

S2 Table. Electron transport activities of photosystem I and II in thylakoid membranes isolated from the WT and *phyB-1* seedlings grown under Rc irradiation for 9 days.

| Photosystem II activity ($\mu\text{mol e}^- \text{ mg Chl}^{-1} \text{ h}^{-1}$) | | Photosystem I activity ($\mu\text{mol e}^- \text{ mg Chl}^{-1} \text{ h}^{-1}$) | |
|---|------------------|--|-----------------|
| WT | <i>phyB-1</i> | WT | <i>phyB-1</i> |
| 487.3 \pm 51.1 | 461.9 \pm 29.4 | 279.2 \pm 14.3 | 242.4 \pm 7.6 |

Each value is the mean of data from 4 samples with their SD.

Procedures

Isolation of thylakoid membranes

Seedlings grown under Rc irradiation for 9 days were harvested, weighed and ground in 8-fold volume of 50 mM Tris HCl buffer (pH 7.5) containing 0.1 M sucrose, 10 mM NaCl, 2.5 mM MgCl₂ and 10 mM sodium ascorbate by a Waring blender. The homogenate was filtered through two layers of cheesecloth and centrifuged at 300 $\times g$ for 30 sec. The supernatant was further centrifuged at 20,000 $\times g$ for 7 min. The precipitate was resuspended with the same buffer and then centrifuged at 20,000 $\times g$ for 7 min. Crude thylakoid was obtained as the resultant precipitate, which was suspended into small volume of a measuring buffer, 50 mM Mes-NaOH (pH 6.5) containing 0.1M sucrose, 10 mM NaCl and 5 mM MgCl₂.

Measurement of PSII activity

Photosynthetic electron transport activity of PSII was estimated from photosynthetic oxygen evolution rate measured by a Clark-type oxygen electrode (Rank Brothers Ltd). Ten μg chlorophyll ml^{-1} of thylakoid in the measuring buffer in a glass chamber with the electrode was irradiated by saturating actinic light from a halogen lamp in the presence of 0.8 mM 2,6-dichlorobenzoquinone (DCBQ) and 1 mM potassium ferricyanide.

Measurement of PSI activity

Photosynthetic electron transport activity of PSI was estimated from decrease of oxygen molecules by re-oxidation of photosynthetically reduced methyl viologen (MV) measured by a Clark-type oxygen electrode (Rank Brothers Ltd). Four μg chlorophyll ml^{-1} of thylakoid in the measuring buffer was exposed to a saturating actinic light in the electrode chamber in the presence of 1 mM KCN, 0.07 mM reduced 2,6-dichloroindophenol (DCIP), 0.1 mM MV, 0.01 mM 3-(3,4-Dichlorophenyl)-1,1-dimethylurea (DCMU) and 0.01 mM carbonylcyanide-3-chlorophenylhydrazone (CCCP).