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Data Article

Data supporting the absence of FNR dynamic photosynthetic membrane recruitment in *trol* mutants



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ABSTRACT

In photosynthesis, the flavoenzyme ferredoxin:NADP⁺ oxidoreductase (FNR) catalyses the final electron transfer from ferredoxin to NADP⁺, which is considered as the main pathway of highenergy electron partitioning in chloroplasts (DOI: http://www.dx. doi.org/10.1111/j.1365-313X.2009.03999.x [1], DOI: http://www.dx. doi.org/10.1038/srep10085 [2]). Different detergents and pH treatments of photosynthetic membranes isolated from the Arabidopsis wild-type (WT) and the loss-of-function mutants of the thylakoid rhodanase-like protein TROL (trol), pre-acclimated to either dark, growth-light, or high-light conditions, were used to probe the strength of FNR-membrane associations. Detergents β -DM (decyl- β -D-maltopyranoside) or β -DDM (n-dodecyl- β -D-maltopyranoside) were used to test the stability of FNR binding to the thylakoid membranes, and to assess different membrane domains containing FNR. Further, the extraction conditions mimicked pH status of chloroplast stroma during changing light regimes. Plants without TROL are incapable of the dynamic FNR recruitment to the photosynthetic membranes.

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Subject area	Biology, Biochemistry
More specific sub- ject area	Protein interactions
Type of data	Western blots
How data was acquired	SDS-PAGE, Western transfer, ECL
Data format	Raw and analysed
Experimental	Arabidopsis thaliana (L.) ecotype Columbia (Col-0, WT) plants and At4g01050
factors	knock-out mutant line, trol, were grown either in dark or under 80 μ mol photons $m^{-2} s^{-1}$ (GL) or 250 μ mol photons $m^{-2} s^{-1}$ (HL), respectively. Intact Arabidopsis chloroplasts were isolated from 3–4 week old plants and analysed by using SDS-PAGE for TROL-FNR complex stability.
Experimental features	For TROL-FNR complex dynamics investigation, thylakoids were isolated from intact Arabidopsis chloroplasts, separated by ultracentrifugation into insoluble and soluble fractions, treated by nonionic detergents at different pH, analysed by SDS-PAGE, Western transfer, immunodecorated with α -FNR antibody, and finally detected by semiquantitative ECL.
Data source location	Zagreb, Croatia
Data accessibility	Data is supplied in this article

Specifications Table

Value of the data

- Data assess alternative membrane binding and release of chloroplast FNR, or its association with different membrane complexes.
- Recruitment of FNR to thylakoids was tested on WT or *trol* Arabidopsis plants pre-acclimated to different light conditions.
- Biological function of FNR-membrane association in the context of photosynthetic electron flow regulation was addressed.

1. Data

The data indicate that the absence of TROL protein influences the dynamic membrane association properties of chloroplast FNR. Thylakoids were isolated from plants acclimated to different light regimes by using buffers of different pH and containing either nonionic detergents decyl- β -D-maltopyranoside (β -DM) (Fig. 1b), or n-dodecyl- β -D-maltopyranoside (β -DDM) (Fig. 1c), or no detergent (Fig. 1a). The inclusion of β -DM or β -DDM was used to probe the stability of FNR binding to thylakoid supramolecular complexes and to assess different membrane domains containing FNR. The dynamism of TROL-FNR interaction was evaluated by quantifying FNR distribution between the membrane and the soluble fractions.

2. Experimental design, materials and methods

2.1. Plant material and growth conditions

Arabidopsis thaliana (L.) ecotype Columbia (Col-0) plants and *At4g01050* knock-out mutant line, *trol* [1], were grown on potting substrate (Stender, Germany) in the growth chamber (Kambič, Slovenia).

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