (red line) forms. Buffers MES (for pH 6–7), PIPES (for pH 7–8), and HEPES (for pH 8–9) were used.

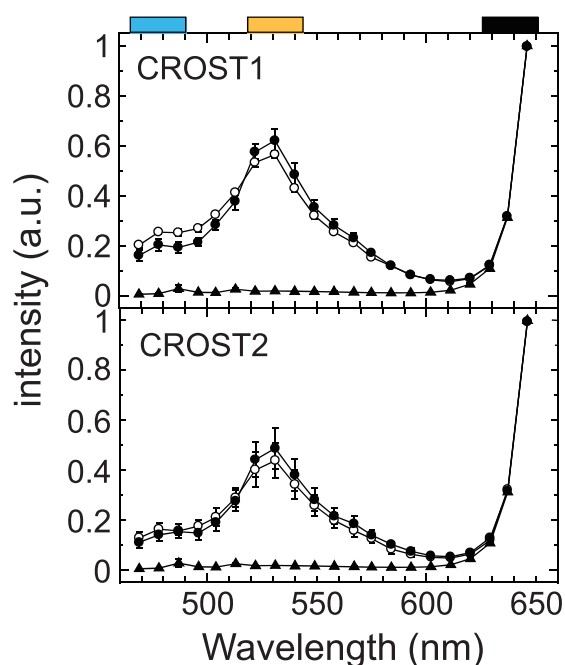
A



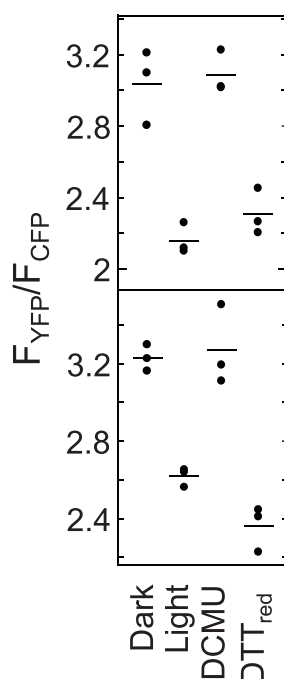
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C



D



E

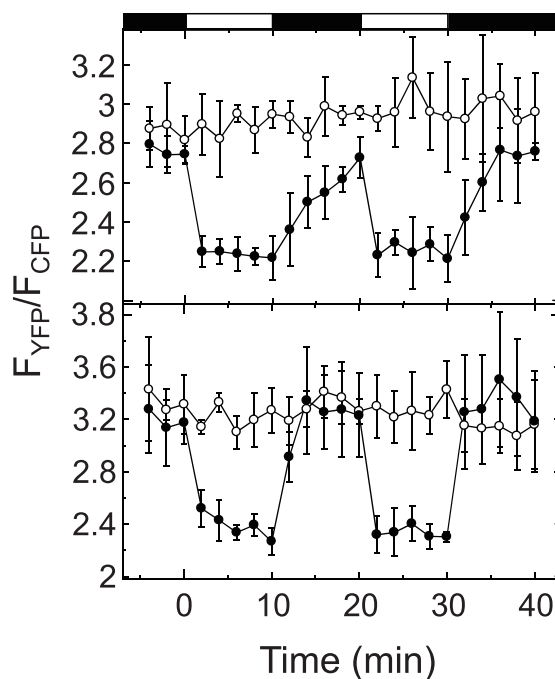


Figure 6. Visualization of intracellular Trx activities. *A*, fluorescence images of transgenic plant leaves expressing CROST1 and CROST2 and those of wild-type leaves were obtained using an Ar 458 nm laser for excitation. The images from 465 to 491 nm for CFP and from 518 to 544 nm for YFP were observed. Chlorophyll auto-fluorescence from 624 to 651 nm was also assessed, and merged images were created (scale bar; 10 mm). *B*, immunoblotting analysis of the expression of CROST1 and CROST2 in leaves using antibody against GFP. Recombinant CROST1, CROST2 and EGFP were used as controls (asterisk). *C*, emission spectra of CROST1 (*upper panel*) and CROST2 (*lower panel*) expressed in chloroplasts were measured using the confocal microscopy under dark (closed cycle) and light (50 mmol photons $\text{m}^{-2} \text{s}^{-1}$, open cycle) conditions. Emission spectra of the wild type *A.thaliana* cells were measured as a control (closed triangle). The obtained spectra were standardized at the chlorophyll auto-fluorescence region (black box) using Least-square method. Cyan and yellow boxes show the region used for the observation of CFP and YFP, respectively. *D*, fluorescent intensity ratios of CROST1 (*upper panel*) and CROST2 (*lower panel*) in the dark and under the light (50 mmol photons $\text{m}^{-2} \text{s}^{-1}$) condition were calculated and plotted. Fluorescent intensity ratios in the presence of 20 mM DTT_{red} and 20 mM DCMU were determined as well. The results of three independent experiments were shown. Bars show the average. *E*, fluorescent intensity ratio changes of CROST1 (*upper panel*) and CROST2 (*lower panel*) under mutual light (50 mmol photons $\text{m}^{-2} \text{s}^{-1}$)/dark conditions in the absence (closed cycle) and presence (open cycle) of 20 mM DCMU were calculated at every 2 min and plotted. The light period is indicated by white rectangles above the graph. The results obtained from three independent experiments were shown with S.D bars.

The thioredoxin (Trx) redox-state sensor protein can visualize Trx activities in the light-dark response in chloroplasts

Kazunori Sugiura, Yuichi Yokochi, Nae Fu, Yuki Fukaya, Keisuke Yoshida, Shoko Mihara and Toru Hisabori

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