p-Chlorophenoxyisobutyric Acid Impairs Auxin Response in Arabidopsis Root

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p-Chlorophenoxyisobutyric acid (PCIB) is known as a putative antiauxin and is widely used to inhibit auxin action, although the mechanism of PCIB-mediated inhibition of auxin action is not characterized very well at the molecular level. In the present work, we showed that PCIB inhibited BA::β-glucuronidase (GUS) expression induced by indole-3-acetic acid (IAA), 2,4-dichlorophenoxyacetic acid, and 1-naphthaleneacetic acid. PCIB also inhibited auxin-dependent DR5::GUS expression. RNA hybridization and quantitative reverse transcriptase-polymerase chain reaction analyses suggested that PCIB reduced auxin-induced accumulation of transcripts of Aux/IAA genes. In addition, PCIB relieved the reduction of GUS activity in HS::AXR3NT-GUS transgenic line in which auxin inhibits GUS activity by promoting degradation of the AXR3NT-GUS fusion protein. Physiological analysis revealed that PCIB inhibited lateral root production, gravitropic response of roots, and auxin-induced accumulation of transcripts of some specific genes with a lag period of 5 to 30 min (Abel and Theologis, 1996). Auxin-dependent degradation of Aux/IAA proteins is a key event for early auxin-dependent gene induction (Leyser, 2002). Aux/IAA proteins interact with auxin response factors (ARFs) that bind to auxin-responsive elements (AuxREs) in the auxin-responsive promoters (Guilfoyle, 1998). Aux/IAA proteins are short-lived proteins, and their stability is regulated by auxin through the ubiquitin-mediated protein degradation pathway (Leyser, 2002). In the absence of auxin, large amounts of Aux/IAA proteins are present, bind to ARFs, and prevent ARFs from activating transcription of auxin-induced genes from AuxREs. In the presence of auxin, degradation of Aux/IAA proteins is promoted, and then ARF proteins are released from Aux/IAA proteins and activate transcription of auxin-responsive genes (Tiwari et al., 2003). Although detailed molecular mechanism of the role of ARF and Aux/IAA proteins and their regulation has been revealed in these recent years, the mechanisms of the very early step of auxin signaling, i.e. how auxin is recognized by plant cell and how ubiquitin proteolysis system is activated, still remains unknown (Benfey, 2002).

The plant hormone auxin (indole-3-acetic acid [IAA]) plays an important role in every aspect of plant growth and development (Thimann, 1977; Davies, 1995). Despite its physiological significance, the molecular mechanism of auxin action is not fully understood yet. One of the earliest events in the auxin action involves the changing of expression pattern of some specific genes with a lag period of 5 to 30 min (Abel and Theologis, 1996). Auxin-dependent degradation of Aux/IAA proteins is a key event for early auxin-dependent gene induction (Leyser, 2002). Aux/IAA proteins interact with auxin response factors (ARFs) that bind to auxin-responsive elements (AuxREs) in the auxin-responsive promoters (Guilfoyle, 1998). Aux/IAA proteins are short-lived proteins, and their stability is regulated by auxin through the ubiquitin-mediated protein degradation pathway (Leyser, 2002). In the absence of auxin, large amounts of Aux/IAA proteins are present, bind to ARFs, and prevent ARFs from activating transcription of auxin-induced genes from AuxREs. In the presence of auxin, degradation of Aux/IAA proteins is promoted, and then ARF proteins are released from Aux/IAA proteins and activate transcription of auxin-responsive genes (Tiwari et al., 2003). Although detailed molecular mechanism of the role of ARF and Aux/IAA proteins and their regulation has been revealed in these recent years, the mechanisms of the very early step of auxin signaling, i.e. how auxin is recognized by plant cell and how ubiquitin proteolysis system is activated, still remains unknown (Benfey, 2002).

p-Chlorophenoxyisobutyric acid (PCIB), also called α-(4-chlorophenoxy)isobutyric acid, 2-(p-chlorophenoxy)-2-methylpropionic acid, or clofibric acid, has been most widely used to inhibit auxin action (e.g. Kim et al., 2000; Xie et al., 2000). Because of the structural similarity of PCIB with a synthetic auxin 4-chlorophenoxyacetic acid (Jönsson, 1961) and its competitive inhibition character to auxin-induced physiological responses, it is believed that PCIB inhibits the auxin action by competing with auxin at the binding site of the auxin receptor (MacRae and Bonner, 1953). If this hypothesis is true, PCIB might be a useful tool to elucidate the mechanism of auxin perception and signal transduction and their role in plant growth and development. However, the detailed molecular mechanism of how PCIB inhibits auxin action is almost unknown. Although there are few reports demonstrating that PCIB inhibits auxin-regulated gene expression (Okamoto et al., 1995;
Klotz and Lagrimini, 1996), the effects of PCIB have never been thoroughly investigated at the molecular level.

We previously designed BA::β-glucuronidase (GUS) transgenic Arabidopsis containing a GUS reporter gene under the control of the auxin-response domains (AuxRDs) of the promoter from the best-characterized primary auxin-responsive gene, pea (Pisum sativum) PS-IAA4/5 (Oono et al., 1998). GUS activity can be observed in the root elongation zone of the BA::GUS transgenic line in response to auxin. Other plant hormones and auxin antagonists did not induce GUS activity in the roots, indicating that the response is highly specific to auxin. This specific response of the BA::GUS transgenic line against auxin can be exploited to find genetic mutations or biologically active compounds that affect the early auxin gene expression (Oono et al., 1998; 2002; Hayashi et al., 2001). Recently, we found that PCIB inhibits the early auxin gene expression during the examination of auxin-related compounds with BA::GUS line. In this paper, we report characterization of PCIB as an auxin response inhibitor and elucidation of the physiological effects of PCIB on Arabidopsis root growth. We show that PCIB inhibited early auxin gene expression and auxin-dependent protein degradation. Our physiological studies reveal that PCIB inhibited gravitropic response, primary root elongation, and auxin-induced lateral root formation in Arabidopsis roots.

RESULTS

PCIB Impairs Auxin-Induced Gene Expression

In the preliminary screen, we examined several auxin-related compounds to investigate their effects on auxin-regulated gene expression using the BA::GUS line. Twenty micromolar of N-(1-naphthyl) phthalamic acid (NPA) and 2,3,5-triiodobenzoic acid (TIBA), auxin efflux inhibitors (Lomax et al., 1995), and 2,4,6-trichlorophenoxyacetic acid (2,4,6-T), an inactive or weak auxin (Jönsson, 1961), showed no effect on GUS activity induced by 0.1 μM IAA, whereas 20 μM PCIB effectively inhibited this activity (Fig. 1A). The inhibitory effect of 20 μM PCIB could be overcome with increasing concentrations of IAA (Fig. 1A). Minimum concentration of PCIB to inhibit 0.1 μM IAA-induced GUS activity was between 0.5 and 2 μM (data not shown). Yokonolide B (YkB) isolated from Streptomyces diastatocromogenes has been recently identified as a new type of inhibitor for auxin signal transduction and strongly inhibited IAA-induced BA::GUS expression (Fig. 1A; Hayashi et al., 2001, 2003). In contrast to PCIB, this expression could not be restored with an application of 10 μM IAA (Fig. 1A).

We previously showed that two putative auxin influx inhibitors, Chromosaponin I (CSI) and 1-naphthoxyacetic acid (1-NOA), inhibited BA::GUS expression induced by IAA (Rahman et al., 2002). To investigate whether PCIB inhibition of IAA-induced BA::GUS expression is also a consequence of inhibition of auxin influx or not, we compared the effect of PCIB with the effects of inhibitors for auxin influx as well as efflux on 1-naphthaleneacetic acid (NAA)-induced BA::GUS expression (Fig. 1B). In contrast to IAA, NAA can enter the cells by bypassing the requirement an auxin influx carrier (Delbarre et al., 1996) and NAA-induced BA::GUS expression is not inhibited by either CSI or 1-NOA (Rahman et al., 2002). Figure 1B showed that 20 μM PCIB partially and 50 μM PCIB completely blocked 1 μM NAA-induced BA::GUS expression. Consistent with the previous report (Rahman et al., 2002), 1-NOA even at 100 μM failed to do so, clearly suggesting that the inhibitory effect of PCIB is not due to the blocking of auxin influx. Interestingly, auxin efflux inhibitor TIBA but not NPA inhibited NAA-induced BA::GUS expression at 50 μM (Fig. 1B), although 20 μM TIBA failed to inhibit 0.1 μM IAA- and 1 μM NAA-induced BA::GUS expression (Fig. 1, A and B), suggesting TIBA may have auxin antagonistic activity at high concentration. Twenty micromolar PCIB also blocked 1 μM 2,4-dichlorophenoxyacetic acid (2,4-D)-induced BA::GUS expression (data not shown).

We also examined several compounds, 2,4-dichloranisole (DCA), maleic hydrazide, trans-cinnamic acid, 3-indole-3-butyric acid (3-IBA), and 4,4,4-trifluoro-3-indole-3-butyric acid (TFIBA) that were reported or proposed to have an auxin antagonistic activity (van Overbeek et al., 1951; Leopold and Klein, 1952; MacRae and Bonner, 1953; Katayama et al., 1995; Tomic et al., 1998), and 1-aminocyclopropane-1-carboxylic acid, a precursor of ethylene (McKeon et al., 1995). Among these compounds, 50 μM TFIBA effectively and 100 μM trans-cinnamic acid slightly inhibited BA::GUS expression induced by 1 μM NAA (data not shown). Other compounds did not show any inhibitory effect on BA::GUS expression induced by 1 μM NAA at the concentration up to 100 μM (data not shown).

To analyze the effects of PCIB on the steady-state level of transcripts of the marker gene regulated by auxin-responsive promoter as well as that of endogenous early auxin genes, 5-d-old seedlings of the BA::mgfp5-ER line (Aspuria et al., 2002) were treated in liquid germination media (GM) containing IAA and/or PCIB for 2 h, and total RNA was extracted. Twenty and 10 μg of total RNA from shoots and roots, respectively, was subjected to hybridization analysis using DNA fragments of mgFP5-ER, IAA1, and IAAt1 (Abel et al., 1995) as probes (Fig. 2). IAA (0.2 μM) increased the accumulation of mRNA for auxin-responsive genes quite effectively in roots and slightly in shoots. Application of 20 μM PCIB reduced the mRNA levels of the BA::mgfp5-ER gene in the presence as well as in the absence of 0.2 μM IAA. The effect of PCIB was much clearer in roots than in
shoots. The similar inhibitory effect of PCIB could be observed not only in the mRNA level of transgene with synthetic promoter but also in the mRNA level of the endogenous auxin-inducible genes, suggesting that PCIB impairs early auxin gene expression at transcriptional level (Fig. 2).

IAA responses in the presence or absence of PCIB were examined by using another auxin-responsive reporter line, DR5::GUS. The DR5::GUS construct consists of seven tandem repeats of an auxin-responsive TGTCTC element, a minimal 35S CaMV promoter, and a GUS coding region. Exogeneously applied auxin induces GUS accumulation strongly in all cells in the primary root meristem of the transgenic DR5::GUS Arabidopsis line (Ulmasov et al., 1997; Sabatini et al., 1999). Therefore, auxin-induced GUS activity in DR5::GUS line can be analyzed quantitatively with higher sensitivity than in BA::GUS line. Application of 20 μM PCIB in the IAA incubation media effectively inhibited auxin-induced GUS activity in roots of DR5::GUS transgenic line (Fig. 3A). Quantitative GUS analysis using whole seedling...
lings also showed that PCIB inhibited IAA-induced GUS activity, although the detectable level of auxin response was obtained at much higher concentrations of IAA than it was required in histochemical experiments (Fig. 3B). This may be because root tissue is much more sensitive to auxin than the aerial tissues that provide the majority of proteins in quantitative GUS analysis.

We also examined pIB-1.6TC transgenic line, which contains a GUS reporter gene with 1.6-kb upstream sequences of cytokinin-responsive gene ARR5 (D’Agostino et al., 2000). GUS expression in 5-d-old pIB-1.6TC seedlings treated with various concentrations of 6-benzyladenine (6-BA) and with or without 20 μM PCIB for 6 h is shown in Figure 3. The intensity of GUS staining was not significantly different with 20 μM PCIB treatment or without (Fig. 3A). The quantitative GUS assay using whole plant extract showed that PCIB had little or no effect on 6-BA-induced GUS activity in pIB-1.6TC transgenic line (Fig. 3B).

Cycloheximide (CHX), a well-characterized protein synthesis inhibitor, is known to stimulate mRNA expression of the primary auxin-responsive genes such as the Aux/IAA genes (Abel et al., 1995). In contrast to auxin, the mRNA induction by CHX is hypothetically caused by the inhibition of synthesis of short-lived transcription repressor proteins (Ballas et al., 1993). To examine whether PCIB affects CHX-induced mRNA induction, real-time reverse transcriptase (RT)-PCR with IAA11 primers were performed. The results clearly showed that PCIB inhibited IAA-induced but not CHX-induced mRNA expression of IAA11 gene, suggesting that PCIB specifically affects the auxin-signaling pathway (Fig. 4).

PCIB Prevents Auxin-Mediated Aux/IAA Protein Degradation

Recently, it was proposed that auxin promotes degradation of Aux/IAA proteins through ubiquitin-proteasome pathway and thereby induces the early auxin response (Zenser et al., 2001; Ramos et al., 2001; Gray et al., 2001; Tiwari et al., 2001). To address PCIB effect on ubiquitin-proteasome-dependent Aux/IAA degradation, we examined GUS activity in the HS::AXR3NT-GUS transgenic Arabidopsis line, which produces GUS protein fused translationally with the N-terminal half of AXR3. Because the N-terminal half of AXR3 is a target of ubiquitin-proteasome-dependent protein degradation system, the GUS activity in the HS::AXR3NT-GUS transgenic line is reduced by incubating with auxin after heat treatment (Gray et al., 2001). HS::AXR3NT-GUS seedlings treated with 1 μM 2,4-D showed decreased GUS activity in root compared with the control seedlings (Fig. 5, A and B). Our results showed stronger GUS activity than those reported by Gray et al. (2001), and more time was required for detecting the degradation of GUS activity. This may be due to the difference of the method to give the heat treatment, a most critical step for controlling basal GUS activity. The degradation of GUS activity by 1 μM 2,4-D was partially relieved by the addition of 50 μM PCIB (Fig. 5, A and B). To confirm that PCIB-induced increase in GUS activity in the presence of 2,4-D is related to the protein stability of the AXR3NT-GUS protein, we examined two control lines, HS::axr3-1NT-GUS, in which the mutation in domain II of AXR3 resulting in an increased stability of the protein, and HS::GUS transgenic lines (Gray et al., 2001). Neither 2,4-D nor PCIB significantly affected GUS activity in these transgenic lines (Fig. 5, A and B). These results suggested that PCIB partially inhibits the auxin-mediated degradation of Aux/IAA protein through ubiquitin-proteasome pathway.
Inhibition Kinetics of PCIB

It has been hypothesized that PCIB inhibits the auxin action by competing with auxin at the binding site of the auxin receptor (MacRae and Bonner, 1953). There are several auxin-binding proteins identified in different subcellular locations (Jones, 1994). If auxin and PCIB simply compete at the extracellular binding site on the cell surface, it is logical to assume that PCIB would inhibit auxin action immediately. To assess this possibility, time course of IAA-induced mRNA level in the presence or absence of PCIB was examined by real-time RT-PCR (Fig. 6). It has been reported that IAA induces mRNA of primary auxin responsible genes very rapidly (Theologis et al., 1985; Koshiba et al., 1995). The response of most Aux/IAA members in Arabidopsis occurs within 5 to 30 min (Abel et al., 1995). Consistent with the previous reports, our results also showed that level of IAA11 and DR5::GUS mRNA were increased by 1 μM IAA within 30 min of incubation (Fig. 6). However, the inhibitory effect of PCIB could not be clearly detected until 1 h after starting the incubation. The presence of the lag period to see the PCIB effect suggests that IAA and PCIB do not simply compete at their extracellular binding site.

Physiological Effects of PCIB on Arabidopsis Root Growth

The results shown above suggested that PCIB inhibited an early step of auxin signaling and prompted us to examine the effects of PCIB at physiological level. Because we found severe physiological effects of PCIB in root but not in aerial part of Arabidopsis seedlings, we focused our study on analysis of root growth.

Figure 7A shows the density of lateral roots (the number of lateral root per centimeter of primary root length) in the seedlings treated for 5 d in the medium containing IAA and/or PCIB. PCIB inhibited exogenous IAA-induced emergence of lateral roots. Higher concentration of PCIB was required to inhibit lateral root emergence that was induced by higher concentration of IAA. During the early lateral root initiation,
auxin is involved in determination of the position and frequency of lateral root, initiation of cell division in pericycle, and maintenance of active lateral root meristem (Celenza et al., 1995; Himanen et al., 2002). To determine which stage PCIB inhibits, we examined two transgenic lines harboring the DR5::GUS and the CycB1;1::GUS genes. The DR5::GUS gene is frequently used to visualize auxin accumulation and auxin response at cellular resolution (Sabatini et al., 1999; Rashotte et al., 2001; Friml et al., 2002). On the other hand, the transcription from the CycB1;1 promoter is closely associated with the G2 and M phases of cell division. The CycB1;1::GUS gene can be used as an indicator for early mitotic event associated with lateral root initiation (Ferreira et al., 1994). Both markers are expressed in the dividing pericycle cells of lateral root root initiation and indicate the initial cells where lateral roots initiation occurs (Ferreira et al., 1994; Casimiro et al., 2001; A. Rahman and Y. Oono, unpublished data). Figure 7, B and C, show the number of the lateral root sites stained with GUS in DR5::GUS and CycB1;1::GUS lines, respectively. In both markers, the number of GUS-staining sites was significantly reduced with increasing concentration of PCIB. These results suggested that PCIB inhibited auxin response and subsequent initiation of cell division in lateral root initiation site.

Because auxin was suggested to be a signal for controlling gravitropic response (Muday, 2001), we examined gravitropic response of Arabidopsis roots in the presence of various concentrations of PCIB. Five-day-old seedlings grown on GM medium were transferred to fresh GM medium containing various concentrations of PCIB and continued to grow in the dark for 3 d with changing the gravity orientation through 135°. Figure 8 shows the distributions of root growth directions of the seedlings. PCIB significantly reduced gravitropic response of Arabidopsis roots from 1 µM and completely inhibited gravitropic response at 20 µM.

Figure 9 shows the length of the primary root of the seedlings treated for 5 d on GM medium containing various concentrations of PCIB. In contrast to counteractive effect of PCIB in lateral root emergence shown in Figure 7A, PCIB inhibited primary root elongation by itself and failed to counteract the effect of 0.1 µM IAA on root elongation (Fig. 9A). We also investigated dose response of IAA in the presence and absence of PCIB and found IAA did not counteract the inhibition of primary root elongation caused by 5 µM PCIB (Fig. 9B). Regarding the relative primary root elongation, 20 µM PCIB reduced primary root elongation to 33% of control in the absence of IAA but 55% of control in the presence of 0.1 µM IAA (Fig. 9A, inset). Furthermore, 0.1 µM IAA reduced primary root elongation to 40% of control in the absence of PCIB but 64% in the presence of 5 µM PCIB (Fig. 9B, inset). These data suggested that primary roots were more resistant to IAA or PCIB in the presence of PCIB or IAA, respectively. The mutants axr1-3, axr2-1, and tir1-1, which are defective in auxin signaling (Leyser et al., 1993; Ruegger et al., 1998; Nagpal et al., 2000; del Pozo et al., 2002), showed partial resistance to PCIB compared with wild type on primary root elongation (Fig. 9C).

On the other hand, the mutants defective in auxin influx and efflux carrier, aux1-7, aux1-22, ein1-1, ein2-1 (Guzman and Ecker, 1990), and ethylene insensitive mutant ein2-1 (Guzman and Ecker, 1990), showed the same response as wild type, suggesting that auxin transport and ethylene are not but auxin signaling is involved in the PCIB-mediated inhibition of the primary root elongation (Fig. 9C).

DISCUSSION

PCIB Inhibits Auxin Response

More than 50 years ago, it was proposed that PCIB acts as an antiauxin (Burström, 1950). PCIB inhibits many auxin-induced physiological effects. For example, PCIB antagonizes IAA-induced inhibition of wheat (Triticum aestivum) root growth (Burström, 1950) and competitively inhibits IAA and 2,4-D induced growth of Azema sp. coleoptile (MacRae and Bonner, 1953).

Because of its structure and inhibition kinetics, PCIB has been believed to inhibit auxin action by competing with auxin at its binding site of the receptor (MacRae and Bonner, 1953). On the other hand, the effect of PCIB was often attributed to the inhibition of auxin transport (Katekar and Geissler, 1980; Tsai and Artea, 1984), the regulation of auxin metabolism (Frenkel and Haard, 1973), or nonspecific toxic effect (Yip and Yang, 1986). Although the effects of PCIB presented in this study such as inhibition of lateral root development, loss of gravitropic re-
sponse, and inhibition of primary root growth are also caused by auxin influx or efflux inhibitors (Fujita and Syono 1996; Luschnig et al., 1998; Casimiro et al., 2001; Malamy and Ryan, 2001; Rahman et al., 2001, 2002), some of the PCIB effects are crucially different from the effects of auxin influx and efflux inhibitors. First, PCIB inhibited not only IAA-induced, but also NAA- and 2,4-D-induced BA::GUS expression. NAA and 2,4-D do not require influx and efflux carrier for moving in and out of the cells, respectively (Delbarre et al., 1996). Second, root growth in Arabidopsis mutants aux1-7, aux1-22 and eir1-1, which are defective in auxin influx or efflux carriers and resistant to auxin transport inhibitors such as NPA and TIBA (Morris, 2000; Luschnig et al., 1998), were not resistant to PCIB. In addition, auxin transport activity in the Arabidopsis inflorescence axis was inhibited by 9-hydroxyfluorene-9-carboxylic acid, NPA, and TIBA but not by PCIB (Okada et al., 1991). PCIB did not affect the auxin influx or efflux during the auxin transport in suspension cultured tobacco cells (Imhoff et al., 2000). All these evidences also support our observation that PCIB does not function as auxin transport inhibitor.

Another explanation for the PCIB-induced inhibition of early auxin gene expression could be that PCIB enhances destruction of auxin. In fact, Frenkel and Haard (1973) reported that PCIB decreased the

Figure 5. PCIB impairs auxin-dependent degradation of Aux/IAA protein stability. A, Seven-day-old HS::AXR3NT-GUS, HS::axr3-1NT-GUS, and HS::GUS transgenic seedlings were incubated in GM liquid media for 120 m at 37°C followed by transferring seedlings into GM liquid media at 23°C. After 20 m, seedlings were transferred into new GM liquid media containing 2,4-D and/or PCIB and incubated for another 120 m. The seedlings were then incubated in GUS-staining buffer containing X-gluc for 18 h for the HS::AXR3NT-GUS line or 1 h for the HS::axr3-1NT-GUS and HS::GUS transgenic lines. B, Excised roots from 7-d-old seedlings were treated with 2,4-D and/or PCIB as in A. Relative GUS activity was measured quantitatively and expressed as percentage of the mock incubation.
have several lines of evidence to suggest that PCIB-mediated inhibition is not simply due to the nonspecific toxic effects. First, applying as little as 10 μM IAA counteracts PCIB-induced inhibition of early auxin gene expression and lateral root emergence. Second, GUS activity, which was observed by non-auxin-specific cytokinin inducible promoter, was not extensively inhibited in the presence of PCIB. Third, 2,4-D-induced reduction of GUS activity in HS::AXR3NT-GUS plants was counteracted by PCIB treatment. All of these results suggest that PCIB-mediated effects that we observed are not a toxic effect but rather support our idea that PCIB specifically inhibits auxin response in Arabidopsis roots.

Figure 6. Time course of the induction of IAA11 and DR5::GUS gene expression by 1 μM IAA in the presence (white circle) or absence (black circle) of 20 μM PCIB. Seven-day-old DR5::GUS transgenic seedlings were treated for the time indicated. Three independent sets of experiments were done and the averages of the values relative to the expression level of the zero time control are shown. Bars represent ± s.e of the average.

Figure 7. Effects of different concentrations of IAA and PCIB on lateral root production. A, Five-day-old seedlings grown vertically were transferred onto GM media containing various concentrations of IAA and/or PCIB and incubated vertically for additional 5 d. The number of emerging lateral roots (average ± s.e) per centimeter of primary root was calculated in each seedling (n = 30). B and C, Five-day-old seedlings grown vertically were transferred onto GM media containing various concentrations of PCIB and incubated vertically for additional 5 d. DR5::GUS (B) or CycB1;1::GUS (C) seedlings were subjected to histochemical GUS staining. Number of GUS-staining sites in pericycle cells was counted as the number of lateral root initiation site under the microscope. Bars represent ± s.e of the average (n = 10). Number of GUS-staining sites in the seedling before transfer is presented as D0.
PCIB Disturbs Auxin Signaling at the Upstream of Auxin-Mediated Aux/IAA Protein Degradation

It is well known that expression of the primary auxin-induced genes is enhanced by not only auxin but also CHX, a protein synthesis inhibitor (Theologis et al., 1985; Abel et al., 1995). This CHX-mediated increase of mRNA expression is explained by decreasing short-lived transcriptional repressor in a consequence of CHX-directed inhibition of protein synthesis (Ballas et al., 1993). Recent evidence suggests that Aux/IAA proteins are the short-lived repressors that regulate expression of primary auxin-induced genes including Aux/IAA genes themselves, and auxin promotes the degradation of Aux/IAA proteins by ubiquitin-mediated process (Leyser, 2002). Thus, although both auxin and CHX induce primary auxin-induced genes including Aux/IAA genes themselves, and auxin promotes the degradation of Aux/IAA proteins by ubiquitin-mediated process (Leyser, 2002). Thus, although both auxin and CHX induce primary auxin-responsive genes by reducing amount of repressor, their acting mechanisms are quite different. Our observation that PCIB inhibited IAA-induced but not CHX-induced expression of IAA11 gene suggested that PCIB specifically acts on auxin signaling.

Several proteins, such as AXR1 and TIR1, which are involved in the ubiquitin-related Aux/IAA degradation and auxin response, are well characterized (Leyser 2002). The functional defects of these proteins cause increasing of the stability of Aux/IAA proteins and reduction in auxin response. The domain II in the N-terminal half of Aux/IAA protein is responsible for auxin-dependent ubiquitin-mediated degradation (Leyser, 2002). The mutations in domain II of Aux/IAA proteins increase the protein stability dramatically and result in a loss of auxin response. The transgenic plants expressing the AXR3NT-GUS, the fusion protein of N-terminal of AXR3 and GUS, are useful
tool to examine the stability of Aux/IAA proteins. Increasing GUS activity has been reported in the axr1 and tir1-1 mutants containing the HS::AXR3NT-GUS gene (Gray et al., 2001). In our experiments, PCIB counteracted the 2,4-D-induced reduction of GUS activity in the HS::AXR3NT-GUS line, suggesting that PCIB acts through stabilizing the Aux/IAA proteins. Furthermore, PCIB did not affect GUS activity of the axr3-1NT-GUS whose stability is no longer regulated by auxin because of the mutation in domain II. These results together with the fact that PCIB did not inhibit CHX-induced IAA11 gene expression suggest that PCIB acts negatively on auxin-signaling pathway upstream of ubiquitin-mediated degradation of Aux/IAA proteins.

The hypothesized PCIB action is that PCIB binds to the auxin receptor and inhibits the auxin action (MacRae and Bonner, 1953). If the receptor was present on the cell surface and compounds could easily access to the receptor without any regulation, inhibitory effect of PCIB could be observed at the same time when auxin effects are detected. However, the presence of a 1-h lag-period to detect the PCIB action indicates that PCIB does not simply compete with auxin in the binding site on the cell surface. This time lag could be attributed by an explanation that the auxin-binding site inhibited by PCIB may be located inside the cell and lag-period may be due to the time required for PCIB entering into the cells. Although there are some evidences that a functional extracellular auxin-binding protein perceives auxin signal (Steffens et al., 2001), the presence of auxin uptake carriers such as AUX1 and altered physiology of aux1 mutants indicate that auxin signal for root growth is perceived intracellularly. The fact that the lack of AUX1 affected the expression level of Aux/IAA genes and BA::GUS transgene (Abel et al., 1995; Oono et al., 1998) strongly suggests that expression level of auxin-regulated gene is also controlled by intracellular auxin. Many functional auxin-binding sites were identified inside the cells (Jones, 1994; Kim et al., 2001). These binding sites may be the targets of PCIB. Alternative explanation for the lag-period of PCIB action could be due to that PCIB does not interact directly to functional auxin-receptor for early auxin gene expression. However, several biochemical experiments clearly showed that PCIB interferes with auxin binding to auxin-binding proteins (e.g. Ray et al., 1977; Moloney and Pilet, 1981; Prasad and Jones, 1991; Sugaya et al., 2000). Further experiments such as molecular genetic analysis will reveal the precise mechanism of PCIB action.

PCIB Influences Arabidopsis Root Growth

Several Arabidopsis mutants and cosuppression transgenic lines such as axr1, tir1, and CSN5G, which are defective in auxin signaling because of loss of the components of ubiquitin-related protein degradation pathway, exhibit fewer lateral roots and reduced sensitivity to the gravity (Schwechheimer et al., 2001). Reduced lateral root formation and less sensitivity to the gravity are also reported in other auxin-signaling mutant, ask1-1 and ecr1-1, respectively (Gray et al., 1999; Dharmasiri et al., 2003). The similar characters were also observed in PCIB-treated seedlings, supporting our idea that PCIB impairs the cellular auxin response and thereby inhibits the initiation of lateral roots and gravity sensing. In contrast, the elongation of primary roots was inhibited by both auxin and PCIB, and these two compounds did not counteract each other. Burström (1950) also reported that 10 μM PCIB reduced the rate of wheat root elongation. Growth of primary roots is consequences of coordinated cell division, cell differentiation, and cell elongation of many different types of cells, all of which are regulated by auxin. Exogenously applied PCIB impairs auxin response but probably would not equally affect to each auxin response in various cells, resulting in disruption of normal cellular organization and inhibition of root growth. It is interesting that the auxin-signaling mutants axr1-3, axr2-1, and tir1-1, but not the auxin influx or efflux mutants aux1s or eir1-1, showed PCIB resistance. This implies that PCIB targets auxin signal pathway to inhibit primary root growth. Evans et al. (1994) demonstrated that axr1 mutants are defective in both promotion and inhibition of primary root growth by low and high concentrations of auxin, respectively. Thus, it might be reasonable that inhibition of auxin response by PCIB is also ineffective in these auxin-signaling mutants. Although further characterization is required, we are speculating that PCIB-mediated inhibition of primary root growth is due to the inhibition of cellular auxin response by PCIB.

Auxin Action Inhibitors Could Be Useful Tools for Molecular Dissection of Auxin Signaling

Application of exogenous auxin transport inhibitors or the mutants having altered response to these inhibitors provide useful information in understanding the molecular mechanism of auxin transport and its role on plant development (Muday and Delong, 2001). On the other hand, although several classes of compounds such as classical antiauxins, xyloglucan oligosaccharides, anion-channel blockers, and phospholipase A₂ inhibitors have been shown to have an inhibitory effect on auxin action (Jönsson, 1961; McDougall and Fry, 1989; Scherer and Arnold, 1997; Thomine et al., 1997), they were less studied at molecular level, and their mode of action was not well understood until lately. Recently, Mauro et al. (2002) reported that oligogalacturonides inhibit the induction of auxin-responsive genes in tobacco. Unlike PCIB, the oligogalacturonides inhibit secondary auxin-responsive genes but not primary auxin-responsive genes, suggesting that oligogalactu-
ronides act on relatively late auxin response. Yokonolide A and YkB 22-membered spiroketal-macrolides were isolated from S. diastatocromagens as inhibitors of early auxin gene expression by screening against BA::GUS expression (Hayashi et al. 2001). The inhibitory effect of YkB on BA::GUS expression is quite strong compared with PCIB (Fig. 1). Further characterization of YkB suggested that YkB blocks degradation of Aux/IAA proteins (Hayashi et al., 2003). PD098059 and U0126, which are mitogen-activated protein kinase inhibitors, are also reported to inhibit BA::GUS expression (Mockaitis and Howell, 2000). More detailed characterization of these auxin response inhibitors including PCIB by combining the pharmacological approaches and molecular genetics, would provide new insights to elucidate the mechanism of auxin signaling.

MATERIALS AND METHODS

Plant Materials and Growth Conditions

All transgenic and mutants lines except the 35S::GUS and pIB-1.6TC lines were derived from Arabidopsis Columbia (Col-0) ecotype. The 35S::GUS and pIB-1.6TC lines were derived from C24 and Wassilewskija ecotypes, respectively (Ferreira et al., 1994; D’Agostino et al., 2000). Transgenic lines containing BA::GUS (BA3), 35S::mGFP5-ER (line 23.1.1), and DR5::GUS are described by Oono et al. (1998), Aspuria et al. (2002), and Umasov et al. (1997). Respectively. BA::GUS, 35S::AXR3NT-GUS, and 35S::axr3-1NT-GUS are described by Gray et al. (2001). Surface-sterilized seeds were plated on GM (one-half-strength Murashige and Skoog salts, 1% [w/v] sodium phosphate, pH 7.0, 10 mM EDTA, 0.5 mM K₃Fe(CN)₆, 0.5 mM K₃Fe(CN)₆, and 0.1% [v/v] Triton X-100) containing 1 mM X-gluc and incubated at 37°C for 15 to 22 h unless indicated. Quantitative GUS fluorometric assay was performed as described by stomp (1992). Protein concentration was determined with a protein assay reagent (Bio-Rad Laboratories, Hercules, CA) and bovine serum albumin (Sigma B-4267) as standard. For RNA hybridization analysis, total RNA was isolated from shoots and roots by the procedure of Theologis et al. (1985), size-fractionated with a 1% (w/v) agarose gel containing formaldehyde, and transferred onto a nitrocellulose membrane (Schleicher & Schuell). Hybridization was performed by using 3²P-labeled probes according to the manual provided by Schleicher & Schuell. The blots were then exposed to x-ray film. DNA fragments for the probes were prepared by digesting plasmids containing cDNA of IAAl and IAA11 (Asbel et al., 1995) with EcoRI/HindIII and EcoRI/Ncol, respectively, and 35S::mGFP5-ER (Aspura et al., 2002) with EcoRI/SacI. The DNA fragment for 175 RNA is described by Zarembinska and Theologis (1993). Digital images of autoradiograms were generated by scanning x-ray films and quantitative analysis of radioactivity of each band was performed with NIH image software.

For real-time RT-PCR, total RNA was isolated from 7- to 8-d-old seedlings using RNeasy Plant Mini Kit (Qiagen, Tokyo) with on-column DNase digestion according to the provided handbook. The real-time RT-PCR was performed on the LightCycler Instrument (Boehringer Mannheim) with the LightCycler-RNA Master SYBR Green I Kit (Boehringer Mannheim) according to the manufacturer’s protocol using the oligonucleotide primer sets (5’-CGCTTCATTTGACGCTTGGG and 5’-GTCGCTGAAAGCTCTTCGTT and 5’-CTCTCTGGACCTTCTGTC and 5’-TCTTCGACTTCCAGAGTTG for EFlaA4 gene). Each reaction (20 µL) was performed with 50 to 100 ng of total RNA, 0.3 µM each of forward and reverse primers, and 3.25 mM MgCl₂. The reactions were incubated at 61°C for 20 m for reverse transcription, followed by denaturation at 95°C for 30 s and PCR with 45 cycles of 1 s at 95°C, 5 s at 55°C, and 15 s at 72°C. The specificity of the PCR amplification was checked with a melting curve analysis program and agarose gel electrophoresis of PCR products. Relative amount of specific mRNA was estimated using the standard cDNA preparation of known size and molecular concentration and normalized to the EFlaA4 mRNA level.

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p-Chlorophenoxyisobutyric Acid Impairs Auxin Response in Arabidopsis Roots

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