

# Chloroplast protein phosphorylation couples plastoquinone redox state to distribution of excitation energy between photosystems

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*In photosynthetic membranes isolated from pea leaves, the redox state of the plastoquinone pool controls both the level of phosphorylation of the chloroplast light-harvesting pigment-protein complex (LHC) and distribution of absorbed excitation energy between the two photosystems. Phosphorylation of LHC polypeptides is proposed as the regulatory mechanism by which photosynthetic systems adapt to changing wavelengths of light.*

IN chloroplasts, non-cyclic photosynthetic electron transport generates ATP and NADPH by the operation of photosystem I and photosystem II acting in series. The energy that drives this process is derived partly from light absorbed by antenna chlorophylls associated closely with each photosynthetic reaction centre and partly from the more peripheral light-harvesting pigments<sup>1</sup> which are organized into specific chlorophyll-protein complexes<sup>2-4</sup>. The most abundant of the chlorophyll-binding proteins in green plants is a 25,000 molecular weight (MW) polypeptide known as the light-harvesting chlorophyll *a/b* protein (LHCP).

In non-cyclic electron transport, the two photosystems must operate at the same rate. If absorbed radiant energy is distributed between the photosystems in such a way that one photosystem is excited more than the other, the excess excitation energy is dissipated, with a consequent loss of photosynthetic quantum efficiency. However, it is clear from studies with algal cells exposed to red and far-red light that chloroplasts can detect such an imbalance in the excitation of the photosystems and at least partially correct it by increasing the proportion of absorbed excitation energy that is distributed to the rate-limiting photosystem<sup>5-8</sup>. These adaptive changes are referred to as state 1-state 2 transitions. State 1 is the adaptive state which results from over-excitation of photosystem I (as, for example, by far-red light) and is characterized by an increase in the proportion of total absorbed excitation energy that is transferred to photosystem II relative to photosystem I. State 2 results from over-excitation of photosystem II by red light and is characterized by increased energy transfer to photosystem I.

The biochemical events underlying state 1-state 2 transitions are poorly understood. However, a related phenomenon that has been extensively studied at the biochemical level is the ability of metal cations to affect the distribution of absorbed excitation energy in isolated thylakoids<sup>9-11</sup>. In the presence of salts of monovalent and divalent metal cations<sup>12</sup>, the light-harvesting chlorophyll-protein complexes that contain LHCP are structurally and functionally associated with photosystem II, whereas in the absence of salts the association between the light-harvesting apparatus and the photosystems is altered and excitation energy transfer to photosystem I increases.

A major factor governing the response of thylakoids to cations *in vitro* is the LHC itself. Experiments using thylakoids have shown that trypsin can selectively remove a surface-exposed segment of the LHCP and that this removal is concomitant with the loss of the ability of cations to affect the distribution of excitation energy<sup>13</sup>. In addition, antibodies raised against isolated, purified LHC were found to react with surface-exposed segments of the membrane-bound complex and to prevent such cation effects<sup>14</sup>. These studies indicate that the

regulatory mechanism controlling the distribution of absorbed excitation energy in chloroplasts involves a surface-exposed segment of the LHCP.

It has recently been established that the LHCP is reversibly phosphorylated on a threonyl residue located within this surface-exposed segment<sup>15-18</sup>. The LHCP belongs to a group of thylakoid polypeptides shown to be phosphorylated by a membrane-bound, light-activated protein kinase<sup>19,20</sup> and dephosphorylated by a membrane-bound phosphatase<sup>17</sup>. Bennett *et al.*<sup>21</sup> have demonstrated that the reversible phosphorylation of LHCP changes the light-harvesting properties of isolated thylakoids: phosphorylation increases (and dephosphorylation decreases) the proportion of absorbed excitation energy transferred to photosystem I.

**Table 1** Activation of thylakoid protein kinase by light and reducing agents

Incubation conditions	Kinase activity (%)
Light	100
Dark	28
Light + DCMU	21
Light + DCMU + DCPIP <sub>2</sub>	18
Dark + dithionite	111
Light + dithionite	112
Light + DCMU + dithionite	121
Dark + TMOH <sub>2</sub>	101
Light + methyl viologen	27
Light + ferricyanide	17

Thylakoid membranes were prepared from 12-20-day-old peas (*Pisum sativum* L. var. Progress no. 9) grown as previously described<sup>3</sup>. Shoots (25 g) were homogenized for 5 s in a blender at low speed in 100 ml of 50 mM Tricine-NaOH (pH 7.8), 0.4 M sorbitol and 5 mM MgCl<sub>2</sub>. The homogenate was filtered through cheesecloth, and centrifuged at 1,000g for 5 min. The pelleted chloroplasts were osmotically shocked in 50 ml of 10 mM Tricine-NaOH (pH 7.8) and 5 mM MgCl<sub>2</sub>. The suspension was centrifuged at 2,800g for 5 min and the thylakoid pellet was resuspended to a final chlorophyll concentration of 1 mg ml<sup>-1</sup> in 50 mM Tricine-NaOH (pH 7.8), 100 mM sorbitol and 10 mM MgCl<sub>2</sub>. Portions (100 µl) of the thylakoid suspension were equilibrated at 20 °C in a final volume of 1 ml of the same buffer containing the indicated additions: 3-(3',4'-dichlorophenyl)-1,1-dimethylurea (DCMU) at 10 µM; 2,6-dichlorophenolindophenol (DCPIP) at 50 µM, maintained in the reduced form (DCPIPH<sub>2</sub>) by sodium ascorbate (2 mM); sodium dithionite (20 mM); tetramethylhydroquinone (TMOH<sub>2</sub>, 500 µM) reduced with sodium borohydride as described elsewhere<sup>23</sup>; methyl viologen (50 µM); potassium ferricyanide (2 mM). Equilibration took place either in the light (15 µE m<sup>-2</sup> s<sup>-1</sup>) or darkness, and, where dithionite was included, reaction mixtures were made anaerobic by bubbling them with nitrogen gas before the addition of thylakoids. Phosphorylation reactions were started by addition of [ $\gamma$ -<sup>32</sup>P]ATP (10 Ci mol<sup>-1</sup>) to a final concentration of 100 µM. The reaction was stopped after 4 min by addition of 0.5 ml of cold trichloroacetic acid (30%, w/v). Acid-stable, precipitable radioactivity was counted by Cerenkov spectrometry. The extent of protein phosphorylation was calculated from the mean activity of duplicated samples. Results are expressed as a percentage of the rate obtained in the light in aerobic conditions (5 nmol phosphate incorporated per mg chlorophyll in 4 min).

This article shows that the activity of the protein kinase is regulated by the redox state of plastoquinone. We propose a molecular mechanism for state 1–state 2 transitions *in vivo* that is consistent with these data and we argue that the presence of cations is merely a prerequisite for these transitions rather than a decisive regulatory factor in itself.

### Activation of the thylakoid protein kinase by light and reducing agents

The protein kinase activity of isolated thylakoids was assayed by measuring the transfer of  $^{32}\text{P}$  from  $[\gamma\text{-}^{32}\text{P}]\text{ATP}$  to total membrane protein (Table 1). Light stimulated the kinase about fourfold compared with a dark-incubated control. The herbicide 3-(3',4'-dichlorophenyl)-1,1-dimethylurea (DCMU), which blocks electron transport between the primary acceptor (Q) of photosystem II and plastoquinone, completely abolished the stimulatory effect of light. The stimulation could not be restored by addition of 2,6-dichlorophenolindophenol (DCPIP) in the reduced form (DCPIPH<sub>2</sub>), which donates electrons to the intersystem chain after plastoquinone. Thus, the kinase seemed to be stimulated by reduction of certain intersystem electron carriers. This idea was supported by the effects of oxidizing and reducing agents on kinase activity (Table 1). Ferricyanide and methyl viologen accept electrons from photosystem I and consequently tend to oxidize the intersystem carriers. These oxidants inhibited light-activation of the kinase. In contrast, the low-potential reductant dithionite<sup>22</sup> activated the kinase in the dark, as did tetramethylhydroquinone (TMQH<sub>2</sub>, a specific donor to plastoquinone in the intersystem electron transport chain<sup>23,39</sup>). Although dithionite was a somewhat more effective activator than light, their effects were equivalent rather than additive or synergistic. It is likely, therefore, that light and dithionite activated the kinase by the same basic mechanism, that is, by reduction of the intersystem carriers.

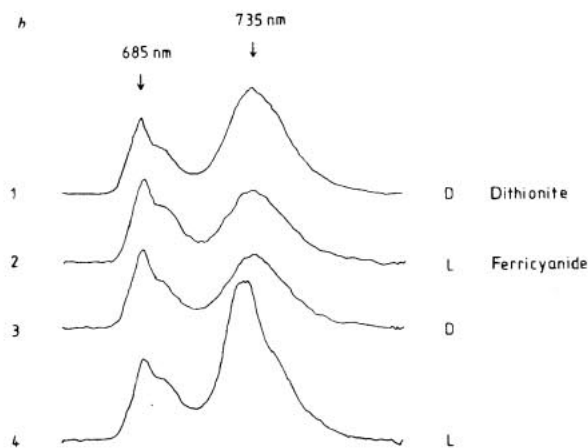
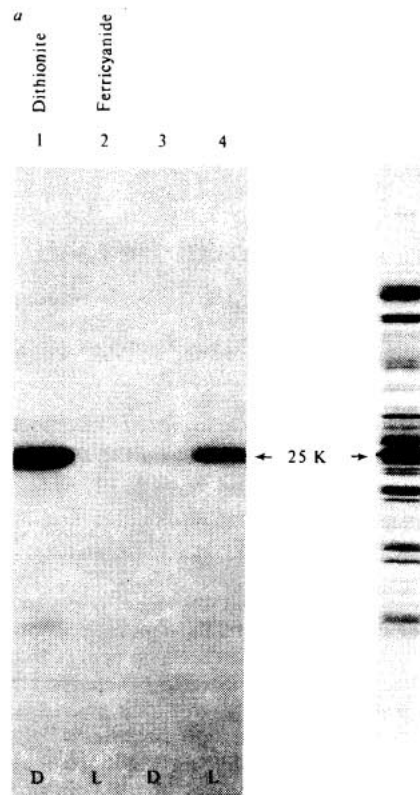
When activated by light, the protein kinase of the thylakoid phosphorylates predominantly the 25,000 MW LHCP<sup>16</sup>. Figure 1a shows that the same protein was phosphorylated when the kinase was activated in the dark by dithionite. Dithionite also stimulated phosphorylation of the 8,000 MW polypeptide reported to be the proton channel of the thylakoid ATP synthase<sup>20</sup>. In contrast, ferricyanide completely inhibited phosphorylation of these thylakoid polypeptides.

### Changes in the distribution of absorbed excitation energy

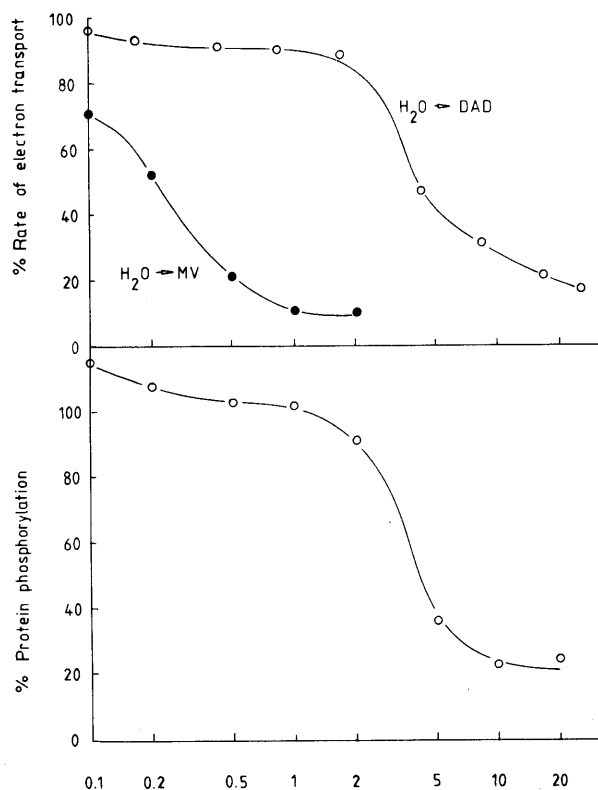
Bennett *et al.*<sup>21</sup> reported that light-induced protein phosphorylation altered the 77 K chlorophyll fluorescence emission spectrum of isolated thylakoids. The finding that emission at 735 nm ( $F_{735}$ ; derived mainly from the chlorophylls of photosystem I) was increased relative to emission at 685 nm ( $F_{685}$ ; derived mainly from light-harvesting chlorophylls associated with photosystem II<sup>1</sup>) was one of several lines of evidence<sup>21</sup> indicating that protein phosphorylation changed the distribution of excitation energy within the pigment bed of the membrane in such a way as to favour distribution to photosystem I at the expense of photosystem II.

If the above conclusion is correct, changes in the distribution of absorbed excitation energy should occur whenever the level of protein phosphorylation is altered regardless of how the latter is brought about. Figure 1b shows that the 77 K chlorophyll fluorescence emission spectrum of isolated thylakoids was changed by ferricyanide and dithionite in accordance with the effects of these reagents on protein phosphorylation. Ferricyanide inhibited the light-dependent increase in  $F_{735}$  while dithionite increased  $F_{735}$  in darkness, thus providing evidence for the control of excitation energy distribution by the redox state of the electron transport chain.

Although the dithionite-induced fluorescence change occurred in the presence of ATP, the data in Fig. 1 do not in themselves indicate that the change required ATP or involved protein phosphorylation. To explore this point in more detail,



**Fig. 1** Effects of light, dithionite and ferricyanide on phosphorylation of LHCP (a) and on 77 K fluorescence emission spectrum of thylakoids (b). a, Thylakoids that had been pre-equilibrated with or without dithionite (10 mM) in darkness and with or without ferricyanide (2 mM) in the light were incubated with  $[\gamma\text{-}^{32}\text{P}]\text{ATP}$  as described in Table 1 legend. Total  $^{32}\text{P}$  incorporation into acid-insoluble material was determined and then each sample was subjected to SDS-polyacrylamide gel electrophoresis, followed by staining (lane 5) and autoradiography (lanes 1–4). The identities of the lanes (together with total protein kinase activity of the thylakoids, expressed as a percentage of the light control) are: lane 1, dark + dithionite (124%); lane 2, light + ferricyanide (33%); lane 3, dark (16%); lane 4, light (100%). The position of the 25,000 MW LHCP is indicated (25 K). b, Thylakoids were pre-equilibrated and incubated as described above except that the dithionite concentration was 20 mM and the ATP was not radioactive. After incubation, thylakoids were quickly diluted 10-fold with a medium containing 50 mM Tricine-NaOH (pH 7.6), 30% (v/v) glycerol and 10 mM  $\text{MgCl}_2$  and then frozen to 77 K. Fluorescence emission spectra were obtained as described previously<sup>21</sup> using  $440 \pm 10$  nm light for excitation. Spectra were normalized to give equal relative fluorescence at 685 nm using a Hewlett-Packard 9825 computer.



**Fig. 2** Inhibition of electron transport and protein kinase activity by 2,5-dibromo-3-methyl-6-isopropyl-*p*-benzoquinone (DBMIB). Whole chain electron transport was measured as oxygen uptake in the presence of methyl viologen (MV, 100  $\mu$ M), sodium azide (5 mM) and ATP (100  $\mu$ M). The control rate in the absence of DBMIB was 23  $\mu$ mol (mg chlorophyll h)<sup>-1</sup>. Photosystem II-mediated electron transport was measured as oxygen evolution in the presence of K<sub>3</sub>Fe(CN)<sub>6</sub> (2.5 mM), diaminodurene (DAD, 1.2 mM) and ATP (100  $\mu$ M). The control rate was 97  $\mu$ mol (mg chlorophyll h)<sup>-1</sup>. All electron transport measurements were carried out at saturating light intensity (1 mE m<sup>-2</sup>s<sup>-1</sup>) in a reaction medium (2 ml) containing 50 mM Tricine (pH 7.8), 100 mM sorbitol, 10 mM MgCl<sub>2</sub> and chloroplasts equivalent to 200  $\mu$ g chlorophyll. Protein kinase activity was measured as described in Table 1 legend. DBMIB was present during the equilibration period before the addition of [ $\gamma$ -<sup>32</sup>P]ATP. Values for phosphorylation were calculated by subtraction of the activity of a dark control and are expressed as a percentage of the activity in the light in the absence of DBMIB.

thylakoids were incubated in various conditions in the presence and absence of ATP and dithionite. Values for the ratio  $F_{735}/F_{685}$  were calculated from the resulting 77 K fluorescence emission spectra (Table 2). An ATP-dependent increase in the ratio was observed only in conditions which also permitted protein phosphorylation. Thus, there was no significant effect of ATP on fluorescence in the dark or in the light in the presence of DCMU, unless dithionite was also present. It is also clear from Table 2 that dithionite exerted two effects on the  $F_{735}/F_{685}$  ratio, one depending on the presence of ATP and the other not. The effect which required both ATP and dithionite represented the change in the distribution of excitation energy that could be attributed to dithionite-activated protein phosphorylation. The effect of dithionite in the absence of ATP may result from a direct alteration in fluorescence yield.

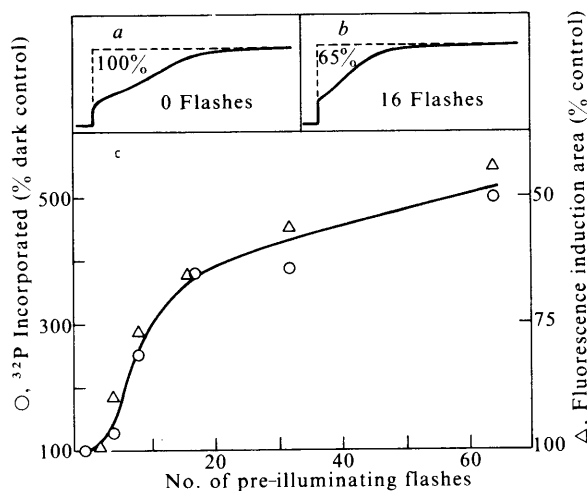
### Activation of protein kinase by reduced plastoquinone

The above data raised the possibility that plastoquinone was the electron transport component whose reduction led to kinase activation. This possibility was tested by the use of 2,5-dibromo-3-methyl-6-isopropyl-*p*-benzoquinone (DBMIB), which inhibits electron transport at two sites—it inhibits oxidation of plastoquinone at a high-affinity site<sup>24</sup> and reduction of plastoquinone at a secondary, low-affinity site<sup>25</sup>. Figure 2 illus-

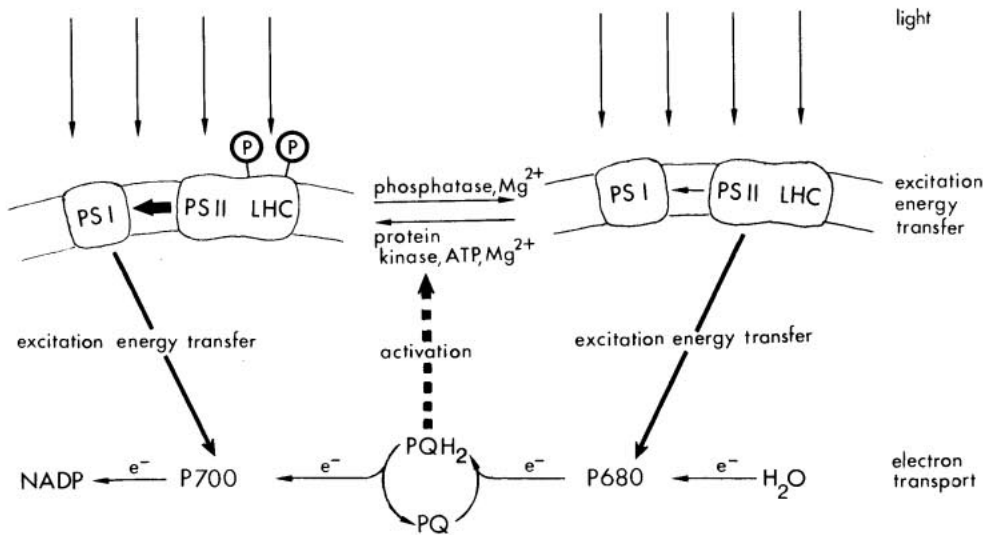
trates both inhibitory activities. At a concentration of 1  $\mu$ M, DBMIB inhibited whole chain electron transport (from water to methyl viologen) by 90%, whereas photosystem II-mediated reduction of diaminodurene (DAD) was strongly inhibited only at concentrations greater than about 10  $\mu$ M (DAD accepts electrons at or before plastoquinone<sup>26</sup>). When light-dependent kinase activity was measured as a function of DBMIB concentration (Fig. 2), strong inhibition was obtained only at concentrations equal to or greater than 10  $\mu$ M. Thus, the electron transport component that links the activity of the protein kinase to the redox state of the intersystem carriers is situated between the primary and secondary sites of action of DBMIB; the most obvious candidate is plastoquinone itself.

To characterize further the possible relationship between the redox state of plastoquinone and kinase activation, we subjected a series of thylakoid samples to short, intense flashes of light. The redox state of the plastoquinone pool was assessed by recording chlorophyll fluorescence induction transients (Fig. 3). The area above the curve of the transient is proportional to the fraction of the plastoquinone pool that is in the oxidized state<sup>27</sup>. As shown in Fig. 3a and b, pre-illuminating flashes decreased the area above the curve of the transient, indicating that the plastoquinone pool had accumulated reducing equivalents. In parallel measurements, [ $\gamma$ -<sup>32</sup>P]ATP was injected in the dark into preflashed samples which were then incubated for 4 min to permit kinase activity assay. As shown in Fig 3c, there was a striking similarity between the extent of kinase activation and that of reduction of the plastoquinone pool, as measured by the decrease in the area above the curve of the transient. Thus, the activity of the kinase is linked to the redox state of plastoquinone.

The previous observation<sup>19</sup> that NADPH and ferredoxin could together activate the protein kinase in darkness may now



**Fig. 3** Thylakoid protein phosphorylation and redox state of plastoquinone as functions of the number of pre-illuminating single-turnover flashes. Thylakoids were diluted with a medium containing 50 mM Tricine (pH 7.8), 100 mM sorbitol, 10 mM MgCl<sub>2</sub> and 10 mM NaF to a chlorophyll concentration of 50  $\mu$ g ml<sup>-1</sup> for protein kinase assays or 5  $\mu$ g ml<sup>-1</sup> for room temperature chlorophyll fluorescence induction transients. The dilution medium had been bubbled with nitrogen before the addition of thylakoids. The samples were pre-illuminated with a flashlamp (General Radio 'Stroboslav', type 1539A) delivering one intense, 8- $\mu$ s flash per s. A fibre-optics bundle carried light from the flash-lamp directly into either cuvettes (for the induction transients) or polyethylene microcentrifuge tubes (for kinase assays). After the indicated number of flashes, fluorescence induction was measured in the cuvettes as previously described<sup>3</sup>. a, b, Direct tracings of oscillographic recordings; broken lines delimit the area above the induction curve, a measure of the proportion of the plastoquinone pool that is in the oxidized state. c, [ $\gamma$ -<sup>32</sup>P]ATP was added to the microcentrifuge tubes for a 2-min, dark, anaerobic incubation which was terminated by addition of trichloroacetic acid to a concentration of 10%. Protein phosphorylation was then measured as described in Table 1 legend. Fluorescence measurements and additions of [ $\gamma$ -<sup>32</sup>P]ATP were made within 5 s of the final pre-illuminating flash.



**Fig. 4** A model for control of distribution of excitation energy in photosynthesis, in which reversible phosphorylation of LHC couples the redox state of plastoquinone (PQ) to the distribution of excitation energy between the photosystems. The mechanism we propose ensures maximal quantum efficiency of non-cyclic electron transport from water to NADP, mediated by P680 and P700, the reaction centres of photosystem II (PSII) and photosystem I (PSI), respectively. Reduction of plastoquinone by photosystem II leads to kinase activation and LHC phosphorylation, and hence to increased excitation of photosystem I. Conversely, oxidation of plastoquinone (PQH<sub>2</sub>) by photosystem I inactivates the kinase; the phosphatase then dephosphorylates LHC and hence excitation of photosystem II is increased relative to that of photosystem I. Since the redox state of plastoquinone will be determined in part by the distribution of excitation energy between the two photosystems, reversible LHC phosphorylation completes a feedback loop by means of which any imbalance in this distribution will tend to be self-correcting.

be explained as an indirect effect, resulting from the ability of reduced ferredoxin to reduce plastoquinone<sup>28,29</sup>. We do not believe that activation of the thylakoid-bound protein kinase is mediated by either the thioredoxin system<sup>30</sup> or the 'LEM' system<sup>31</sup>, both of which have been implicated in the control of stromal enzyme activity and require electron transport via components on the reducing side of photosystem I.

### Regulation of photosynthetic efficiency by reversible protein phosphorylation

The involvement of plastoquinone in the activation of the thylakoid protein kinase is probably of fundamental importance in the regulation of photosynthesis, for it provides a physiological mechanism for an adaptation which will correct any imbalance in the distribution of excitation energy between the two photosystems (Fig. 4).

The operation of this mechanism may be illustrated by reference to state 1-state 2 transitions. In green algae and higher plants, that is, in photosynthetic organisms containing LHCP, exposure to red light (of wavelength <680 nm) will excite photosystem II to a greater extent than photosystem I, and thereby lead to reduction of plastoquinone. The protein kinase will then be activated, LHCP will become phosphorylated and an increased proportion of absorbed excitation energy will be transferred from the light-harvesting complex to photosystem I at the expense of photosystem II. Thus, the biochemical event underlying the transition to state 2 is phosphorylation of the LHCP. Conversely, under far-red light (of wavelength >680 nm), photosystem I will be excited to a greater extent than photosystem II, plastoquinone will be oxidized and the kinase will be inactivated. Any phosphorylated LHCP will be dephosphorylated by the thylakoid phosphatase and excitation energy transfer to photosystem I will be decreased. Thus, the biochemical event underlying the transition to state 1 is dephosphorylation of the LHCP.

It has often been suggested that state 1-state 2 transitions reflect changes in the cation content of the stroma surrounding the thylakoids<sup>9-11,32</sup>, a suggestion based on the effect of cations on the distribution of excitation energy in isolated thylakoids<sup>9-11</sup>. One problem with this idea is that state 2 transitions would have to be correlated with a massive loss of cations from the stroma (to either the thylakoid or the cytoplasm), but there is no reason to believe that the cation concentration of the stroma changes greatly when higher plants or algae are transferred from far-red light to red light or vice versa. However, it should not be assumed that cations are not involved in the regulation of excitation energy distribution *in vivo*. Two roles need to be emphasized. First, cations such as Mg<sup>2+</sup> ions are required to establish the pattern of excitation energy distribution seen in dephosphorylated membranes, including the close structural and functional association between the LHCP complex and photosystem II<sup>10,12,33</sup>. The second role for Mg<sup>2+</sup> ions derives from the fact that these cations are cofactors of both the kinase and the phosphatase (Fig. 4)<sup>17,19,20</sup>. Thus, the presence of cations should be thought of as a prerequisite for state 1-state 2 transitions rather than as a regulatory factor *per se*.

The mechanism depicted in Fig. 4 provides an explanation for the original results of Bonaventura and Myers<sup>5</sup> and of Murata<sup>6</sup> and also for more recent contributions to the study of state 1-state 2 transitions. Several authors have drawn attention to the involvement of the high energy state of the membrane in these transitions<sup>9,34,35</sup>. This would certainly be required for the generation of ATP consumed in protein phosphorylation (transition to state 2). Duysens<sup>36</sup> suggested that the redox state of a component of the intersystem electron transport chain controls these transitions. Plastoquinone would clearly fulfil this role. Horton and Black<sup>37,38</sup> have recently established state 1-state 2 transitions *in vitro*. The transition to state 2 required ATP and was controlled by a membrane-bound factor with the redox potential of plastoquinone. The idea that protein phosphorylation might provide the connection between plastoquinone and ATP is fully in agreement with our present results.

It is important to recognize that state 1-state 2 transitions are also observed in organisms that lack LHCP. For example, Murata<sup>6</sup> and Ried and Reinhardt<sup>8</sup> have shown that these transitions occur in red algae, which contain phycobiliproteins instead of LHCP complexes. Ried and Reinhardt<sup>8</sup> conclude

**Table 2** Effects of light, dithionite, DCMU and ATP on the ratio of 77 K fluorescence emissions at 735 nm and 685 nm

Incubation conditions*	$F_{735}/F_{685}$		
	+ATP	-ATP	(a)/(b)†
Dark	1.00	0.95	1.05
Light	1.90	1.20	1.58
Dark + dithionite	1.45	1.10	1.32
Light + dithionite	1.85	1.30	1.42
Light + DCMU	0.85	0.90	0.94
Light + DCMU + dithionite	1.90	1.30	1.46

\* As described in Fig. 1b; DCMU concentration, 10 μM.

† Values greater than 1 indicate an ATP-dependent change in excitation energy distribution in favour of photosystem I.

from their studies that the transitions are controlled by the redox state of a component of the intersystem electron transport chain. It will be interesting to discover whether state 1-state 2 transitions in the red algae are brought about by a mechanism analogous to that shown in Fig. 4; that is, a mechanism in which reversible phosphorylation of the phycobiliproteins is controlled by the redox state of plastoquinone.

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*Note added in proof:* Another piece of evidence for the mechanism in Fig. 4 has now been obtained. Potentiometric redox titrations of plastoquinone, of LHCP phosphorylation and of ATP-dependent chlorophyll fluorescence quenching give identical results<sup>40</sup>.

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