A type III ACC synthase, ACS7, is involved in root gravitropism in Arabidopsis thaliana

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Abstract

Ethylene is an important plant hormone that regulates developmental processes in plants. The ethylene biosynthesis pathway is a highly regulated process at both the transcriptional and post-translational level. The transcriptional regulation of these ethylene biosynthesis genes is well known. However, post-translational modifications of the key ethylene biosynthesis enzyme 1-aminocyclopropane-1-carboxylate (ACC) synthase (ACS) are little understood. In vitro kinase assays were conducted on the type III ACS, AtACS7, fusion protein and peptides to determine whether the AtACS7 protein can be phosphorylated by calcium-dependent protein kinase (CDPK). AtACS7 was phosphorylated at Ser216, Thr296, and Ser299 by AtCDPK16 in vitro. To investigate further the function of the ACS7 gene in Arabidopsis, an acs7-1 loss-of-function mutant was isolated. The acs7-1 mutant exhibited less sensitivity to the inhibition of root gravitropism by treatment with the calcium chelator ethylene glycol tetraacetic acid (EGTA). Seedlings were treated with gradient concentrations of ACC. The results showed that a certain concentration of ethylene enhanced the gravity response. Moreover, the acs7-1 mutant was less sensitive to inhibition of the gravity response by treatment with the auxin polar transport inhibitor 1-naphthylphthalamic acid, but exogenous ACC application recovered root gravitropism. Altogether, the results indicate that AtACS7 is involved in root gravitropism in a calcium-dependent manner in Arabidopsis.

Key words: 14-3-3, ACS, calcium, CDPK, ethylene, phosphorylation, root gravitropism.

Introduction

Gravity is one of the most important environmental cues that control growth direction (Morita, 2010). Shoots generally grow upward (i.e. negative gravitropism) and roots grow downward (i.e. positive gravitropism). It is unknown how plants receive and respond to the gravity signal. One widely accepted theory is that starch-accumulating amyloplast movement along the gravity vector within gravity-sensing cells (statocytes) is a likely trigger of subsequent intracellular signalling (Morita, 2010). However, several studies have demonstrated that starch is important but not essential for gravity sensing (Strohm et al., 2011; Wolverton et al., 2011). The receptor that responds to gravity has not yet been identified. In the process of gravity perception, changes in the gravity vector are transduced into multiple intracellular signals [i.e. cytosolic pH, inositol 1,4,5-triphosphate (InsP₃), and cytosolic calcium concentration] (Fasano et al., 2001; Plieth et al., 2002; Perera et al., 2006). The plant hormone auxin was identified as one of these signalling molecules, and its redistribution is polar. Auxin has been thought to be involved in gravitropic responses, based on the Cholodny–Went theory. Auxins were the first group of plant growth regulatory substances to be discovered (Went, 1928). Indole-3-acetic acid (IAA) was the first native representative of the group that was identified. Auxins are known to be involved in the regulation
of basic growth processes, such as cell division and cell elongation at the tissue, organ, and whole-plant levels. The auxin molecule functions as a mobile signal between cells, tissues, and organs, and is involved in the spatial and temporal coordination of plant morphogenesis and responses to the environment in planta. Auxin is not only a mobile molecule; it also participates in downstream signalling. Many developmental processes appear to depend on the local asymmetric distribution of auxin molecules (Tanaka et al., 2006). These include embryo development and apical–basal axis formation in Arabidopsis thaliana (Friml et al., 2003), pattern formation and root development (Bilou et al., 2005), organ formation (Benkova et al., 2003), and phototropism and gravitropism (Blakeslee et al., 2004).

In addition to auxin, the phytohormone ethylene has been shown to be involved in the gravity response in plants (Madlung et al., 1999). In tomatoes, root penetration into the soil required cross-talk between ethylene and auxin (Santisree et al., 2011). A mutation of the Arabidopsis gene ARG1, which is involved in root gravitropism, showed resistance to ethylene but increased sensitivity to auxin (Sedbrook et al., 1998). The ARG gene encodes a membrane protein that is homologous with bacterial transporter proteins (Utsuno et al., 1998) as a polar auxin transporter (Chen et al., 1998). Another Arabidopsis mutant, clg1, showed resistance to ethylene in root gravitropism (Ferrari et al., 2000). In Arabidopsis, flavonoid accumulation by ethylene was found to be involved in root gravitropism (Buer et al., 2006). Additionally, the Arabidopsis mutant rh1 showed resistance to ethylene in roots and was found to be involved in root gravitropism (Fortunati et al., 2008). In six plant species tested, an ethylene biosynthesis inhibitor inhibited root curvature (Hoson et al., 1996). In Arabidopsis, a recent study showed that ethylene and gravity can affect root skewing and waving (Oliva and Dunand, 2007). The Arabidopsis mutant alh1 was shown to link the cross-talk between ethylene and auxin in the gravity response in roots (Vandenbussche et al., 2003). Moreover, acs6 and acs9 mutants showed defective hypocotyl length in the gravity response (Tsuschisaka et al., 2009). However, it is still controversial whether ethylene plays a positive or negative role in modulating the gravity response (Philosoph-Hadas et al., 1996; Buer et al., 2006).

Ethylene is a plant hormone involved in many plant growth and developmental processes, including seed germination, leaf and flower senescence and abscission, cell elongation, fruit ripening, nodulation, and responses to a wide variety of stressors (Crocker and Knight, 1908; Wang et al., 2002; Yoo et al., 2009). The biosynthesis of ethylene has been well documented in plants (Yang and Hoffman, 1984; McClellan and Chang, 2008). Ethylene is derived from the amino acid methionine, which is converted to S-adenosylmethionine (AdoMet) by S-adenosylmethionine synthetase. AdoMet is then converted to l-aminocyclopropane-1-carboxylic acid (ACC) and S′-deoxy-5′-methylthioadenosine (MTA) by the enzyme l-aminocyclopropane-1-carboxylase synthase (ACS) (Adams and Yang, 1979; Lin et al., 2009), which is the rate-limiting step in ethylene biosynthesis. MTA is recycled to methionine through the Yang cycle, which allows high rates of ethylene production without depleting endogenous methionine. ACC is converted to ethylene, CO₂, and cyanide by ACC oxidase (ACO). The cyanide produced by this reaction is detoxified into β-cyanoalanine by the enzyme β-cyanoalanine synthase, preventing toxicity to plants under conditions of high ethylene biosynthesis.

ACS proteins play an important role in the ethylene biosynthesis pathway. The enzyme catalyses the conversion of AdoMet to ACC, and this reaction requires pyridoxal-5′-phosphate (PLP) as a cofactor. In most plant species, ACS is encoded by a multigene family that is regulated by various environmental and developmental factors (i.e. cytokinin, auxin, root hair development, fruit ripening, wounding, and pathogens). Arabidopsis has eight genes that encode active ACS proteins and an additional gene that encodes a catalytically inactive enzyme, ACS1 (Liang et al., 1992). The Arabidopsis ACS2, ACS6, ACS7, and ACS9 genes can be induced by hypoxia (Peng et al., 2005). Zhang’s group found that ACS2, ACS6, ACS7, ACS8, and ACS11 were involved in Botrytis cinerea-induced ethylene biosynthesis in Arabidopsis (Li et al., 2012). Based on the C-terminal sequences, ACS proteins in Arabidopsis can be divided into three main types (Chae and Kieber, 2005). Type I proteins have an extended C-terminus that contains three conserved serine residues that are targets for phosphorylation by mitogen-activated protein kinase 6 (MPK6) (Liu and Zhang, 2004) and a conserved serine residue that is a phosphorylation site for calcium-dependent protein kinase (CDPK; Tatsuki and Mori, 2001; Sebastià et al., 2004). Type II proteins have a shorter C-terminus that has only the CDPK phosphorylation site. Type III proteins have a very short C-terminal extension that lacks both phosphorylation sites. ACS proteins can act as homo- or heterodimeric proteins, similar to other PLP-dependent enzymes, and their ability to form active heterodimers might act to increase the versatility of ethylene responses (Tsuschisaka and Theologis, 2004), which enhances the ability to regulate ethylene production after exposure to different developmental and environmental stimuli.

Details on ACS turnover have been derived from studies of Arabidopsis ethylene-overproducing (Eto) mutants (Chae and Kieber, 2005). The eto mutants produce 10- to 40-fold more ethylene in the dark compared with the wild-type (WT) seedlings, and adopt a triple-response morphology (i.e. a morphology that etiolated seedlings adopt in the presence of ethylene) in the absence of exogenous ethylene application (Guzman and Ecker, 1990; Kieber et al., 1993). The cloning of ETO1 revealed that it encodes an E3 ligase component, a BTB/TPR protein. ETO1 binds to type II ACS proteins but not type I or type III ACS proteins (Wang et al., 2004; Yoshida et al., 2005, 2006). The disruption of ETO1 resulted in increased stability of the type II ACS protein ACS5 (Chae et al., 2003) and consequently increased ethylene biosynthesis. The stability of ACS proteins is also regulated by protein phosphorylation. Treatment of tomato cells with the protein kinase inhibitors K-252a and staurosporine inhibited the elicitor-dependent induction of ACS and ethylene biosynthesis (Grosskopf et al., 1990; Felix et al., 1991) through a mechanism that most probably involves increased
turnover of the ACS protein (Spanu et al., 1994). In tomato cells, ACS2 was shown to be phosphorylated by CDPK from extracts of wounded tomato fruit (Tatsuki and Mori, 2001). The protein stability of ACS2 was found to be regulated by CDPK phosphorylation (Kamihoshihara et al., 2010). The target of CDPK phosphorylation was the conserved serine residue Ser460 at the C-terminal region of the ACS protein. A novel CDPK phosphorylation motif was identified in the C-terminal domain of type II ACS proteins (Sebastià et al., 2004). The current model proposes that the phosphorylation of type I and type II ACS proteins blocks the ability of ETO1/EOL proteins to bind and inhibit the ubiquitination of these ACS proteins for their degradation by the 26S proteasome (Wang et al., 2004; Chae and Kieber, 2005).

The regulation of ethylene biosynthesis and ACS stability are also controlled by mitogen-activated protein kinases (MAPKs). In tobacco, a stress-induced MAPK (SIPK) is involved in the response to different stressors, including pathogen- and ozone-induced ethylene biosynthesis. The closest homologue of SIPK in Arabidopsis is MPK6. Therefore, MPK6 was used in an in vitro kinase assay. The results showed that MPK6 can phosphorylate ACS2 and ACS6 in vitro, and transgenic plants that overexpress a phosphomimetic mutant of ACS6 showed increased ethylene production (Liu and Zhang, 2004). These results indicate that a pathway similar to the SIPK pathway in tobacco operates in Arabidopsis and that MPK6 phosphorylates ACS proteins, thereby decreasing their turnover and increasing ethylene biosynthesis after pathogen stress. A possible CDPK- and MPK6-regulated pathway was recently proposed by Ludwig et al. (2005). These findings highlight the complexity of phosphorylation-regulated signalling and ethylene biosynthesis in plants in response to different stressors.

Calcium is a ubiquitous secondary messenger in eukaryotic cells. In plants, intracellular calcium levels can modulate many growth and developmental processes, including plant hormones, light, gravity, and biotic and abiotic stress (Batistic and Kudla, 2012). Unlike most other ions, calcium does not freely diffuse within cells (Trewavas, 1999). Plants have multiple calcium stores, including in apoplasts, vacuoles, the nuclear envelope, the endoplasmic reticulum (ER), chloroplasts, and mitochondria. Different stimuli can trigger calcium efflux from specific organelles. After calcium is released, different calcium sensors that have an EF-hand motif that can specifically bind calcium [i.e. the EF-hand-containing proteins calmodulin (CaM) and CDPK] can recognize specific calcium signals in specific places and transduce them into downstream effects, including altered protein phosphorylation and gene expression patterns (Sanders et al., 1999, 2002; Dodd et al., 2010; Perochon et al., 2011; Liese and Romeois, 2012).

In Arabidopsis, the C-terminal domain in type I and type II ACS proteins can be phosphorylated by CDPK and MAPK. This phosphorylation of type I and type II ACS proteins blocks the ability of ETO1/EOL proteins to bind, thus inhibiting the ubiquitination of these ACS proteins and their degradation by the 26S proteasome. The degradation of type III ACS7 was recently found to also be mediated by the 26S proteasome (Lyzenga et al., 2012), but how ACS7 protein activity and stability are regulated is still unknown. The present study investigated whether the type III ACS protein ACS7 is phosphorylated by CDPK in vitro. An in vitro kinase assay was conducted to determine the phosphorylation of ACS7 by CDPK and identify the phosphorylation site in ACS7. Additionally, the protein–protein interaction between ACS7 and 14-3-3 was confirmed using two independent methods. A T-DNA insertion knockout mutant, acs7-1, was also identified and used to investigate the previously uncharacterized functions of ACS7 in Arabidopsis. It was found that ACS7 is involved in root gravitropism in Arabidopsis.

### Materials and methods

#### Plant growth conditions

Seeds from WT Arabidopsis thaliana plants (Columbia and Wassilewskija ecotype) were sterilized with chlorine for 3 h with 100 ml of 6% bleach and 3 ml of 10 N sulphuric acid (H2SO4) and then spread onto plates that contained half-strength Murashige and Skoog medium and 0.5% sucrose. The seeds were placed in the dark for 2 d at 4 °C and then incubated in growth chambers for 16 h with 100 μmol photon m−2 s−1 light and 8 h in the dark at 23 °C. After seed germination, all of the plants were transferred to a 9F walk-in growth chamber under a short-day photoperiod condition (8 h light/16 h dark) or a long-day photoperiod condition (16 h light/8 h dark).

#### Preparation of E. coli DH5α competent cells for transformation

Escherichia coli competent cells were prepared. A single colony of DH5α was inoculated into 5 ml of LB medium (1% tryptone, 0.5% yeast extract, and 1% NaCl) with shaking at 37 °C for 16 h. The cells were then added to a 1 litre flask that contained 500 ml of Super Optimal Broth medium with shaking at 37 °C for 3 h. The cells were centrifuged at 2700 g (Beckman Coulter J2-MC, Brea, CA, USA) for 10 min at 4 °C and gently resuspended in 130 ml of 0.1 M CaCl2 solution. The cells were then centrifuged at 2700 g for 10 min at 4 °C, and the supernatant was removed. Finally, the cells were resuspended in 4 ml of TB buffer and separated in a 1.5 ml tube. DH5α competent cells were placed on ice, and 1 μl of plasmid was added to thaw the cells. The cells were placed on ice for 30 min and heat-shocked for 90 s at 42 °C. The cells were then placed on ice for 2 min. LB medium (1 ml) was added to let the cells recover at 37 °C for 30 min. After recovery, the cells were spread on plates and incubated overnight at 37 °C.

#### Purification of glutathione S-transferase (GST)-tagged protein

Escherichia coli was incubated in 40 ml of 2× YT medium with 200 μg of ampicillin at 37 °C for 16–18 h overnight, and 400 ml of 2× YT medium was then added at 28 °C for 3 h. After incubation, 220 μl of isopropyl β-d-thiogalactopyranoside (IPTG; 1 M stock, 230 mg ml−1) was added to a final concentration of 0.5 mM and incubated for 3 h at 37 °C. The cells were centrifuged at 6000 rpm (Beckman Coulter J2-MC) for 30 min at 4 °C. The supernatant was discarded, and 20 ml of lysis buffer was then added for 15 min. The cells were transferred to a 50 ml Falcon tube and stored at −80 °C. The cells were incubated in water (~40 °C), and a sonicator (Misonix XL2020, Farmingdale, NY, USA) was used to break the cells. After sonication, the cells lysate was centrifuged at 10 000 rpm for 30 min at 4 °C. The supernatant was then transferred to a new 50 ml Falcon tube. GST beads (1 ml) were added and washed with GST binding buffer three times, and the solution was shaken for 1–3 h in a cold room. The GST beads were centrifuged, and 10 ml of GST binding
buffer was added to wash the beads three times. TRIS buffer (10 ml, 50 mM, pH 7.5) was added to wash the beads, and the solution was transferred to a biospin column (Bio-Rad, Hercules, CA, USA) to allow the GST beads to adhere to the column. TRIS buffer (1.5 ml, 50 mM, pH 8.0) that contained 10 mM glutathione (30 mg per 10 ml) was used to elute the protein, and the eluate was collected by a centrifugal filter (Amicon Ultra 10K, Millipore, Billerica, MA, USA).

6His-SUMO–ACS7 recombinant protein purification

6His-SUMO–ACS7 recombinant protein was incubated in 40 ml of LB medium that contained 50 μg ml⁻¹ ampicillin and shaken at 37 °C overnight. The overnight culture was inoculated with 400 ml of LB medium with shaking at 37 °C. The cells were raised to an absorbance of optical density (OD) 0.4–0.6 (mid-log phase) at 600 nm and then induced with 0.5 mM IPTG and allowed to continue to grow for 3 h. The cells were then harvested by centrifugation at 5520 g (Beckman Coulter J2-MC) for 30 min. The cell pellet was resuspended in lysis buffer and stored at −80 °C. The cells were thawed in 40 °C water, sonicated, and centrifuged at 20 400 g for 30 min (Beckman Coulter J2-MC). The supernatant was incubated with pre-washed Ni-NTA resin (GE) for 2 h at 4 °C and washed with a first wash buffer and second wash buffer. 6His protein was eluted with 1.5 ml of elution buffer. The eluate was collected with a centrifugal filter (Amicon Ultra 10K, Millipore) for buffer exchange. The fusion protein was resuspended in phosphate-buffered saline (PBS) buffer. Plasmid maps are shown in Supplementary Fig. S1 available at JXB online.

Protein quantification using the Bradford assay

The protein concentration measurement was based on the Bradford method using Protein Assay Dye (Bradford, 1976; catalog no. 500-0006, Bio-Rad). Protein assay dye (100 μl) was mixed with 900 μl of dH₂O and added to different concentrations of bovine serum albumin to reconstitute the standard curve. Sample absorbance was read at 595 nm.

Fusion peptide design and construction

Approximately 50–60 nucleotide long forward and reverse primers were used to self-ligate in a temperature gradient and had a sticky-end Ascl and BamHI restriction enzyme recognition site. After phosphorylation by polynucleotide kinase at the 5’ end for 30 min, the double-stranded primer was constructed into an NRV vector and transformed into BL21 for fusion protein expression.

In vitro kinase assay

The in vitro kinase assay was performed according to a modified method described previously (Curran et al., 2011). ATP (50 μM, spiked with 1.25 μCi of [γ-³²P]ATP) was added to begin the kinase reaction at a final volume of 10 μl, which consisted of 300 ng of purified CDPK, 3 μg of fusion protein substrate, and standard kinase reaction buffer. The reactions were incubated for 15 min at room temperature and stopped by SDS sample buffer. All of the samples were loaded onto a 12% SDS–PAGE loading well for electrophoresis, and labelling signals were normalized to the amount of protein determined from Coomassie Brilliant Blue-stained gels after running SDS–PAGE.

Protoplast isolation

Arabidopsis protoplasts were isolated according to a modified method described previously (Yoo et al., 2007). The leaves from 4-week-old plants were excised and subjected to an enzyme solution for 2 h at room temperature. The enzyme solution that contained protoplasts was filtered with a miracloth and centrifuged at 100 g (Kubota 2420, Japan) to pellet the protoplasts in a 15 ml tube for 3 min. The supernatant was removed, and the protoplasts were washed three times in W5 solution. The protoplasts were resuspended in Mmg solution with 2.5 × 10⁵ protoplasts in 1 ml before polyethylene glycol (PEG)-mediated transformation.

Plasmid construction and transformation for transient expression

The open reading frame (ORF) of AtACS7 was amplified using designated primers from cDNA. The amplified ORF was inserted into the p2YGW7 vector (Invitrogen). The method of transiently expressed plasmid transformation was performed as previously described (Yoo et al., 2007). Plasmids (10 μg) and 200 μl of protoplasts were added to a 15 ml tube and gently mixed. PEG solution (200 μl) was then added and incubated at room temperature for 10 min. The PEG solution that contained protoplasts was diluted with 1 ml of W5 solution and gently mixed. Protoplasts were centrifuged at 100 g (Kubota 2420) to pellet the protoplasts for 3 min. The supernatant was removed, and the protoplasts were washed twice with W5 solution. The protoplasts were resuspended with 1 ml of W5 solution in each well of a 6-well tissue culture plate and incubated at room temperature. After 12–16 h, yellow fluorescent protein (YFP) fluorescence was detected with a confocal microscope (SP5, Leica, Microsystems, Germany). Plasmid maps are shown in Supplementary Fig. S1 at JXB online.

Plasmid construction of bimolecular fluorescence complementation (BiFC) and transformation of plasmids for BiFC analysis

BiFC analyses were performed according to a modified method described previously (Yoo et al., 2007; Lee et al., 2012). The ORF of AtACS7 was amplified using designated primers from cDNA. The amplified ORF was inserted into the pEarlyGate201-YN vector or pEarlyGate202-YC driven by the 35S promoter and fused to YFP-N or YFP-C in-frame. Plasmids (10 μg; YFP-N and YFP-C) and 200 μl of protoplasts were added to a 15 ml round-bottomed tube and gently mixed. PEG solution (110 μl) was added and incubated at room temperature for 10 min. The PEG solution that contained protoplasts was diluted with 550 μl of W5 solution and gently mixed. Protoplasts were centrifuged at 100 g (Kubota 2420) to pellet the protoplasts for 3 min. The supernatant was removed, and the protoplasts were washed twice with W5 solution. Protoplasts were resuspended with 1 ml of W5 solution in each well of a 6-well tissue culture plate and incubated at room temperature. After 12–16 h, YFP fluorescence was detected by a confocal microscope. Plasmid maps are shown in Supplementary Fig. S1 at JXB online.

Site-directed mutagenesis

Site-directed mutagenesis was performed using the QuickChange Lightning kit (Stratagene, La Jolla, CA, USA). Two complementary oligonucleotides that contained the desired mutation, flanked by an unmodified nucleotide sequence, were designed. Mutated nucleotides were amplified by PCR. The DpIII restriction enzyme (2 μl) was then directly added to each amplification reaction and incubated at 37 °C for 5 min to digest the parental supercoiled double-stranded DNA. The DNA treated with 2 μl of DpnI was transformed into the DH5α competent cells.

SDS–PAGE

SDS–PAGE was performed according to a modified method described previously (Laemmli, 1970). A gel preparation system (Bio-Rad) was used to prepare a 4% stacking gel and 12% resolving gel. The samples were supplemented with sample buffer, heated at 95 °C for 5 min, and subjected to SDS–PAGE at a constant voltage of 100 V until the protein dye left the gel.
Quartz crystal microbalance (QCM) sensor washing and analysis

QCM was performed according to a modified method described previously (Matsumana and Ueda, 2010). A volume of 500 μl of 1% SDS was added to the sensor and left for 3 min. Double-distilled H2O (ddH2O) was used to wash the sensor. Piranha solution (3 μl; 99% H2SO4:30% H2O2; 3:1) was added to the sensor and left for 5 min. ddH2O was used to wash the sensor. This procedure was repeated twice. The sensor was placed into an AFFINIX QN tube with liquid nitrogen. The leaves were ground into a powder, and centrifuged for 10 min. Chloroform (200 μl) was added into the 1.5 ml tube. Isopropanol (500 μl) was added and centrifuged at 13 000 rpm for 10 min. The pellet was washed with 70% ethanol and resuspended in ddH2O. Genomic DNA extraction buffer and 50 μl of 20% SDS were added and centrifuged at 13 000 rpm for 10 min. Chloroform (200 μl) was added into the 1.5 ml tube and centrifuged for 10 min at 13 000 rpm. A total of 800 μl of the supernatant was moved to a new 1.5 ml tube. Isopropanol (560 μl) was added and centrifuged at 13 000 rpm for 10 min. The pellet was washed with 70% ethanol and resuspended in ddH2O. Genomic DNA was used to determine the homozygosity of the T-DNA insertion in the acs7-1 mutant using PCR.

Genomic DNA extraction

The leaves of 3-week-old plants were excised and placed into a 1.5 ml tube with liquid nitrogen. The leaves were ground into a powder, and 750 μl of Genomic DNA extraction buffer and 50 μl of 20% SDS were added. The 1.5 ml tube was heated at 65 °C for 10 min. After heating, 250 μl of 5 M potassium acetate was added and placed on ice for 10 min. Chloroform (200 μl) was added into the 1.5 ml tube and centrifuged for 10 min at 13 000 rpm. A total of 800 μl of the supernatant was moved to a new 1.5 ml tube. Isopropanol (560 μl) was added and centrifuged at 13 000 rpm for 10 min. The pellet was washed with 70% ethanol and resuspended in ddH2O. Genomic DNA was used to determine the homozygosity of the T-DNA insertion in the acs7-1 mutant using PCR.

RNA extraction

RNA was isolated using RE佐TM C&T reagent (Protech, Taipei, Taiwan) and converted to cDNA using a high-capacity cDNA reverse transcription kit (Applied Biosystems, Foster City, CA, USA). The leaf sample was ground into a powder with liquid nitrogen, and 1 ml of RE佐TM C&T and 200 μl of chloroform were added. The sample was centrifuged at 13 000 rpm for 15 min and moved to a new 1.5 ml tube. Isopropanol (500 μl) was added and centrifuged at 13 000 rpm (Sigma 1-15K, St Louis, MO, USA) for 10 min. The pellet was washed with 75% ethanol and resolved by diethylpyrocarbonte (DEPC)-treated H2O. Total RNA (2 μg) was subjected to cDNA synthesis. The same amount of cDNA was used for PCR analysis.

Root curvature and gravity-sustaining response ratio measurement

The root gravity response was measured as previously described (Sukumar et al., 2009) (Supplementary Fig. S2 at JXB online). Three-day-old seedlings were transferred to either control agar (0.8%) or agar supplemented with multiple chemicals [i.e. EGTA (Sigma), LaCl3, LiCl, ACC, 1-naphthylphthalamic acid (NPA), and ruthenium red (RR)] at the indicated concentrations. After 12–24 h of vertical growth, the plates were rotated 90° counterclockwise. Photographs of the plants were taken at specific time points (Time 0) after reorientation using a digital camera. The root tip curvature (in degrees) after reorientation (the angle difference of before and after reorientation) was measured every 12 h by ImageJ software, and the gravity-sustaining response ratio was determined using the percentage of roots that grew toward the direction of the new gravity vector.

Results

AtACS7 can be phosphorylated by AtCDPK16 in vitro

A recent study found that type I and type II ACS proteins can be CDPK substrates (Sebastià et al., 2004). Although the type III ACC synthase ACS7 has a shorter C-terminal domain and does not have a predicted CDPK phosphorylation site, ACS7 may still be a substrate of CDPK. AtCDPK1, AtCDPK16, AtCRK3, and AtCDPK34 were used for the in vitro kinase assay against the full-length ACS7 recombinant protein 6His-SUMO–ACS7. G-NR, G-Dii19-2-2 WT, and G-Dii19-2-2 mutant (MT) fusion peptides were used as controls for the kinase assay as described previously (Curran et al., 2011). The kinase assay results indicated that CDPK16 can phosphorylate the recombinant protein 6His-SUMO–ACS7 (in vitro (Fig. 1).

The possible phosphorylation sites in ACS7 were further investigated. Liquid chromatography−mass spectrometry/LC-MS/MS was used to identify the peptides in the ACS7 recombinant protein phosphorylated by CDPK. According to the LC-MS/MS results and conserved Ser/Thr residues of ACS7 in different plant species, 13 fusion peptides that contained possible phosphorylation sites were designed as candidates for the in vitro kinase assays (Supplementary Table S1 at JXB online). After the kinase assay, it was found that the fusion peptides P5-2, which contains the peptide V(294)GT(296)IYS(299)YNDNV(304), and P11-2, which contains the peptide V(207)RGVLIT(213)NPS(216)NPL(219), were labelled in the 32P autoradiogram (Fig. 2). This result indicates that P5-2 and P11-2 can be phosphorylated by CDPK16 in vitro.

A previous study showed that CDPK phosphorylates substrates in a calcium-dependent manner (Hetherington and Trewavas, 1984). Because the fusion peptides P5-2 (VGTIYSYNDNV) and P11-2 (VRGVLITNPSNPL) were shown to be phosphorylated by CDPK16, experiments were carried out to determine whether this phosphorylation is calcium dependent. Figure 3A shows that when calcium ions were added in the kinase reaction buffer, P5-2 and P11-2 phosphorylation signals were detected. When calcium ions were depleted in the reaction buffer, phosphorylation was abolished. This confirms that P5-2 and P11-2 phosphorylation is calcium dependent.

To confirm that the in vitro phosphorylation of AtACS7 fusion peptides was due to AtCDPK16 activity, a control experiment was carried out. Two autophosphorylation sites (Ser274 and Ser541) of the AtCDPK16 recombinant fusion protein (Hegeman et al., 2006) were mutated by site-directed mutagenesis. The mutated AtCDPK16 recombinant fusion protein (G-CDPK16-6His MT) was used in the in vitro kinase assay against AtACS7 fusion peptides. The results showed that the autophosphorylation of the mutated recombinant protein (G-CDPK16-6His MT) was greatly reduced (Fig. 3B; Supplementary Fig. S3 at JXB online). In addition, the phosphorylation of the AtACS7 fusion peptides was greatly reduced (Fig. 3B; Supplementary Fig. S3). This indicated that the phosphorylation of AtACS7 fusion peptides in vitro resulted from AtCDPK16 but not from other bacterial kinases. The P5-2 and P11-2 peptides each contain one serine and one threonine residue. Furhter experiments were carried out to determine which amino acid is the target for CDPK16 or whether both can be phosphorylated by CDPK16. To address this issue, a point mutation was...
created in the two peptides using site-directed mutation of either serine or threonine to alanine (Supplementary Table S1 at JXB online). For P11-2, the results indicated that only Ser216 was recognized by CDPK16 (Fig. 4). To confirm the phosphorylation sites, the fusion peptide P4-5, which has 10 amino acids that overlap with P11-2, was used for the kinase assay. A point mutation of either serine or threonine to alanine was designed to verify the phosphorylation site in P11-2. The results showed that only Ser216 was recognized by CDPK16-6His in vitro. G-NR is a fusion protein, with GST fused with a peptide of nitrate reductase (TLKRTASTPFM), and this peptide is known to be recognized by G-AtCDPK1-6His; G-NRV is a vector-only protein; G-Di19-2-2WT is also a fusion protein in which GST is fused with a peptide (DVLKSEQKEMSYREDPY); this peptide can be recognized by G-AtCDPK16-6His, and G-Di19-2-2MT is similar to G-Di19-2-2WT but with serine mutated to alanine. The molecular weights of all the fusion proteins are ~55 kDa and that of SUMO–ACS7 is ~63 kDa; the peptide phosphorylation signal is marked with a arrowhead, and the kinase autophosphorylation signal is ~100 kDa (right panel).

For P5-2, a fusion peptide with a single point mutation of either threonine or serine to alanine and double mutations of both serine and threonine to alanine was used to perform the kinase assay. The results showed that both Thr296 and Ser299 were recognized by CDPK16 (Fig. 5). Specifically, the phosphorylation of Ser299 was also supported by the MS/MS data (Supplementary Fig. S4 at JXB online). In conclusion, based on the in vitro assay, three AtCDPK16 phosphorylation sites, Ser216, Thr296, and Ser299, were identified. These results suggest that AtACS7 may be involved in calcium signalling.

The subcellular localization of AtACS7 is in the cytosol

ACS proteins have been found to be cytosolic proteins (Yip et al., 1991). To confirm further whether AtACS7 is actually localized in the cytosol, transient expression of ACS7 protein in protoplasts was detected. The 35S::YFP:ACS7 plasmid was transferred to Arabidopsis protoplasts, and confocal microscopy was used to observe the localization of the recombinant YFP fused with ACS7. As expected, ACS7 protein was localized in the cytosol based on the transient assay (Fig. 6).

Protein–protein interaction between ACS7 and 14-3-3ω was confirmed using BiFC and QCM

14-3-3 proteins have been regarded as scaffold proteins that can bind to phosphorylated proteins (Sehnke et al., 2002). A previous study showed that ACS7 can be a client of 14-3-3ω in Arabidopsis (Chang et al., 2009; Yoon and Kieber, 2013). Whether 14-3-3ω physically interacts with ACS7 and modulates its functions was investigated. Previous studies showed that ACS7 can form a homodimer to perform its functions (Tsuchisaka and Theologis, 2004), and fusion proteins ACS7-YN and ACS7-YC were used as a positive control (Supplementary Fig. S5B at JXB online). ACS7 formed homodimers as expected (Tsuchisaka and Theologis, 2004). Additionally, NR peptide, which contains a 14-3-3 interaction site, interacted with 14-3-3ω as expected. Based on the transient expression results, ACS7 physically interacted with 14-3-3ω in the cytosol (Supplementary Fig. S5E, F). Both ACS7-YN and ACS7-YC interacted with 14-3-3ω-YN.

To investigate the functional relevance of the identified phosphorylation sites, ACS7-YN and ACS7-YC, each with Ser216 point-mutated to alanine, were tested for interactions

![Fig. 1. SUMO–AtACS7 was phosphorylated by AtCDPK16 in vitro.](image)

Recombinant protein 6His-SUMO–ACS7 was used as a substrate, and four kinds of recombinant kinases (G-AtCDPK1-6His, G-AtCDPK16-6His, G-AtCRK3-6His, and G-AtCDPK34-6His) were used to perform the kinase assay in vitro. The result indicates that 6His-SUMO–AtACS7 can only be phosphorylated by G-AtCDPK16-6His in vitro. G-NR is a fusion protein, with GST fused with a peptide of nitrate reductase (TLKRTASTPFM), and this peptide is known to be recognized by G-AtCDPK1-6His; G-NRV is a vector-only protein; G-Di19-2-2WT is also a fusion protein in which GST is fused with a peptide (DVLKSEQKEMSYREDPY); this peptide can be recognized by G-AtCDPK16-6His, and G-Di19-2-2MT is similar to G-Di19-2-2WT but with serine mutated to alanine. The molecular weights of all the fusion proteins are ~55 kDa and that of SUMO–ACS7 is ~63 kDa; the peptide phosphorylation signal is marked with a arrowhead, and the kinase autophosphorylation signal is ~100 kDa (right panel).
using BiFC. ACS7-YN and ACS7-YC, each with Thr296 and Ser299 double-mutated to alanine, were also tested. Both BiFC results showed that the mutation of these phosphorylation sites resulted in an altered subcellular localization pattern. The dimer formation signal was weaker and not evenly distributed in the entire cytoplasm as it was in the WT (Fig. 7C–E), suggesting that the phosphorylation of AtACS7 may have important functions.

To confirm further the protein–protein interaction between ACS7 and 14-3-3ω, QCM analysis was performed. The recombinant protein G-14-3-3ω was first coated on the sensor, and then 6His-SUMO–ACS7 protein was injected into the PBS buffer. If the recombinant protein 6His-SUMO–ACS7 can physically interact with G-14-3-3ω, then the sensor frequency would change. The results showed that 6His-SUMO–ACS7 interacted with G-14-3-3ω in vitro (Supplementary Fig. S6F). The molecular weight of the fusion peptide is ~55kDa; peptide phosphorylation signals are marked with arrowheads, and the kinase autophosphorylation signal is ~100kDa (right panels).

Fig. 2. Peptide P5-2 and P11-2 were phosphorylated by CDPK16 in vitro. The fusion proteins (GST fused with small peptides) were designed to perform the kinase assay and determine which serine or threonine can be phosphorylated by G-CDPK16-6His. P5-2 contains the peptide V(294)GTIYSYNDNV(304); and P11-2 contains the peptide V(207)RGVLITNPS(216)NPL(219). G-Di19-2-2WT phosphorylated by G-CDPK16-6His is a positive control; G-Di19-2-2MT is a negative control; and 6His-SUMO–ACS7 is a full-length ACS phosphorylation control. The results indicate that P5-2 (in B) and P11-2 (in D) can be phosphorylated by CDPK16. The interaction assay was performed with phosphorylated 6His-SUMO–ACS7 and G-14-3-3ω. The dissociation constant (K_d) was 9.29 ± 1.99 × 10^{-9} M (Supplementary Fig. S6C). A previous study showed that 14-3-3 is a scaffold protein that can interact with its clients that have already been phosphorylated (Wu et al., 1997). To determine whether the interaction between ACS7 and 14-3-3ω is phosphorylation dependent and whether the interaction is caused by ACS7 phosphorylation catalysed by CDPK16, an interaction assay between phosphorylated 6His-SUMO–ACS7 and G-14-3-3ω was performed. 6His-SUMO–ACS7 was phosphorylated by CDPK16 in vitro in advance. The results showed that phosphorylated 6His-SUMO–ACS7 can physically interact with G-14-3-3ω in vitro (Supplementary Fig. S6B), and the K_d was 5.05 ± 1.96 × 10^{-9} M (Supplementary Fig. S6D). The QCM results of tag-only controls did not show any interaction, as expected (Supplementary Fig. S6E, F). Altogether, the results are consistent with the previous discovery that ACS7 forms a protein complex with 14-3-3ω protein (Chang et al., 2009; Yoon and Kieber, 2013). The interaction between ACS7 and 14-3-3ω protein appeared to
be both phosphorylation dependent and phosphorylation independent.

**Identification of acs7-1 loss-of-function mutant line**

To study the functions of AtACS7 in planta, an acs7-1 mutant line was ordered from the Arabidopsis Biological Research Center (http://abrc.osu.edu/). In this mutant line, T-DNA was inserted into the third exon of the AtACS7 gene (Supplementary Fig. S7A at JXB online). The background of the mutant line was Wassilewskija (WS). Genomic PCR was used to confirm the T-DNA insertion site. The T-DNA primer GBK5-F paired with the gene primer acs7-1-R, and the gene primer acs7-1-F paired with acs7-1-R were used to perform PCR. It was found that only the T-DNA primer GBK5-F with acs7-1-R flanked the DNA fragment, and acs7-1-F with acs7-1-R did not (Supplementary Fig. S7B). The results showed that the acs7-1 mutant was homozygous.

The acs7-1 mutant is less sensitive to inhibition by a calcium chelator and the channel blocker LiCl in the gravity response

A previous study indicated that acs7-1 has multiple phenotypes (i.e. lower ethylene emission, larger cotyledons and true leaves, and longer primary root length; Dong et al., 2011). Additionally, the acs7-1 mutant exhibited hypersensitivity to abscisic acid (ABA) during seed germination (Dong et al., 2011). However, no studies of which the authors are aware have discussed the aspect of calcium in the acs7-1 mutant. The phenotype of the acs7-1 mutant line has been widely described (Tsukihisa et al., 2009; Dong et al., 2011). The acs7-1 mutant exhibits early flowering,
slightly reduced ethylene production, larger cotyledons, and enhanced salt tolerance. Because little is known about the relationship between ethylene biosynthesis and calcium (Raz and Fluhr, 1992; Philosoph-Hadas et al., 1996), several calcium channel blockers (i.e. LiCl, LaCl$_3$, and RR), a calcium chelator (i.e. EGTA), and the CaM antagonist chlorpromazine (CPZ) were used to investigate the relationships between them. AtACS7 can be phosphorylated by CDPK in vitro, and it is hypothesized that AtACS7 may be involved in calcium signalling. Previous studies showed that calcium ions are important for the gravity response. When the WT seedlings are treated with calcium channel blockers or a calcium chelator, the plants may lose their gravity response (Hasenstein and Evans, 1988; Friedman et al., 1998; Toyota et al., 2008). To investigate the relationship between acs7-1 and calcium, the calcium channel blockers LiCl, LaCl$_3$, and RR, the calcium chelators EGTA, and the CaM antagonist CPZ were used to determine whether calcium ions influence acs7-1. The results indicated that LaCl$_3$ and CPZ did not cause significant differences between the

Fig. 4. Phosphorylation of Ser216 was confirmed by site-directed mutagenesis. In order to identify the phosphorylation residue in P11-2, site-directed mutagenesis was performed. P4-5 is a peptide which has a 10 amino acid sequence overlapping with P11-2. P11-2 S→A is a serine to alanine mutation peptide in P11-2; P11-2 T→A is a threonine to alanine mutation peptide; P11-2 TS→AA is a double mutation peptide of serine and threonine to alanine. The results indicated that only serine (Ser216) but not threonine (Thr213) in P11-2 can be phosphorylated by G-CDPK16-6His in vitro. Peptide phosphorylation signals are marked with arrowheads.
WT and the acs7-1 mutant (Supplementary Figs S8, S9 at JXB online). Surprisingly, EGTA treatment caused a loss of the gravity response in the WT but not in the acs7-1 mutant. Figure 8C–E shows that acs7-1 can retain gravitropism, but the WT almost lost gravitropism. Moreover, 10 mM LiCl treatment had the same effect as EGTA treatment (Fig. 8B, D, E). Collectively, the results indicate that ACS7 is involved in root gravitropism in a calcium-dependent manner in Arabidopsis.

The acs7-1 mutant is less sensitive to ethylene in the triple response

To determine whether the acs7-1 mutant displays ethylene-related phenotypes that are different from the WT in etiolated seedlings, the hypocotyl length was measured in the dark for 5 d and analysed by ImageJ software. The triple-response results showed that the acs7-1 mutant is less sensitive to ethylene compared with the WT (Fig. 9).

Gravity response under optimal ACC concentration

Ethylene has been suggested to regulate the gravity response. Some studies indicate that it plays a negative role in gravitropism (Buer et al., 2006), but other studies suggest a positive role in gravitropism (Philosoph-Hadas et al., 1996). To investigate the effects of ethylene on gravitropism, gradient concentrations of ACC were introduced in the medium. Both the WT and the acs7-1 mutant responded to certain concentrations of ACC that positively regulated the gravity response (WT at 1 μM ACC; acs7-1 at 0.1 μM ACC). A specific concentration of ethylene with 0.01 μM ACC negatively regulated the gravity response in both the WT and the acs7-1 mutant (Fig. 10).
The *acs7-1* mutant is less sensitive to NPA-mediated inhibition of the gravity response

Some auxin polar transport inhibitors, including NPA and 2,3,5-triiodobenzoic acid (TIBA), were found to inhibit auxin polar transport. For example, TIBA can compete with auxins for translocation across the plasma membrane (Depta and Rubery, 1984), and NPA was found to disturb the polarity of cell division (Thomson et al., 1973). Therefore, NPA has long been used as an inhibitor of auxin polar transport (Thomson et al., 1973). The gravity response is greatly reduced by NPA treatment. To investigate the effects of NPA on the gravity response in the WT and the *acs7-1* mutant, different concentrations (12 μM and 2 μM) of NPA were tested. The WT showed a smaller root curvature and lower gravity response than the *acs7-1* mutant (Fig. 11A–D) in response to NPA treatment. This indicates that the *acs7-1* mutant was less sensitive to the inhibitory effect of NPA. Exogenous ACC (0.1 μM) was added to the medium with NPA, which greatly reduced the effect of NPA and recovered root gravitropism (Fig. 11A–D). These results suggest that AtACS7 is involved in the root gravity response through an unknown cross-talk relationship between auxin and ethylene.

**Discussion**

Figure 1 shows the phosphorylation of AtACS7 catalysed by AtCDPK16 *in vitro*. However, this result is different from the previous expectation that type III ACS proteins have no predicted CDPK phosphorylation sites and may not be a substrate of CDPKs. To date, only one *Arabidopsis* AtCDPK16 substrate, AtDi19-2, has been documented (Curran et al., 2011). The present study showed that AtCDPK16 can phosphorylate AtACS7 *in vitro*, but whether AtACS7 is an AtCDPK16 substrate requires further studies. Several consensus CDPK phosphorylation motifs are known (Harper and Harmon, 2005). The first consensus phosphorylation motif is ϕ_{2-3}-X_{4-5}-Basic_{3-2}-X_{3-4}-S, in which the underlined S is phosphorylated, X is any residue, and ϕ is a hydrophobic...
The second consensus phosphorylation motif is Basic$_{3}$-Basic$_{3}$-X$_{7}$-Basic$_{2}$-ϕ$_{3}$-X$_{4}$-X$_{3}$-X$_{2}$-S-X$_{1}$-Basic$_{2}$. The third consensus phosphorylation motif is ϕ$_{3}$-R$_{3}$-ϕ$_{2}$-S-ϕ$_{1}$-X-K$_{3}$-R$_{4}$+2, which was defined as an ACS motif in a previous study (Sebastià et al., 2004). The identified phosphorylation site of AtACS7, Thr296, perfectly matches the first consensus phosphorylation motif. The phosphorylation motif S-X-R was recently identified from the AtCDPK16 substrate AtDi19-2 (Curran et al., 2011). In the present study, Ser216 and Ser299 were found to be phosphorylated by CDPK16 in ACS7 in vitro (Figs 2, 4, 5) but did not conform to any of these consensus phosphorylation motifs. Ser216 and Ser299 appear to be newly identified CDPK phosphorylation sites with an unknown CDPK phosphorylation motif. Whether Ser216, Thr296, and Ser299 are indeed phosphorylated by CDPK in vivo needs further confirmation.

Fig. 8. Multiple calcium chelator and channel blockers affected the gravity response in acs7-1. The WT (WS) and the acs7-1 mutant were grown in half-strength Murashige and Skoog (1/2MS) medium for 3 d and transferred to 1/2MS medium which contained (A) control (normal conditions), (B) 10 mM LiCl, (C) 0.25 mM EGTA and other calcium channel blockers, and grown for 1 d. The gravity vector is downwards as the arrowhead indicates. The root curvature (degree) and ratio of gravity response (%) were measured in both the WT and the acs7-1 mutant after 24 h (D and E). The average and SE are presented (n=60) for multiple calcium chelators or channel blockers in three independent experiments in (C), and the ratio of the gravity response is presented (n=60) for multiple calcium chelators or channel blockers in (D). RR, ruthenium red. Significant differences between normal conditions and treatment groups are indicated by * at $P < 0.05$ and by ** at $P < 0.01$ by Student’s t-test.

The Arabidopsis genome has eight genes that encode active ACS proteins and an additional gene that encodes a catalytically inactive enzyme, ACS1 (Liang et al., 1992). Previous studies showed that ACS1 was inactive because it lacked the highly conserved tripeptide Thr-Asn-Pro (TNP) between Ile204 and Ser205 in the ACS conserved region that was responsible for binding the cofactor PLP (Liang et al., 1995; Yamagami et al., 2003). The introduction of TNP into ACS1 restores ACS activity, whereas its removal from enzymatically active ACS2 results in a loss of activity (Liang et al., 1995). In the sequence alignment of plant ACS proteins (data not shown), it was found that ACS7 has a tripeptide, Thr-Asn-Pro (TNP), between Ile212 and Ser216. Ser216 was phosphorylated by CDPK16 in the in vitro kinase assay (Figs 2, 4), and the phosphorylation of Ser216 in ACS7 can probably influence the interaction between ACS7 and its cofactor PLP and finally change the enzyme activity. To investigate
whether phosphorylation can influence enzyme activity, an in vitro enzyme activity assay was performed according to previous studies (Li et al., 1996; Chae et al., 2003). The results indicated that non-phosphorylated ACS7 had a $K_m$ value of 27.43 μM, whereas phosphorylated ACS7 had a lower $K_m$ value (13.65 μM; data not shown). These results suggest that phosphorylated ACS7 has a higher catalytic ability, and the phosphorylation of ACS7 catalysed by CDPK16 may enhance ACS7 activity. However, such a possibility requires further studies.

ACS gene expression in light-grown mature Arabidopsis plants was previously studied using reverse transcription–PCR (RT–PCR) (Yamagami et al., 2003), in which the expression of all Arabidopsis ACS genes was surveyed in plant roots, leaves, stems, flowers, and siliques. These researchers found that ACS7 was expressed only in roots. It was confirmed here that ACS7 is a cytosolic protein (Fig. 6), suggesting that ACS7 protein is mainly localized in the root cytosol. Another study used a green fluorescent protein (GFP) fused to nine kinds of AtCDPKs (Dammann et al., 2003). The authors found that AtCDPK16 was localized to the root plasma membrane in a thin layer at the periphery of the cell. Further studies are needed to confirm whether ACS7 can be phosphorylated by CDPK16 in plant roots in vivo.

Lithium chloride is known to suppress root curvature in Pisum sativum seedlings (Belyavskaya, 2001). In Arabidopsis, the root cap comprises four tiers of columella cells that originate from initial columella cells. Columella cells have a specific polarity, in which the nucleus and ER are localized to the proximal side of the root meristem and periphery of the cell, respectively. In addition to the peripheral arrangement of the ER, its function as an intracellular calcium reservoir has been the preferred hypothesis for a possible gravity-sensing mechanism in columella cells, in which amyloplasts that come into contact with the peripheral ER can trigger the release of calcium stored in the ER (Perbal and Driss-Ecole, 2003). Many signalling molecules and secondary messengers, such as calcium (Plieth and Trewavas, 2002) and InsP$_3$ (Perera et al., 2001), and pH (Johannes et al., 2001) have been implicated in linking gravity sensing to the initiation of a differential growth response. Very rapid gravity-specific changes in transcript abundance have been documented in gravistimulated root tips of Arabidopsis (Kimbridge et al., 2004) and whole seedlings (Moseyko et al., 2002). InsP$_3$ is well known to induce calcium release from intracellular calcium reservoirs. Changes in InsP$_3$ levels have been reported after the gravistimulation of pulvini in maize (Perera et al., 1999). Lithium chloride has been reported to be a phosphatidylinositol cycle inhibitor (Belyavskaya, 2001; Dieck et al., 2012). It can suppress InsP$_3$ cycling and signalling by inhibiting inositol-1-phosphatase (Belyavskaya, 2001), leading to a reduction in the amount of InsP$_3$. When plants are treated with LiCl, some plant organs, such as tonoplasts and the ER, that are reported to have IP$_3$-sensitive calcium channels may be blocked, leading to a reduction of calcium ion efflux from these organs. When the gravity vector changes, InsP$_3$ may bind to the ER or tonoplast InsP$_3$-sensitive calcium channels, which ultimately leads to calcium ion efflux from these organs. These calcium ions are captured by some calcium sensors, such as CDPK and CaM, and pass the signals to downstream effectors, ultimately leading to the plant gravity response.

Several mutant lines have been shown to be related to calcium signalling in root gravitropism in plants. A mutation of the calmodulin agr-3 gene exerted a reduced root gravitropism phenotype in Arabidopsis (Sinclair et al., 1996). Spalding's group found that a mutation of the Glutamate Receptor-Like 3.3 gene (i.e. a mammalian calcium channel orthologue) showed a root gravitropism phenotype (Miller et al., 2010). Moreover, an Arabidopsis calcium-dependent
binding protein AtCLB gene overexpression line exhibited reduced root gravitropism, but a T-DNA insertion mutant exhibited greater root gravitropism than the WT (de Silva et al., 2011). However, little is known about whether the ACS gene is involved in calcium signalling. Additionally, these mutant lines expressed a root gravitropism phenotype without EGTA treatment. In contrast, the present study observed root gravitropism with EGTA treatment. Therefore, the root gravitropism phenotype observed here appeared to be distinct from others.

Auxin polar transport is involved in many developmental processes, such as vascular differentiation and tropic growth (Luschnig et al., 1998; Mattson et al., 1999; Rashotte et al., 2000). The auxin polar transport inhibitor NPA was used to investigate the role of the ACS7 gene in the gravity response using a T-DNA knockout acs7-1 mutant line and it was found that acs7-1 is more resistant to NPA-mediated inhibition (Fig. 11). The data indicate that the inhibition of the gravity response by NPA was greatly attenuated by the addition of 0.1 μM exogenous ACC (Fig. 11). Collectively, these findings suggest cross-talk between auxin and ethylene in root gravitropism regulated by AtACS7. Cross-talk between the phytohormone auxin and ethylene in root gravitropism has been reported previously. In 1990, Estelle’s group found that an auxl mutant exhibited a defect in root gravitropism and resistance to both auxin and ethylene in Arabidopsis (Pickett et al., 1990). In 2003, an alhl mutation revealed cross-talk between ethylene and auxin in Arabidopsis (Vandenbussche et al., 2003). However, how auxin engages in cross-talk with ethylene to regulate root gravitropism through AtACS7 in Arabidopsis requires further investigation.

The rice type II ACC synthase OsACS1 was found to interact with rice 14-3-3 proteins in a yeast two-hybrid assay (Yao et al., 2007). These authors found that the C-terminal domain of OsACS1 (RSvSCP) was predicted to be phosphorylated by CDPK through a Mode-I 14-3-3 recognition
motif. They suggested that OsACS1 phosphorylation catalysed by OsCDPK may interact with rice 14-3-3 proteins, which can prevent the substrate ubiquitin adaptor protein ETO1 from binding to produce 26S proteasome degradation. Interactions with 14-3-3 proteins on the CDPK phosphorylation site may prevent ETO1 protein binding, and this may be a reason why the phosphorylation of ACS proteins catalysed by CDPK enhances ACS protein stability. In fact, the phosphorylation status of AtACS6 affected the stability of AtACS6 (Skottke et al., 2011). Moreover, the stability of AtACS5 was also found to be regulated by phosphorylation and 14-3-3 binding (Yoon and Kieber, 2013). A recent study found that ACS7 is turned over in a 26S proteasome-dependent manner and that ACS7 degradation requires the E3 ligase XBAT32 (Lyzenga et al., 2012). Whether phosphorylation and 14-3-3 binding are involved in the regulation of AtACS7 degradation requires further investigation.

Apart from ACS, ACO is another important enzyme in the ethylene biosynthesis pathway that catalyses the oxidation of ACC to ethylene. ACO is encoded by a small multigene family that is usually composed of three to four members. In the Arabidopsis genome, six genes that encode ACOs were identified in silico by Babula et al. (2006). Recently, ACO2 and ACO4 proteins were found, which may interact with 14-3-3 proteins in a yeast two-hybrid system (Jaspert et al., 2011). Together with the present results, this indicates that 14-3-3 proteins may regulate the ethylene biosynthesis pathway by modulating ACS and ACO proteins. A previous study showed that an ACO promoter can respond to IAA, bending stress, and gravity (Yuan and Dean, 2010). Together, ACS proteins and ACO may synergistically participate in the gravity response by regulating ethylene concentrations in plant cells.

Supplementary data

Supplementary data are available at JXB online. Figure S1. Plasmid maps used in the present study. Figure S2. Gravity response measurement method. Figure S3. Bar graph of the phosphorylation signal in the in vitro kinase assay using mutated AtCDPK16. Figure S4. MS/MS fragmentation pattern of a phosphorylated peptide (VGTIYSYNNDNVVR) of ACS7 recombinant protein. Figure S5. Protein–protein interaction between ACS7 and 14-3-3o using BiFC analysis. Figure S6. Quartz crystal microbalance (QCM) and kinetic analysis of protein–protein interaction between ACS7 and 14-3-3o recombinant proteins. Figure S7. Isolation of the T-DNA insertional mutant acs7-1. Figure S8. Gravity response of the WT and acs7-1 mutant affected by 0.5 mM LaCl3. Figure S9. The effect of 50 μM CPZ on the WT and acs7-1 mutant. Table S1. Information on fusion peptides.

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