Light-dependent Translocation of a Phytochrome B-GFP Fusion Protein to the Nucleus in Transgenic Arabidopsis

Rumi Yamaguchi,* Masanobu Nakamura,* Nobuyoshi Mochizuki,* Steve A. Kay,‡ and Akira Nagatani*

*Department of Botany, Graduate School of Science, Kyoto University, Kyoto 606-8502, Japan; and ‡National Science Foundation Center for Biological Timing, Department of Cell Biology, Scripps Research Institute, La Jolla, California 92037

Abstract. Phytochrome is a ubiquitous photoreceptor of plants and is encoded by a small multigene family. We have shown recently that a functional nuclear localization signal may reside within the COOH-terminal region of a major member of the family, phytochrome B (phyB) (Sakamoto, K., and A. Nagatani. 1996. Plant J. 10:859–868). In the present study, a fusion protein consisting of full-length phyB and the green fluorescent protein (GFP) was overexpressed in the phyB mutant of Arabidopsis to examine subcellular localization of phyB in intact tissues. The resulting transgenic lines exhibited pleiotropic phenotypes reported previously for phyB overexpressing plants, suggesting that the fusion protein is biologically active. Immunoblot analysis with anti-phyB and anti-GFP monoclonal antibodies confirmed that the fusion protein accumulated to high levels in these lines. Fluorescence microscopy of the seedlings revealed that the phyB-GFP fusion protein was localized to the nucleus in light grown tissues. Interestingly, the fusion protein formed speckles in the nucleus. Analysis of confocal optical sections confirmed that the speckles were distributed within the nucleus. In contrast, phyB-GFP fluorescence was observed throughout the cell in dark-grown seedlings. Therefore, phyB translocates to specific sites within the nucleus upon photoreceptor activation.

Key words: green fluorescent protein • nuclear targeting • photomorphogenesis • phytochrome • signal transduction

Light is an important environmental stimulus which plants must perceive and to which they must respond. Plants use light signals to regulate various developmental processes such as seed germination, de-etiolation, and floral induction (Kendrick and Kronenberg, 1994). For this purpose, plants have evolved several different photoreceptors. Among them, phytochrome is the best characterized. Phytochrome is a soluble chromoprotein consisting of a apoprotein of 120 kD and covalently attached linear tetrapyrrole chromophore (Furuya, 1993; Quail et al., 1995). Phytochrome is a ubiquitous photoreceptor in the plant kingdom, the origin of which can be traced back to cyanobacteria (Kehoe and Grossman, 1996; Hughes et al., 1997; Yeh et al., 1997).

Phytochrome undergoes photoreversible conversion between two spectrally distinct forms, a red light absorbing form (Pr) and a far-red light absorbing form (Pfr). Only the Pfr form is believed to be biologically active. Red light activates phytochrome by converting it from the Pr to Pfr form. Conversely, far-red light cancels the effects of red light. In this way, phytochrome acts as a molecular switch. The phytochrome protein is comprised of two domains. The NH₂-terminal portion, to which the chromophore is attached, confers the spectral properties characteristic of phytochrome. The COOH-terminal portion is involved in dimerization of the molecule and transfer of the signal to downstream components (Quail, 1997).

Phytochrome has been studied intensively since its discovery in 1959 (Sage, 1992). Nevertheless, little was known about the initial step of the phytochrome signal transduction until recently. However, two recent studies hint at how phytochrome transduces the light signal to downstream components. Firstly, a phytochrome-interacting factor, PIF3, has been identified through a yeast two-hybrid screen (Ni et al., 1998). Interestingly, PIF3 is a nuclear-localized basic helix-loop-helix protein. Hence, a direct interaction between phytochrome and a transcriptional regulator might be involved in the signaling pathway within the nucleus. Secondly, phytochrome appears to be a light-regulated serine/threonine kinase (Yeh and Lagarias, 1998), and therefore may transmit light perception via protein phosphorylation.

Address correspondence to Akira Nagatani, Department of Botany, Graduate School of Science, Kyoto University, Sakyo-Ku, Kyoto 606-8502, Japan. Tel.: 81-75-753-4123. Fax: 81-75-753-4126. E-mail: nagatani@physiol.bot.kyoto-u.ac.jp

1. Abbreviations used in this paper: DIC, differential interference contrast; GFP, green fluorescent protein; Pfr, far-red light absorbing form of phytochrome; phy, phytochrome.
A wide range of physiological and developmental processes is under the control of phytochrome. Accordingly, phytochromes are expressed in various tissues throughout the life cycle of plants (Nagatani, 1997). The mode of phytochrome action varies substantially (Mancinelli, 1994). For example, some responses are induced by a relatively low fluence of red light, whereas prolonged irradiation with far-red light is required for some responses. The rate of escape from the red/far-red reversibility varies substantially depending on responses. The diversity of phytochrome action can be explained in part by the multiple molecular species. Phytochrome is known to be encoded by a small multigene family (Mathews and Sharrock, 1997). In Arabidopsis, the complete family consists of five members, phytochromes (phy) A–E, that are encoded by respective genes (PHYA–E). A nalysis of mutants deficient in phyA and phyB suggests that the modes of their action are different (Furuya and Schaefer, 1996; Shimomura et al., 1996). It has also been shown that phyC (Halliday et al., 1997; Qin et al., 1997), phyD (Aukerman et al., 1997), and phyE (Devlin et al., 1998) are functionally different from phyA and phyB. These findings imply that different molecular species of phytochrome may transduce light signals via distinct mechanisms.

To elucidate the signal transduction mechanisms of different phytochromes, it is essential to know the sites of their action within the cell. Since biochemical analysis has indicated that phyA resides in the cytoplasm in darkness, it has been assumed that phytochrome action takes place in the cytoplasm (Nagatani, 1997). In accordance with this notion, phyA and phyB, and probably other phytochromes, are soluble proteins. In addition, microbeam irradiation experiments in green algae and fern gametophytes have indicated that phytochrome, which mediates various cellular responses in these systems, resides in the cytoplasm (Wada et al., 1993). Although associations of phyA with isolated organelles have been reported repeatedly, the biological relevance of these observations remains obscure (Pratt, 1994).

More recently, we have produced transgenic Arabidopsis expressing fusion proteins consisting of GFP and COOH-terminal fragments of phyB (Sakamoto and Nagatani, 1996). The CAT staining from the fusion proteins is observed in the nucleus, suggesting that a functional nuclear localization signal may reside in the phyB sequence. Furthermore, we have confirmed that a substantial fraction of total cellular phyB is recovered in the isolated nuclei. Interestingly, the level of nuclear phyB is substantially reduced by the dark adaptation of plants. On the basis of these findings, we have proposed that phyB translocates to the nucleus upon photoactivation (Sakamoto and Nagatani, 1996; Nagatani, 1997). However, we could not exclude the possibility that those observations were due to technical artifacts.

In this work, the green fluorescent protein (GFP) of the jelly fish was fused to phyB and expressed in the phyB mutant of Arabidopsis to determine its intracellular localization in vivo. Since GFP is relatively small and tolerates protein fusion, it has been shown to be potentially useful as a fluorescent tag (Chiu et al., 1996). The fluorescence emission of GFP does not require any cofactor or substrate, which enables us to observe its fluorescence without making any pretreatment of the tissue. The result was transgenic lines exhibited pleiotropic phenotypes reported previously for the phyB overexpressing plants, indicating that the phyB-GFP fusion protein is biologically active. Fluorescent microscopic observation revealed that the fusion protein was localized to the nuclear region in the light. Confocal microscopic analysis confirmed that the fusion protein was indeed inside the nucleus. The effects of light on the nucleocytoplasmic partitioning of phyB were then examined. In dark-grown seedlings, fluorescence was observed throughout the cell. Treatment of the seedlings with continuous red light induced accumulation of phyB-GFP fusion protein in the nucleus. Hence, we suggest that phyB translocates to the nucleus upon light stimulation.

Materials and Methods

Plant Materials

The phyB-5 mutant (R edd et al., 1993) of Arabidopsis thaliana (ecotype, Landsberg) was used as the host for transformation. Arabidopsis thaliana (ecotype Landsberg) and the phyB-5 mutant were used as controls for physiological, immunochromical, and microscopic experiments.

Plasmid Construction and Transformation

A full-length PHYB cDNA clone was isolated from an Arabidopsis (ecotype Columbia) cDNA library. Cloned PHYB cDNA was almost identical to a previously reported sequence (accession number X 17942, submitted by Dr. R. Sharrock, Montana State University, Bozeman, MT) except that a C to T substitution at the base position 971, which does not cause amino acid difference, was detected. To construct the PHYB-GFP fusion sequence, PHYB translational termination codon (TAG) was replaced with an oligonucleotide sequence (GGAGGGTACGATG) by PCR. This oligonucleotide introduces a unique ClaI restriction site at its 3' terminus. The GFP clone (blue-sGFP-TG-nos KS) (Chiu et al., 1996) was a kind gift from Dr. J. Sheen (Massachusetts General Hospital, Boston, MA). A unique ClaI restriction site that shortly precedes the ATG start codon of the GFP gene. The PHYB and GFP clones were ligated at the ClaI restriction site to generate PHYB-GFP translational fusion. As a result, an oligonucleotide sequence (GGGGID-KLD) was inserted between the constitutive cauliflower mosaic virus 35S promoter and the Nos terminator of Agrobacterium tumefaciens transformation vector pBI1-Hyg/35S-NosT (Fig. 1a). This PHYB-GFP chimeric cassette was inserted into the pBI-Hyg vector (Chiu et al., 1996) to generate E. coli clones (blue-sGFP-TG-nos KS) (Chiu et al., 1996) was used as the host for transformation. Arabidopsis thaliana (ecotype Landsberg) and the phyB-5 mutant were used as controls for physiological, immunochromical, and microscopic experiments.

Growth Conditions and Light Treatments

For growth of plants, seeds were sown on 0.6% agar plates containing the Murashige-Skoog medium with 2% (wt/vol) sucrose and grown under continuous white light from fluorescent tubes (FLR 40S/W; Hitachi). The plants were then transplanted to pots containing vermiculite and grown to maturity under continuous white light from fluorescent tubes. For the immunochromical detection of the fusion protein, rosette leaves were harvested from 3-wk-old plants. For the hypocotyl assay and microscopic observation, seeds were sown on agar plates containing Murashige-Skoog salt mixture without sucrose. The plates were placed at 4°C for 12 h and then irradiated with continuous white light for 12 h at 23°C to induce germination. For the hypocotyl assay, seedlings were grown for 5 d under continuous red light (6.0 W m⁻²) from red fluorescent tubes (FL20S/R; National) or in darkness. For microscopic observation, seedlings were
grown for 5 d under continuous white light (15 W m\(^{-2}\)) from fluorescent tubes (FLR 40SW/M-B; Hitachi) or in darkness.

**Immunological Experiments**

To detect the phyB-GFP fusion protein and the authentic phyB, 0.1 g of rosette leaves was glass homogenized in the presence of 0.1 ml of the phytochrome extraction buffer (100 mM Tris-HCl, 2 mM DTT, 5 mM EDTA, pH 8.3) containing proteinase inhibitor cocktails for general use (P2714; Sigma Chemical Co.) and for fungal and yeast extracts (P8215; Sigma Chemical Co.) at the concentrations recommended by the manufacturer. Debris was removed by centrifugation. Proteins were concentrated from the crude homogenate by ammonium sulfate precipitation. The precipitated protein was dissolved in the SDS-PAGE sample buffer and subjected to immunoblot analysis (Sakamoto and Nagatani, 1996). Antibodies used were an anti-phyB mAb, mBA2 (Shinomura et al., 1996), and an anti-GFP mAb (Clontech). Molecular weight markers (prestained SDS molecular weight standard mixture) were from Sigma Chemical Co.

**Microscopic Observation**

Arabidopsis seedlings were soaked in 2 \(\mu\)g ml\(^{-1}\) Hoechst No. 33342 (Sigma Chemical Co.) solution made in H\(_2\)O for visualization of the nucleus in some experiments. Epidermal layers including cortex were peeled from the hypocotyl and placed on glass slides. For the other parts of seedlings, whole organs were placed on glass slides and pressed gently. The specimens were observed using an Olympus BX 60 microscope equipped with \(\times 20\), \(\times 40\), and \(\times 100\) objectives, differential interference contrast (DIC) optics, and a 100-W mercury arc light source. Fluorescence was filtered using UV (U-MWU) or FITC (U-MNIBA) filter sets (Olympus).

For confocal microscopy, trichomes were removed from the surface of cotyledons with a razor blade and placed on glass slides. Root tips were placed on glass slides without any pretreatment. The specimens were observed using an inverted laser scan microscope (LSM 410 invert; Carl Zeiss Jena) equipped with \(\times 40\) and \(\times 63\) objectives. The laser scan images were obtained with a combination of 488 nm laser excitation and 515 nm long-pass emission filter (LPS515; Carl Zeiss Jena). Sequential images from different focus planes were recorded automatically.

**Results**

**phyB-GFP Is Biologically Active in Transgenic Plants**

To examine biological activity and intracellular localization of the phyB-GFP fusion protein, the phyB-5 mutant of Arabidopsis was transformed with a vector harboring the 35S::PHYB-GFP construct. The resulting transgenic lines, PBG-5 and PBG-7, exhibited an overall dwarfing of mature plants under continuous white light (Fig. 1, b–e). They flowered a few days later than the wild-type under the conditions tested. Similar phenotypes, which are opposite to those of the phyB-deficient mutants (Reed et al., 1993), have been reported in phyB overaccumulating plants (Wester et al., 1994). Hence, the phyB-GFP fusion protein is likely to be fully functional.

It is known that inhibition of hypocotyl elongation by continuous red light is mediated primarily by phyB (Quail et al., 1995). To confirm the biological activity of phyB-GFP further, heterozygous progeny of the PBG-5 plant.
was examined for this response. The seedlings were grown under continuous red light for 5 d and hypocotyl lengths were determined. As shown in Fig. 2, a short population segregated from a longer one at about a 3:1 ratio. Hypocotyl lengths in the longer population matched well with those in the parental phyB mutant. In contrast, the shorter seedlings were significantly shorter than the wild-type seedlings, which is consistent with the phyB overexpression phenotypes reported by other groups (Wagner et al., 1991; McCormac et al., 1993). Cosegregation of the short phenotype with the expression of phyB-GFP was then examined. As expected, all the short seedlings exhibited GFP fluorescence whereas no fluorescence was observed in the longer seedlings (Fig. 2).

The seedling phenotype was examined in darkness as well. As is the case with the phyB overexpressing plants (Wagner et al., 1991), no clear segregation of shorter seedlings was observed in the PBG-5 heterozygous progeny (Fig. 2). The average hypocotyl lengths in the fluorescent and nonfluorescent populations were indistinguishable. Hence, phyB-GFP was suggested to be not only biologically but also photochemically active.

**Immunoblot Analysis Confirms Accumulation of phyB-GFP**

To examine the accumulation of phyB-GFP fusion protein in the transgenic plants, immunoblot analysis was performed. Proteins were extracted from rosette leaves of the PBG-5 plants and probed with anti-phyB and anti-GFP antibodies (Fig. 3). The anti-phyB mAb detected a major band of ~143 kD in the PBG-5 extracts (Fig. 3, left). The size was consistent with the expected mass of the phyB-GFP fusion protein. A band at the same size was detected with the anti-GFP antibody (Fig. 3, right), confirming that the band represented the phyB-GFP fusion protein. The higher intensity of the phyB-GFP band compared with that of the authentic phyB indicated that the phyB-GFP was overaccumulated in the transgenic plants. A similar result was obtained for the other transgenic line, PBG-7 (data not shown).

In addition to the major 143-kD band, a weak band of ~123 kD was detected in the PBG-5 plants (Fig. 3). The intensity of the band was comparable to that of authentic phyB. To confirm that the fragment is larger than the authentic phyB (117 kD on the blot), extracts from the PBG-5 and the wild-type plants were mixed and probed with the anti-phyB antibody. As expected, the two bands were separated on the blot. Since the fragment was not detected with anti-GFP antibody, it is speculated that proteolysis of phyB-GFP within the GFP portion yielded this fragment. In accordance with this, minor bands around 20 kD were detected on the anti-GFP blot. In the absence of the protease inhibitor cocktails, fragmentation was much more severe (data not shown). Hence, the 123-kD fragment is likely to be produced by the residual proteolytic activity in the extract during the extraction procedure.

**phyB-GFP Localizes to the Nucleus in the Light**

Intracellular localization of the phyB-GFP fusion protein in the PBG-5 seedlings was examined. Epidermal layers including cortex were peeled from the light-grown seedlings and observed under a fluorescence microscope. At lower magnification, bright green spots of GFP fluores-
Fluorescence microscopy was used to observe the localization of phyB-GFP. The fluorescence images showed that the spots were located within the nuclei, and their positions matched well with those revealed by Hoechst staining. Another transgenic line, PBG-7, showed similar fluorescence images (data not shown). At higher magnification, the phyB-GFP fluorescence was speckled within the nuclear region. The apparent size of each speckle appeared to be around 1 μm. The number of speckles per nucleus varied, with most cases containing 5–10 speckles.

The intracellular localization of phyB-GFP was examined in other parts of the seedling. As shown in Figure 5, fluorescence was observed in leaf, root, and root hair cells. The speckles were observed in all parts of the cell types examined. To determine the spatial distribution pattern of the speckles within the nucleus, optical sectioning was performed using a confocal microscope. In this case, the speckles appeared to be distributed more or less evenly in the nucleus. In most cases, at least 24 spots were recognized. This may be due to the large size of the trichome nucleus. The images clearly demonstrated that the size of each speckle varied substantially even within one nucleus.

**phyB-GFP Is Distributed throughout the Cell in Darkness**

A previous study suggested that the nuclear localization of phyB is light dependent (Sakamoto and Nagatani, 1996). In accordance with this, weak fluorescence was observed throughout the cell in dark-grown seedlings. This weak fluorescence was observed only in the peripheries but also in the nuclear region. It was difficult to conclude that phyB-GFP exists inside the nucleus.
nucleus even by confocal observation (data not shown).

Intracellular distribution of phyB-GFP in the light and darkness was compared in root tip cells with a confocal microscope. As shown in Fig. 7 j, the speckles of fluorescence were observed in light-grown seedlings. In contrast, relatively uniform fluorescence was observed in the peripheries of the cells in dark-grown seedlings (Fig. 7 i), which provided further evidence that phyB-GFP was distributed outside the nucleus and throughout the cell in darkness.

Red Light Induces Nuclear Accumulation of phyB-GFP

The time course of nuclear accumulation of phyB-GFP during the dark to light transition was followed. The PBG-5 dark-grown seedlings were transferred under continuous red light. As shown in Fig. 8, nuclear fluorescence was not clear at time 0 (Fig. 8, a and b). After 2 h in red light, the intensity of the nuclear GFP signal was increased (Fig. 8, c and d). However, fluorescence remained detectable in the periphery of the cells. Speckles in the nucleus were rarely observed at this time point, although a few tiny spots were detected in some cases. After 4 h in red light, many small speckles were observed (Fig. 8, e and f). Fluorescence in the cell periphery was greatly reduced. After 6 h in red light, the speckles became larger but the number per nucleus was reduced (Fig. 8, g and h). Hence, translocation of phyB-GFP to the nucleus appeared to be completed within 4–6 h in hypocotyl cells under continuous red light. In the course of these experiments, we noticed that the translocation took longer in root cells, although the reason for this was not clear (data not shown).

Discussion

phyB-GFP Is Biologically Active

It is known that phytochromes overexpressed in transgenic plants are biologically active. Transgenic Arabidopsis expressing exogenous phyB exhibits increased sensitivity to red light (Wagner et al., 1991; McCormac et al., 1993). In this study, we have demonstrated that the plants expressing phyB-GFP show similar light-dependent phenotypes (Figs. 1 and 2). Since the expression level of phyB-
GFP was comparable to those reported for the phyB overspeaking plants (Wagner et al., 1991), it is concluded that phyB-GFP is as active as authentic phyB. Furthermore, the fusion protein was expressed in the phyB-deficient background in the present study, confirming that the presence of endogenous phyB is not required for correct functioning of the phyB-GFP fusion protein.

In the PBG-5 seedlings, a proteolytic fragment of the fusion protein was detected (Fig. 3). However, its level was as low as the authentic phyB. Furthermore, the fragmentation might have occurred during the extraction. Even if the fragment existed in vivo, it would not contribute to the fluorescence. The immunoblot analysis suggests that the fragment resulted from proteolysis within the GFP portion, which would cause the loss of fluorescence. Likewise, it is unlikely that the fragment alone caused the phyB overexpression phenotypes, although it might have contributed to the phenotype to some extent.

**phyB-GFP Is Localized to the Nucleus**

Although associations of phyA with various organelles have been reported, only a small portion of the total cellular phyA was recovered in those cases (Pratt, 1994). It is also known that the Pfr form of phyA tends to associate with particulate material under certain cell extraction conditions (Quail, 1983). Thus, it had remained obscure whether phytochrome indeed resides within organelles in vivo. More recently, we have shown that COOH-terminal fragments of phyB fused to GUS are localized to the nucleus (Sakamoto and Nagatani, 1996). Furthermore, a substantial amount of endogenous phyB has been detected in isolated nuclei. On the basis of these findings, we had tentatively proposed that phyB translocates to the nucleus to mediate the light responses (Sakamoto and Nagatani, 1996; Nagatani, 1997). However, this observation could have been due to a cryptic nuclear localization signal that is exposed only in the context of the fusion protein. It is also difficult to exclude the possibility that phyB detected in the isolated nuclei might be due to contamination during specimen preparation and staining.

This study provides more dramatic evidence for phyB nuclear localization. Localization of the phyB-GFP fusion protein in the nucleus has been observed in intact live cells without any pretreatment (Figs. 4 and 5). The optical sectioning by a confocal microscope clearly indicates that the fluorescence is distributed inside the nucleus (Fig. 6). Furthermore, the fusion protein appears to be fully functional as a photoreceptor (see above), suggesting that the structure of phyB is preserved in the phyB-GFP fusion context. However, it should be noted here that phyB-GFP is overexpressed under control of the 35S promoter. Hence, there remains the possibility that ectopic expression contributes somewhat to the observed intracellular distribution. To address this question, we are now screening for transgenic lines with lower accumulation levels.

In the light, fluorescence of the phyB-GFP protein was observed mainly in the nuclei in all the cell types examined. Hence, the nucleus is likely to be the major site of the phyB action, although it remains possible that a minor fraction of phyB-GFP is present in other compartments of the cell and might contribute to the overall response. All the phenotypes observed in this study, such as the shorter hypocotyls, overall dwarfing in mature plants, and late flowering, can ultimately be explained by alteration in gene expression patterns. Hence, it is an intriguing possibility that phyb-GFP translocates to the nucleus to affect the transcription of target genes. Although phytochrome appears to have neither DNA-binding nor transactivation domains, it could interact with other factors that directly regulate transcription. PIF3, a nuclear-localized basic helix-loop-helix protein that binds to the COOH-terminal domain of phytochrome (Ni et al., 1998), is a potential candidate for such a factor.

**phyB-GFP Speckles**

Interestingly, the phyB-GFP fusion protein forms speckles in the nucleus. The sizes of the speckles are mostly <1 μm. However, the size varies even within one nucleus under continuous light (Fig. 6). The number of speckles per nucleus also varies. Interestingly, the size of the speckle gradually increases during the dark to light transition (Fig. 8). Conversely, the number of speckles decreases.

Speckled structures similar to one observed in this study have been reported in animal cells (Lamond and Earnshaw, 1998). Factors involved in the processing and transcription of RNA are found in those speckled structures in the nucleus. The promyelocytic leukemia (PML) nuclear body is another example of such a structure. However, the biological relevance of those structures remains unclear, although they may function as a repressor of transcription (Singer and Green, 1997; Lamond and Earnshaw, 1998). In plant cells, COP1, which is a negative regulator of plant photomorphogenesis or light responses (von Arnim and Deng, 1996), has been shown recently to form speckles in the nucleus (Ang et al., 1998). Hence, those speckles observed in plant cells might represent the site where the photoreceptor and other nuclear factors such as COP1 and PIF3 interact with each other to mediate light signals. Identification of proteins present in the phyB speckles is awaited.

It should be noted here that the phyB-GFP speckles could be due to an artifact caused by overaccumulation of the fusion protein at nonphysiological concentration. It is known that phytochrome in general tends to form aggregates in vitro (Quail, 1983). Hence, as with all studies using GFP fusion proteins, it is difficult at present to exclude this possibility. However, the light-dependent nature of the nuclear translocation supports the view that this distribution is indeed relevant to phytochrome function. Detailed analysis of transgenic lines that accumulate the fusion protein at lower levels is awaited. A search for mutations that abolish the speckles would greatly help to answer the question. For example, it has been shown in onion cells that GFP-HY5 fails to form speckles in the absence of COP1 expression (Ang et al., 1998). It will be of interest to investigate the effect of removing known light transduction components upon the subnuclear distribution of phyB.

**phyB-GFP Is Distributed throughout the Cell in Darkness**

In dark-grown seedlings, the phyB-GFP fusion protein ap-
peared to be distributed evenly throughout the cytoplasm (Fig. 7). This is consistent with the previous observation that phyB is not detected in the nuclei isolated from the dark-adapted rosette leaves (Sakamoto and Nagatani, 1996). In addition, a similar result has been obtained in dark-grown pea seedlings (Nagatani, A., unpublished observations). However, the detailed distribution of phyB-GFP could not be determined because of the resolution limit within the small Arabidopsis cells. At present, it is not clear whether the fusion protein is excluded completely from the nucleus in darkness. Although fluorescence was observed in the nucleus region, this could be due to fluorescence from the cytoplasm surrounding the nucleus. It would be helpful to isolate intact cells or protoplasts to determine the localization pattern in greater resolution.

To observe GFP fluorescence, the seedlings received relatively intense actinic blue light. In theory, this could alter the localization pattern of phyB-GFP. The apparently nuclear fluorescence observed in the dark-grown seedlings might be due to this effect. However, the distribution of phyB-GFP did not change significantly during the observation period (10–20 min). This is consistent with the relatively slow kinetics of phyB-GFP accumulation in the nucleus induced by continuous red light (Fig. 8). In addition, no change was observed in light-grown seedlings during the observation. Hence, the actinic blue light does not appear to disturb the distribution pattern at least during the observation period of 10–20 min.

Light-induced Nuclear Accumulation of phyB-GFP

Continuous red light induced accumulation of phyB-GFP in the nucleus (Fig. 8). Relatively slow kinetics of this process is consistent with the observation made in pea seedlings (Nagatani, A., unpublished observations). PhyB was not detected in the nuclei isolated from dark-grown pea seedlings. However, treatment of the seedlings with continuous red light induced nuclear localization of phyB. The level of nuclear phyB reached a plateau ~4 h after the onset of light treatment.

Phytochrome is known to regulate expression of various genes, of which CAB is the best characterized (Terzaghi and Cashmore, 1995). In Arabidopsis, the time course of the CAB gene induction by a red light pulse has been examined at high time resolution by using luciferase as a reporter (Millar et al., 1992; Millar and Kay, 1996; Anderson et al., 1997). These results indicate that the induction is multiphasic. An acute response occurs in Arabidopsis with the peak at 2 h after the pulse treatment. Subsequently, the expression oscillates under control of the biological clock. Namely, the levels fall to a trough at 6.5–8 h and peaks again at 15.5 h. A nalysis of the phyB mutant has demonstrated that phyB contributes to both the acute and clock-dependent responses (Anderson et al., 1997).

As shown in Fig. 8, it takes ~4 h for phyB-GFP to complete the translocation from the cytoplasm to the nucleus, which is significantly slower than the acute response of the CAB gene expression. This may indicate that the level of nuclear phyB attained 2 h after the onset of light might be sufficient to induce the acute response. It is also possible that phyB-GFP migrates more slowly than the authentic phyB. Alternatively, the signal transduction from phyB to the downstream components may take place in the cytoplasm in the early phase of the response. Interestingly, it seems here that the speckle formation is not required for the acute response of the CAB gene expression. As most, no speckles could be observed 2 h after the onset of red light treatment (Fig. 8).

It is more probable that the clock-dependent induction of the CAB gene expression is under control of nuclear-localized phyB. The extent of clock-dependent expression is maximal under continuous light. In such a condition, phyB-GFP is localized to the nucleus almost exclusively (Figs. 4–6 and 8). In addition, we have confirmed in pea seedlings that a pulse of red light can induce long-lasting accumulation of phyB in the nucleus (Nagatani, A., unpublished observations). It has been proposed that the circadian clock confines the ability of light to induce CAB expression (Kay and Millar, 1992). Hence, it is an intriguing possibility that phyB and components of the biological clock directly interact with each other within the nucleus.

Possible Functions of phyB in the Nucleus

Together with the previous report (Sakamoto and Nagatani, 1996), the present results provide compelling evidence that the nucleus is at least one of the sites of phyB action. In animal and yeast cells, many signal transduction factors are known to translocate to the nucleus upon receipt of the signal (Nagatani, 1998). For example, steroid hormone receptors are targeted to the nucleus upon binding of the hormonal ligands (Angelsdorff et al., 1995). Therefore, it is not surprising that phyB translocates to the nucleus upon light stimulation.

The det, cop, and fus mutants of Arabidopsis exhibit constitutive photomorphogenic phenotypes in darkness (von Arnim and Deng, 1996; Fankhauser and Chory, 1997). The DET1, COP1, COP9, and FUS6 proteins can be localized to the nucleus. It is especially interesting that the nuclear localization of COP1 is light dependent (von Arnim and Deng, 1996). Conversely, the hy5 mutant is impaired in the light signal transduction. The HY5 gene has been cloned recently (Oyama et al., 1997). The gene encodes a putative transcription factor which is constitutively localized to the nucleus (Chattopadhyay et al., 1998). The HY5 protein binds to the light-responsive promoters of the CAB and CHS genes (Ng et al., 1998; Chattopadhyay et al., 1998). More recently, a putative transcription factor, PIF3, has been identified as a phytochrome-interacting protein (Ni et al., 1998). Hence, it is an attractive possibility that phyB interacts with those proteins in the nucleus to modify transcription of the target genes.

The biological relevance of the phyB-GFP speckles in the nucleus is unknown as discussed above. The time course analysis in this study suggests that speckles are not required for the acute response of the CAB gene expression (Fig. 8). However, the speckled structure may contribute to long-term effects of phyB. It is intriguing here that GFP-COP1 fusion protein forms speckles in the nucleus (Ng et al., 1998). Furthermore, GFP-HY5 is recruited to the speckles if it is coexpressed with COP1. Hence, the speckles of COP1 may represent the site where nuclear factors interact with each other to mediate light
signals. In this connection, it would be particularly interesting to know whether phyB colocalizes with other factors in the nucleus.

Concluding Remarks

As discussed above, the present results suggest that phyB translocates to the nucleus upon light stimulation. In the nucleus, phyB may interact with other nuclear factors to transduce the light signal to alter transcription of target genes. However, the kinetics of the light-induced accumulation of phyB-GFP in the nucleus is clearly too slow to explain rapid phytochrome responses such as light-induced changes in intracellular Ca$^{2+}$ levels (Roux, 1994). Hence, phyB might be functioning not only in the nucleus but also in the cytoplasm. Alternatively, different molecular species of phytochromes may function at different sites within the cell. In this connection, it will be important to know whether other molecular species are localized to the nucleus.

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