Thylakoid protein phosphorylation in evolutionally divergent species with oxygenic photosynthesis

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Abstract Phosphothreonine antibody was used to explore reversible thylakoid protein phosphorylation in vivo in evolutionally divergent organisms with oxygenic photosynthesis. Three distinct groups of organisms were found. Cyanobacteria and red algae, both with phycobilisome antenna system, did not show phosphorylation of any of the photosystem II (PSII) proteins and belong to group 1. Group 2 species, consisting of a moss, a liverwort and a fern, phosphorylated both the light-harvesting chlorophyll \textit{ahl} proteins (LHCII) and the PSII core proteins D2 and CP43, but not the D1 protein. Reversible phosphorylation of the D1 protein seems to be the latest event in the evolution of PSII protein phosphorylation and was found only in seed plants, in group 3 species. Light-intensity-dependent regulation of LHCII protein phosphorylation was similar in group 2 and 3 species, with maximal phosphorylation of LHCII at low light and nearly complete dephosphorylation at high light.

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Key words: Photosystem II; Light-harvesting chlorophyll \textit{ahl} protein; Thylakoid protein phosphorylation; Lower plant; Cyanobacterium

1. Introduction

Several thylakoid proteins of higher plants undergo reversible light-dependent phosphorylation [1,2]. These phosphorylated proteins mainly belong to the photosystem II (PSII) complex, including the heterodimeric reaction center polypeptides D1 and D2, the internal chlorophyll \textit{a} antenna polypeptide CP43 and the 9 kDa \textit{psbH} gene product as well as the two major polypeptides of the light-harvesting chlorophyll \textit{ahl} binding complex (LHCII). Reversible phosphorylation of all these proteins occurs at the N-terminal threonine residue on the stromal side of the thylakoid membrane [3,4]. Additionally, a chill-induced light-dependent phosphorylation of a minor chlorophyll \textit{ahl} binding protein CP29 has been shown to occur in maize leaves [5]. Reversible phosphorylation of PSII proteins in lower photosynthetic organisms is less well characterized. Phosphorylation of D1, D2 or CP43 in red algae or cyanobacteria has not been reported (for reviews, see [6,7]), yet the phenomenon has been supposed to be involved in regulatory processes of PSII function [6,8]. Chlorophyll \textit{b}-possessing lower eukaryotic organisms, like green algae and mosses, also appear to differ from higher plants with respect to PSII core protein phosphorylation [9,10]. On the other hand, LHCII polypeptide phosphorylation has been established also for green algae [11–13].

Reversible LHCII polypeptide phosphorylation has been implicated in the regulation of excitation energy distribution between the two photosystems [6,14] while the physiological significance of PSII core protein phosphorylation is not yet fully resolved. Reversible D1 protein phosphorylation of higher plants has been shown to play an important role in the regulation of D1 protein turnover and hence the repair of PSII centers under photoinhibitory conditions [15,16]. PSII repair cycle also involves a conversion of PSII dimers to monomers [17]. The stability of PSII dimers is possibly controlled by reversible phosphorylation of PSII core proteins thus providing another level of regulation for D1 degradation [18]. Thylakoid protein phosphorylation, however, may not only be involved in a short-term regulation of photosynthetic processes but possibly also has a crucial role in physiological and metabolic signalling within chloroplasts or even in the cross-talk between the chloroplast and the nuclear genomes.

Recent application of the phosphothreonine antibody for studies of thylakoid protein phosphorylation has made it possible to investigate the dynamics of thylakoid protein phosphorylation in intact plants [19]. In the present study we have addressed a question on phosphorylation of PSII proteins and its response to the irradiance level in evolutionally divergent oxygenic photosynthetic organisms, which has been a matter of confusion in attempts to reveal the physiological role of PSII protein phosphorylation.

2. Materials and methods

\textit{Synechocystis} sp. PCC 6803 wild type cells were grown in BG-11 medium under continuous light of 50 \textmu mols photons m$^{-2}$ s$^{-1}$ at 20°C. \textit{Marchantia polymorpha}, a liverwort, and \textit{Adiantum tenerum}, a fern, were grown in a greenhouse under low irradiance (30 \textmu mols photons m$^{-2}$ s$^{-1}$) in a 12/12 h light rhythm. Pumpkin (\textit{Cucurbita pepo} \textit{L.}) and winter rye (\textit{Secale cereale} \textit{L.}) were grown at 300 \textmu mols photons m$^{-2}$ s$^{-1}$ in a 16/8 h light rhythm at 20°C for pumpkin and at 20°C/13°C for rye. \textit{Ceramium tenuicorne}, a red alga, was collected from its natural habitat (the coast of the Baltic Sea in southwestern Finland) and exposed to different light treatments during the same day.

For analysis of the steady-state level of thylakoid protein phosphorylation, the photosynthesizing tissue of different species was harvested in the end of the 12-h dark period, after 4 h in growth light conditions and after a high-light treatment of intact plants. Rye plants were additionally treated at several intermediary light intensities. High-light treatment of the higher plants was given under a metalhalide lamp (HQI-T 250 W/daylight) at a PFD of 2000 \textmu mols photons m$^{-2}$ s$^{-1}$ for 4 h. Less severe light treatments were given for low-light grown organisms; \textit{Synechocystis}, \textit{Ceratodon} and \textit{Adiantum} were illuminated for 4 h and \textit{Ceramium} and \textit{Marchantia} for 1 h at a PFD of...
500 μmol photons m⁻² s⁻¹. After different light treatments, samples were immediately frozen in liquid nitrogen and stored at −80°C.

Thylakoid membranes of Synechocystis were isolated according to Tyyşjärvi et al. [21] in the presence of 10 mM NaF to inhibit thylakoid protein dephosphorylation. For other species the thylakoids were rapidly isolated by homogenizing the tissue in ice-cold isolation buffer (350 mM sucrose, 25 mM Tris-HCl, pH 8.5, 5 mM Mg-acetate, 5 mM K-acetate, 10 mM NaF). After filtration through Miracloth, thylakoids were collected by centrifugation at 6000 g, washed in 25 mM Tris-HCl, pH 8.5, 10 mM Mg-acetate, 100 mM K-acetate, 10 mM NaF and finally suspended in the isolation buffer and stored at −80°C. The chlorophyll content of plants with LHCII antenna was determined according to Porra et al. [22] while the method of Arnon [23] was used to determine chlorophyll a content in Synechocystis and Ceramium, possessing a phycobilisome antenna as a light-harvesting system for PSII.

Thylakoid polypeptides were separated by SDS-PAGE essentially according to Laemmli et al. [24], using 15% acrylamide and 6 M urea in the separation gel. Thylakoid samples equivalent to 2.5 μg chlorophyll for Ceratodon, Marchantia and Adiantum, 1.5 μg of chlorophyll for pumpkin and rye were loaded in the wells. Polypeptides were electrophoresed to a PVDF membrane (Immobilon P, Millipore). After blocking with 1% BSA (fatty acid free, Sigma), thylakoid phosphoproteins were immunodetected with a commercial polyclonal phosphothreonine antibody (Zymed Laboratories Inc.) using an Immuno-Lite Assay Kit (BioRad). D1, D2, CP43, CP29 and LHCII were immunodetected with protein-specific antibodies. The D1-specific antibody was raised against the amino acids 234–242 of the D-E loop of D1 protein and the D2-specific antibody against the amino acids 230–245 of the D-E loop of D2 protein in Synechocystis 6803 (Research Genetics, Huntsville, AL). Protein-specific antibodies against CP43, CP29 and LHCII were generous gifts from Dr. Roberto Barbato, Dr. David Simpson and Dr. Stefan Jansson, respectively.

3. Results

3.1. Reversible phosphorylation of thylakoid proteins in angiosperms

Pumpkin and rye responded to changing light conditions with clear modifications in the steady-state phosphorylation level of the thylakoid proteins (Fig. 1), and essentially similar results were also obtained for pea (data not shown) and spinach [19]. The main phosphoproteins were identified as P-CP43, P-D2, P-D1 and P-LHCII, with protein-specific antibodies (Fig. 1). Detection of the 9 kDa phosphoprotein was not consistently successful. In pumpkin, nearly complete dephosphorylation of the phosphoproteins took place during the dark period and the level of phosphorylation of all the phosphoproteins increased after a shift to growth light (Fig. 1A). During high-light treatment, the PSII core protein phosphoproteins further increased whereas LHCII was almost completely dephosphorylated, as also shown earlier [19]. Fig. 1B depicts in more detail the steady-state phosphorylation pattern of thylakoid proteins in intact rye plants under different irradiance levels. Phosphorylation of D1 increased with increasing light intensity and did not show saturation in the light regime (up to 2000 μmol photons m⁻² s⁻¹) used in

![Image](image-url)

Fig. 1. Reversible thylakoid protein phosphorylation in higher plants. A: Steady-state phosphorylation level of thylakoid proteins in pumpkin leaves after incubation in darkness, in growth light (300 μmol photons m⁻² s⁻¹) and in high light (2000 μmol photons m⁻² s⁻¹). On the right hand side, an immunoblot (i) with protein-specific antibodies demonstrating the migration of CP43, D2, P-D1, D1 and LHCII. B: Thylakoid phosphoproteins in rye plants illuminated under different PFDs. On the right hand side, an immunoblot (i) showing the migration of CP29. The corresponding light intensities are indicated above the lanes.

![Image](image-url)

Fig. 2. Thylakoid protein phosphorylation in differentially light-treated lower plants with LHCII antenna. Ceratodon, Marchantia and Adiantum were exposed to darkness, to growth light and to high light, and the PSII phosphoproteins were analyzed with phosphothreonine-specific antibody. The corresponding light intensities are indicated above the lanes. For quantitative comparison of immunolabelling, a thylakoid sample of rye, illuminated at 600 μmol photons m⁻² s⁻¹, was run in parallel with the lower plant samples. On the right hand side, an immunoblot (i) with protein-specific antibodies demonstrating the migration of CP43, D2, P-D1, D1 and LHCII in Adiantum (A) and rye (R).
this experiment. Phosphorylation of D2 and CP43 responded to increasing irradiance very similarly to that of the D1 protein. As in pumpkin [19], LHCII showed maximal phosphorylation well below the growth light intensity (300 μmol photons m⁻² s⁻¹) and then sharply declined as irradiance increased. Interestingly, exposure of rye plants to high irradiance (2000 μmol photons m⁻² s⁻¹) also resulted in an accumulation of a 33 kDa phosphoprotein in the thylakoid membrane. This polypeptide cross-reacted with a CP29-specific antibody (Fig. 1B) and possibly represents the phosphorylated form of the 29 kDa chlorophyll ab binding protein [5]. It should be noted that in our gel system the phosphorylated and non-phosphorylated forms of CP29 could not be separated.

3.2. Thylakoid phosphoproteins in lower eukaryotic photosynthetic organisms with LHCII antenna

No distinct reversible D1 protein phosphorylation was detected in lower eukaryotic oxygenic organisms (Fig. 2). Hardly any signal was detected in the thylakoids of Ceratodon (moss) or Marchantia (liverwort) and only a faint immunoresponse in the region where P-D1 migrates in higher plants was obtained with the phosphothreonine antibody in Adiantum (fern). Moreover, this phosphoprotein was not recognized by the D1-specific antibody, which cross-reacted with both the non-phosphorylated and phosphorylated forms of D1 in the higher plants (Fig. 2). Neither did different irradiance conditions induce any D1 phosphorylation in these species. Opposite to the D1 protein, both D2 and CP43 were clearly phosphorylated in Ceratodon, Marchantia and Adiantum (Fig. 2). In Ceratodon and Marchantia, D2 and CP43 remained phosphorylated also in darkness and at low light, and the phosphorylation level hardly increased with increasing light intensity. However, in Adiantum, D2 and CP43 were almost completely dephosphorylated under growth light conditions and the level of phosphorylation clearly increased as the plant was shifted from growth light to high light, and a surprisingly high level of phosphorylation was also detected in dark-adapted plants.

Ceratodon, Marchantia and Adiantum all possess chlorophyll ab complexes as a light-harvesting system for PSII and also showed phosphorylation of LHCII polypeptides. Notably, changing light conditions induced similar reversible phosphorylation of LHCII in all chlorophyll b possessing organisms (chlorophyll ab ratios are given in Table 1): also in these lower organisms, the phospho-LHCII was present almost exclusively at growth light conditions. Hardly any immunoresponse with the phosphothreonine antibody was detectable after incubation of these species at high light or in darkness (Fig. 2).

3.3. Thylakoid phosphoproteins in phycobilisome-containing oxygenic organisms

No phosphorylation of PSII core proteins was detected in vivo in the cyanobacterium Synechocystis or in the red alga Ceramium (Fig. 3). The phosphothreonine antibody gave some cross-reaction with a 31 kDa and a 24 kDa polypeptide in Synechocystis (Fig. 3A). The level of phosphorylation of the 31 kDa protein was highest in darkness and somewhat less phosphorylation was observed both at low and high irradiances. Phosphorylation of the 24 kDa polypeptide was totally independent of the light conditions. In Ceramium, the phosphothreonine antibody gave an immunoresponse with a 67 kDa polypeptide (data not shown) and with 31 kDa, 22.5

Table 1
Chlorophyll ab ratios of the species with LHCII antenna

| Species             | Chlorophyll ab |
|---------------------|----------------|
| Ceratodon purpureus | 2.7            |
| Marchantia polymorpha | 2.6          |
| Adiantum tenerum    | 2.7            |
| Rye                 | 3.8            |
| Pumpkin             | 3.8            |
kDa, 15 kDa polypeptides (Fig. 3B). As in *Synechocystis*, also in *Ceramium*, strong phosphorylation of the 31 kDa protein was evident in darkness but at low and high irradiances the protein was completely dephosphorylated. The 22.5 kDa protein was phosphorylated under low light conditions while a complete dephosphorylation was observed under high light conditions and in darkness. The 15 kDa (Fig. 3B) and the 67 kDa polypeptides (data not shown) were phosphorylated in a light-independent manner. Both in *Synechocystis* and *Ceramium*, the protein-specific antibodies cross-reacted with only one form of D1, D2 and CP43, none of which co-migrated with the 31 kDa phosphoprotein (Fig. 3).

4. Discussion

Oxygencic photosynthetic organisms can be divided into three groups with respect to their PSII core protein phosphorylation. Cyanobacteria and red algae, with apparent lack of the phosphorylation of PSII core polypeptides D1, D2 and CP43, belong to group 1 organisms. Although light-dependent phosphorylation of both thylakoid and soluble proteins has previously been reported in cyanobacteria [25,26], all these experiments, like our present approach with polyclonal phosphothreonine antibody (Fig. 3), have failed to reveal any phosphorylation of PSII core proteins in cyanobacteria. Similarly, no phosphorylated PSII core proteins could be detected in the thylakoids of a red alga (Fig. 3). Prokaryotic prochlorophytes apparently also belong to this group [27,28], and it is worth noting that the chlorophyll *a/b* proteins present in this group have recently been shown not to belong to the Lhc gene family encoding the LHCCI proteins of chloroplasts [29].

Group 2 organisms consist of green algae [9], mosses, liverworts and probably also ferns. The most distinguishable feature of these species was the apparent lack of D1 protein phosphorylation (Fig. 2). Notably, all the species in group 2 exhibited phosphorylation of D2 and CP43. Clear variance, however, was evident inside this group with respect to the light regulation of D2 and CP43 phosphorylation. Only in the fern, a strong light-intensity-dependent phosphorylation of both D2 and CP43 could be clearly demonstrated.

Group 3 organisms, consisting of gymnosperms [10] and angiosperms (Fig. 1), both monocots and dicots, showed distinct reversible phosphorylation of all three PSII core phosphoproteins D1, D2 and CP43. Opposite to the saturation of PSII core protein phosphorylation already at moderate light intensities in pumpkin leaf discs [16], a linear increase in the phosphorylation of D1, D2 and CP43 with increasing light intensity up to 2000 μmol photons m−2 s−1 was apparent when intact rye plants were illuminated (Fig. 1).

It is noticeable that all organisms in groups 2 and 3, with a capacity to phosphorylate at least D2 and CP43, also possess reversible LHCCI phosphorylation (Figs. 1 and 2) [11,12,30]. In contrast, no phosphorylated PSII core proteins were detected in the eukaryotic red alga that possesses a phycobilisome antenna instead of LHCCI (Fig. 3). It is therefore conceivable that the PSII core protein phosphorylation is associated with the mechanism of light harvesting into PSII and raises a question whether the phosphorylation of LHCCI is the origin for PSII core protein phosphorylation. LHCCI phosphorylation has apparently evolved first, together with grana stacking, to regulate the distribution of excitation energy between the two photosystems. Grana stacking, although sometimes very irregular in group 2 species, is typical for species with reversible LHCCI phosphorylation. Whether the phosphorylation of D2 and CP43 evolved independently of LHCCI phosphorylation cannot be answered at the moment. Indisputably, however, the reversible D1 protein phosphorylation is the latest event in thylakoid protein phosphorylation and present exclusively in seed plants, both angiosperms and gymnosperms.

The lack of D1 protein phosphorylation in group 2 organisms suggests at least two different kinases in phosphorylation of thylakoid proteins, one for LHCCI, D2 and CP43 and the other for D1 protein phosphorylation. However, the regulation of LHCCI phosphorylation distinctively differs from that of D2 and CP43, as evidenced by the differential response of the phosphorylation of these proteins to increasing light intensity (Fig. 1) and by the phosphorylation studies with electron transport inhibitors and *cyt hol* mutants [31,32]. Therefore it is possible that even three different kinases are involved in reversible phosphorylation of thylakoid proteins in seed plants. Moreover, the kinase responsible for phosphorylation of CP29 is possibly distinct from the kinase(s) involved in phosphorylation of other PSII proteins. Despite intense research, none of the thylakoid-associated kinases have yet been isolated and characterized.

The absence of PSII core protein phosphorylation in cyanobacteria and red algae clearly demonstrates it unnecessary for the proper function of PSII. Reversible PSII core protein phosphorylation may instead have evolved as a regulatory mechanism for the acclimation of PSII function to changing conditions in more developed photosynthetic organisms. Reversible D1 protein phosphorylation has been attributed a distinct role in PSII photoinhibition-repair cycle in higher plants [16,33] with strict lateral heterogeneity of the thylakoid membrane. PSII core protein phosphorylation in general has been suggested to stabilize PSII dimers and has also been discussed in terms of the regulation of PSII repair [18]. It should be noted, however, that the dimerization [34] and stability of cyanobacterial PSII does not involve phosphorylation of any of the PSII core proteins.

Interestingly, the regulation of LHCCI phosphorylation with respect to irradiance level is principally similar in all organisms, with maximum phosphorylation at low light and nearly complete dephosphorylation in darkness and at high irradiances (Figs. 1 and 2). The inactivation of LHCCI kinase at high light intensities [19] was a general feature for all chlorophyll *b* containing photosynthetic organisms studied here and has earlier been reported also for *Chlamydomonas* cells [12]. Indeed, the complex regulation mechanism for LHCCI polypeptide phosphorylation seems to have remained principally similar throughout the eukaryotic organisms with chlorophyll *a/b* binding LHCCI polypeptides, whereas the phosphorylation of PSII core proteins shows much more versatility.

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