INTRODUCTION

Bacteria sense their population density and act in concert using signal molecules of low molecular weight to monitor their local population density and to coordinate their collective behavior in a process called “quorum sensing” (QS). N-acyl-homoserine lactones (AHLs) are the primary QS signals among Gram-negative bacteria. AHL-mediated QS plays an essential role in diverse bacterial physiological processes. Recent evidence shows that plants are able to sense bacterial AHLs and respond to them appropriately. However, little is known about the mechanism by which plants perceive and transduce the bacterial AHLs within cells. In this study, we found that the stimulatory effect of 3OC6-HSL on primary root elongation of Arabidopsis was abolished by the calmodulin (CaM) antagonists N-(6-aminohexyl)-5-chloro-1-naphthalene sulfonamide (W-7) and trifluoperazine (TFP). Western-blot and ELISA analysis revealed that the concentration of CaM protein in Arabidopsis roots increased after treatment with 1 μM 3OC6-HSL. Results from quantitative RT-PCR demonstrated that the transcription of all nine CaM genes in Arabidopsis genome was up-regulated in the plants treated with 3OC6-HSL. The loss-of-function mutants of each AtCaM gene (AtCaM1-9) were insensitive to 3OC6-HSL stimulation of primary root elongation. On the other hand, the genetic evidence showed that CaM may not participates the inhibition of primary root length caused by application of long-chained AHLs such as C10-HSL and C12-HSL. Nevertheless, our results suggest that CaM is involved in the bacterial 3OC6-HSL signaling in plant cells. These data offer new insight into the mechanism of plant response to bacterial QS signals.

Keywords: quorum sensing, calmodulin, N-acyl-homoserine lactone, signal transduction, plant development
On the other hand, some AHLs or AHL-producing bacteria induced resistance against microbial pathogens (Schuegger et al., 2006; Schikora et al., 2011; Schenk et al., 2012, 2014; Zarkani et al., 2013). 3OC14-HSL has been shown to induce resistance against biotrophic and hemibiotrophic pathogens in Arabidopsis and barley (Schikora et al., 2011). Similarly, resistance against Pseudomonas syringae induced by Sinorhizobium meliloti in Arabidopsis plants depended on the accumulation of AHL (Zarkani et al., 2013). Transcriptomic and proteomic analysis showed that treatments of plant with bacterial AHLs elicit significant changes in gene transcription and protein expression (Mathesius et al., 2003; Ortiz-Castro et al., 2008; Miao et al., 2012; Schenk et al., 2014). Arabidopsis mutants like drr1 and dhmi1 that are tolerant or hypersensitive to alkamides from plants show alterations in primary and lateral root development and in root responses to C10-HSL, indicating that medium-chained AHLs and alkamides act through common signaling mechanisms (Morquecho-Contreras et al., 2010; Pelagio-Flores et al., 2013).

All these data indicated that AHLs might mediate the interaction between bacteria and their host plant (Hartmann et al., 2014). However, the signal transduction pathways of AHLs in plant cells are widely unexplored so far. The identification and investigation of those components in signaling will be crucial to understand the mechanism of plant response to bacterial AHLs.

Calcium is an ubiquitous second messenger in plants. Intracellular cytosolic free Ca\(^{2+}\) concentration ([Ca\(^{2+}\)]\(_i\)) often shows significant elevation in response of plant to various environmental stimuli (Knight et al., 1991; van der Luit et al., 1999). Ca\(^{2+}\) signals are sensed and translated into proper cellular response by diverse Ca\(^{2+}\) binding proteins and their downstream targets. Calmodulin (CaM) is an acidic Ca\(^{2+}\) binding protein that possesses EF hand motifs, a helix-loop-helix structure for binding one Ca\(^{2+}\) ion. Upon Ca\(^{2+}\) binding, CaM undergoes conformational changes that promote either its own catalytic activity or its interaction with target proteins (Chin and Means, 2000; Snedden and Fromm, 2001; Cheval et al., 2013). Ca\(^{2+}\)-CaM signaling has been implicated in various developmental and adaptation responses. Many exogenous and endogenous factors including light, temperature, drought and salt stress, pathogen-derived molecules and phytohormones provoke elevation in [Ca\(^{2+}\)]\(_i\) of plants (Sander et al., 1999). Likely, the level of CaM was up-regulated by heat-shock (HS) in maize seedlings (Gong et al., 1997). Pharmacological analysis revealed that CaM participated in signal transduction in lemon in response to bacterial pathogen Alternaria alternata (Ortega et al., 2002). Studies of plants demonstrated the presence of multiple CaM genes that encode numerous CaM isoforms (Lee et al., 1995; Takezawa et al., 1995; Yang et al., 1996; Snedden and Fromm, 2001; Yamakawa et al., 2001; Zielinski, 2002). Braam and Davis (1990) reported that the expression of many plant CaM genes are induced by rain, wind and touch. Among of eight CaM genes in potato plants, PCM1 showed an increase in expression upon touching and PCMs during tuberization (Takezawa et al., 1995). Soybean CaM genes SCAm-4 and SCAm-5 are reported to be required for defense against pathogen attack (Heo et al., 1999). AtCaM3, one of nine Arabidopsis CaM genes, has been shown to be involved in Ca\(^{2+}\)-CaM mediated HS signal transduction pathway (Liu et al., 2003, 2005; Zhang et al., 2009). Recently, we have reported evidence for a transient elevation in [Ca\(^{2+}\)]\(_i\) upon the exposure of Arabidopsis roots to C4-HSL (Song et al., 2011). Given that CaM, as a Ca\(^{2+}\) sensing protein, translates Ca\(^{2+}\) to downstream protein targets in numerous signal transduction cascades, CaM might be involved in bacterial AHL signaling in plant cells. However, there is no direct evidence for the participation of CaM in AHL-mediated primary root elongation in plants.

In this paper, we first demonstrated the involvement of CaM in 3OC6-HSL mediated primary root growth in Arabidopsis using pharmacological approaches. Then we observed the increased level of CaM protein and CaM gene transcription after treatment with 3OC6-HSL. Furthermore, we used T-DNA knockout mutants to provide genetic evidence for the role of individual CaM gene in primary root elongation regulated by 3OC6-HSL. A possible regulatory model of Ca\(^{2+}\)-CaM in AHL signaling in plant cells is discussed.

**MATERIALS AND METHODS**

**PLANT MATERIALS AND GROWTH CONDITIONS**

*Arabidopsis thaliana* (L.) cv. Columbia-0, the T-DNA insertion null mutants *cam1* (CS872565), *cam5* (SALK_007371C), *cam6* (SALK_071609C), *cam7* (SALK_074336C), *cam8* (SALK_022524C) and *cam9* (SALK_040392) purchased from ABRC (http://abrc.osu.edu), and *cam2*, *cam3*, and *cam4* generously gifted from Prof. Sujuan Cui in Hebei Normal University were used in this study. All seeds were surface-sterilized by 75% (v/v) ethanol for 30 s and 20% (v/v) NaClO for 5 min. After five washes with sterile distilled water, seeds were germinated on agar plates containing Murashige and Skoog (MS) medium (Murashige and Skoog, 1962) adjusted to pH 5.8. The plates were stratified at 4°C for 2 days and then placed in a plant growth chamber with a photoperiod of 12 h of light, 12 h of darkness, light intensity of 100 μmol·m\(^{-2}\)·s\(^{-1}\) and temperature of 22 ± 2°C. For primary root growth assay, the seeds were grown vertically for 3 days on MS agar plates, and the seedlings with similar root length were transferred to 1/2 MS agar plates containing the indicated compounds. For qRT-PCR assay, a hydroponic system was employed to cultivate Arabidopsis. Seedlings germinated on MS agar plates for 10 days were transplanted into a sterile plastic basin (18 by 11 cm) containing 500 ml of sterile Hoagland medium. After 7 days, the seedlings were used for compounds treatments. Plants grown under these conditions were more vigorous than those grown in potting media, uniformly absorbing solution with AHL. To ensure that the hydroponic system is free of bacterial contamination, the medium and compound solution were sterilized by passing them through a 0.22-μm filter just prior to use. The untreated plants were taken as the control. After harvest, an aliquot of the growth medium was plated on bacterial growth medium and incubated overnight to check for contaminations that occurred during the experiment. Any contaminated samples were discarded. For mutants identification assay, seedlings germinated on MS agar plates for 10 days were transplanted into vermiculite media and watered Hoagland medium once a week. After 1 month, the leaves of seedlings were harvested for RT-PCR.
COMPOUNDS
N-(β-ketocaproyl)-DL-homoserine lactone (3OC6-HSL), N-decanoyl-DL-homoserine lactone (C10-HSL) and N-dodecanoyl-DL-homoserine lactone (C12-HSL) were purchased from Sigma-Aldrich (Taufkirchen, Germany), stored dry and diluted as 10 mM stock solutions in dH2O or ethanol just prior to use. Trifluoperazine (TFP) was purchased from Sigma (USA) and diluted as 10 mM stock solutions in methanol. N-((6-aminohexyl)-5-chloro-1-naphthalene-sulfonamide (W-7) was purchased from Santa Cruz Biotechnology (USA) and diluted as 10 mM stock solutions in dH2O. N-((6-aminohexyl)-1-naphthalene-sulfonamide hydrochloride (W-5) was purchased from Tokyo Chemical Industry (TCI, Japan) and diluted as 10 mM stock solutions in dH2O. All compound solutions were sterilized by passing them through a 0.22-μm filter.

PHARMACOLOGICAL ASSAY
CaM antagonists TFP, W-7 and W-5 were used for pharmacological assay. W-5 is the dechlorinated analog of W-7 and is often used as control compound for understanding the specificity of W-7. After germination for 3 days, Arabidopsis (Columbia, Col-0) seedlings with similar root length were transferred to 1/2 MS medium plates containing the indicated concentrations of 3OC6-HSL or/and CaM antagonists. The untreated plants were taken as the control. Seedlings were grown vertically with the same growth condition as described above, and primary root length was assessed with a ruler after 7 days. The concentration of TFP used in this study ranged from 1 to 50 μM, while the concentration of W-7 and W-5 ranged from 30 to 200 μM. 1 μM 3OC6-HSL was used in all experiments. The results were normalized to the root length of control samples. Four to six independent experiments were done and data shown are the average of 30 samples.

IMMUNODETECTION OF CaM
Seventeen-day-old Arabidopsis Col-0 (Columbia, Col-0) seedlings by hydroponic culture were treated with 1 μM 3OC6-HSL. The untreated plants were taken as the control. Roots were harvested at 0, 1, 2, 3, 6, 12, and 24 h after 3OC6-HSL treatment and powdered with liquid nitrogen. Total protein were extracted with protein extracting buffer [50 mM Tris-HCl (pH 7.5), 5 mM EDTA, 5 mM EGTA, 10 mM DTT, 10 mM Na3VO4, 10 mM NaF, 1 mM PMSF, 2 μg/ml Pepstain, 2 μg/ml Leupeptin, 2 μg/ml Aprototin, 10% (v/v) NP-40, 10 mM β-Glycerophosphate] after 14,000 rpm centrifuge for 20 min at 4°C. Protein was quantified by the Bradford method using bovine serum albumin as the standard (Bradford, 1976). For ELISA analysis, 2 μg total proteins were coated in the 96-well plate, and the polyclone antibody against wheat CaM (generous gift from Prof. Daye Sun in Hebei Normal University) and HRP-labeled goat anti-rabbit IgG antibody (Bioworld Technology Inc., St. Louis Park, USA). The images were detected by Biodlight ECL Chemiluminescent HRP Kit (Bioworld Technology Co. Ltd, St. Louis Park, USA) and scanned by BIO-RAD ChemiDoc TM MP Imaging System (Bio-Rad Laboratories, Inc. USA). Actin was used as the internal quantification control by using plant actin mAb (Bioworld Technology Inc., St. Louis Park, USA).

QUANTITATIVE REAL-TIME PCR (qRT-PCR)
Seventeen-day-old seedlings were cultured in Hoagland medium with or without 1 μM 3OC6-HSL. Roots of the treated plants were harvested at 0, 1, 2, 3, 6, 12, and 24 h after 3OC6-HSL treatment and total RNA was extracted from the treated and untreated roots using the RNAiso Plus reagent (TaKaRa, Dalian, China) according to the manufacturer’s instructions. The cDNA was synthesized using the PrimeScript® RT Reagent Kit with gDNA Eraser (TaKaRa, Dalian, China) according to the manufacturer’s instructions. For the relative quantification of gene expression, the comparative Ct method (Livak and Schmittgen, 2001) was used with the 7500 Real Time PCR System (Applied Biosystems, Foster City, CA, USA). For qRT-PCR, received cDNA was diluted 15 times in diethylpyrocarbonate-treated water. PCR amplification was done in a total volume of 20 μl containing 5 μl of diluted cDNA, 0.4 μl of each primer (10 μM), and 10 μl of SYBR Premix Ex Taq (TaKaRa, Shiga, Japan). The following qRT-PCR thermal cycling program was employed: 10 s at 95°C and 40 cycles for 5 s at 95°C and 31 s at 60°C. The amount of target was normalized to the endogenous reference gene Actin 2. Values were normalized to the 0 h time point. For technical control, each qRT-PCR experiment was repeated four times on the same 96-well plate. Each data point represents the average of three independent experiments. A 1.5-fold increase (ratio > 1.5) or 1.5-fold decrease (ratio < 0.8) in expression in the treated plants compared with the untreated plants was considered as up-regulation or down-regulation related to 3OC6-HSL response. The specific primers for qRT-PCR are shown in Supplementary Table 1.

IDENTIFICATION OF CaM NULL MUTANTS
The leaves of one-month-old wild type (Columbia, Col-0) and CaM mutants (cam1-cam9) seedlings cultured in vermiculite were harvested respectively and total RNA was extracted using the RNAiso Plus reagent (TaKaRa, Dalian, China) according to the manufacturer’s instructions. The cDNA was synthesized using the PrimeScript® RT Reagent Kit with gDNA Eraser (TaKaRa, Dalian, China) according to the manufacturer’s instructions. For RT-PCR, amplification was done in a total volume of 20 μl containing 1 μl of received cDNA, 1 μl of each primer (10 μM), 2 μl of 10×PCR buffer, 2 μl of dNTP (2.5 mM each), 0.2 μl of rTaq DNA Polymerase (5 units/μl, TaKaRa, Shiga, Japan) and 12.8 μl of sterilized distilled water. The following RT-PCR thermal cycling program was employed: initial denaturation for 5 min at 94°C, 30 cycles for 40 s at 94°C, 50 s at 52°C and 1 min at 72°C, and then extension for 10 min at 72°C. Actin 2 was used as the endogenous reference gene. The primer pairs were designed at 5'-UTR and 3'-UTR respectively for containing ORF sequence of each CaM genes and avoiding homologs mismatches between the nine CaM genes (Supplementary Table 2).
ANALYSIS OF PRIMARY ROOT GROWTH

The seeds of wild type (Columbia, Col-0) and mutants Atcam1-9 (cam1-cam9) germinated at MS agar plate vertically. After 3 days, seedlings with similar root length were transferred to 1/2 MS agar plates containing 1 μM 3OC6-HSL or 50 μM C10-HSL or 50 μM C12-HSL. Because of the space limitation in one plate, the seedlings of WT and each three mutants were grown side by side vertically in one plate with the same growth condition as described above. The equivalent dH2O-treated plants and ethanol-treated plants were taken as the solvent control. Primary root length was assessed with a ruler. In addition, the fresh weight of seedlings were assessed with an electronic balance (accuracy of 10,000 g) after 7 days. Each experiment included at least 30 seedlings for each genotype and treatment. Data shown are the average of four independent experiments.

For dissecting the root elongation, the length of root meristem zone and root elongation zone were measured on the 7-days-old seedlings fixed in ethanol: acetic acid (3:1) and mounted in choral hydrate. Microscopy was performed on a Leica DM4000 B (CMS GmbH, Wetzlar, Germany) equipped with a Leica DFC420C camera. Root meristem zone length was assessed as the distance between the quiescent center and the first elongating cell with a vacuole. The length of elongation zone was measured as the distance from the first elongating cell to the first cell with a root hair. The number of the cortical cells in root meristem and elongation zone was obtained by counting the cortical cells under the high-resolution images of the root tips using Image J software (http://rsb.info.nih.gov/ij/). The length of cortical cells in the root meristem or elongation zone was calculated by dividing the root meristem or elongation zone length by the number of the cortical cells in root meristem or elongation zone. At least 20 seedling were processed, and at least three independent experiments were conducted, giving the similar statistically significant results.

STATISTICAL ANALYSIS

For all experiments, the overall data were statistically analyzed in the DPS v7.05 program. Univariate and multivariate analyses with a Duncan’s new multiple range method were used for testing different root growth responses of different ecotype, wild-type, and CaM mutant lines to 3OC6-HSL or to CaM antagonists.

RESULTS

CaM ANTAGONISTS BLOCK THE IMPACT OF 3OC6-HSL ON PRIMARY ROOT LENGTH

A previous study showed that 3OC6-HSL promotes the primary root elongation of wild-type Arabidopsis (Jin et al., 2012; Liu et al., 2012). Root growth is controlled by the cell division in meristem zone and the cell expansion in elongation zone. In order to dissect the promoting effect of 3OC6-HSL on primary root elongation, we measured the length of meristem and elongation zones in primary roots 7 days after application of 3OC6-HSL to Arabidopsis root systems. The results revealed that the primary roots of plants treated with 3OC6-HSL had both longer meristem and elongation zones than those of solvent-treated wild type plants (Figures 1A,D). The difference in the length of elongation zone (30.2%) was much more pronounced than the difference in the length of meristem zone (13.6%). To further pinpoint the process whereby 3OC6-HSL functions, we counted the number and calculated the length of cortical cells in the meristem and elongation zone. We found that treatment with 3OC6-HSL caused a 30% increase in the number of cortical cell in the meristem zone compared to the solvent-treated roots while no significant increase in the length of cortical cells in the meristem zone was observed after 3OC6-HSL treatment, indicating that the increase in meristem size is mainly due to the increased number of cortical cells in root meristem (Figures 1B,C). On the other side, analysis for root elongation zone demonstrated that the elongated root elongation zone induced by 3OC6-HSL contained similar number of cortical cells but longer cells than the control, suggesting an increase in cell expansion (Figures 1E,F). Collectively, these results suggest that 3OC6-HSL promotes primary root growth via changes in cell division and elongation.

Previously we demonstrated that Ca²⁺ signal is involved in AHL signaling in plant cells (Song et al., 2011). To further analyze the role of CaM in AHL-mediated primary root growth, the effects of different concentrations of CaM antagonists, such as trifluoperazine (TFP) and N-(6-aminohexyl)-5-chloro-1- naphthalene-sulphonamide (W-7), on the length of primary roots was first investigated. It was found that the growth of primary roots in the plants only treated with higher concentration W-7 (Figure 2A) or TFP (Figure 2B) was significantly reduced; however, the length of primary roots did not significantly change in the plants treated with 0–50 μM concentration of W-7 (Figure 2A) or with 0–1 μM concentration of TFP (Figure 2B). W-5 is the dechlorinated analog of W-7 and is often used as control compound for understanding the specificity of W-7. The result shows that no deleterious effect on root elongation was observed for W-5 at the tested concentration (Figure 2A). Therefore, 50 μM for W-7 and 1 μM for TFP have been considered as a threshold concentration and chosen for subsequent experiments.

Subsequently, we investigated the effect of 3OC6-HSL on root elongation with or without antagonists. Three-day-old seedlings were transferred to 1/2 MS medium plates containing 1 μM 3OC6-HSL and 50 μM W-7 or 1 μM TFP and cultivated for additional 7 days. The plates containing 50 μM W-5 were taken as control for W-7 treatment. The results showed that the 3OC6-HSL-stimulated primary root growth was completely suppressed by addition of 50 μM W-7 while 50 μM W-5 did not affect the effect of 3OC6-HSL on primary root elongation, indicating that the effect of W-7 was attributable to CaM inhibition (Figure 3). Likewise, 1 μM TFP blocked the increase of primary root length in response to 3OC6-HSL as well (Figure 3). These data suggested that CaM has an important role in AHL-stimulated primary root elongation.

TREATMENT WITH 3OC6-HSL PROMOTES THE ACCUMULATION OF THE CaM PROTEIN

Pharmacological analysis implicates that CaM is required for 3OC6-HSL-mediated primary root elongation (Figure 3). Whether the level of CaM in plant cells is influenced by treatment with 3OC6-HSL has not been resolved. To address this point, we examined the level of CaM protein in Arabidopsis roots reacting to 3OC6-HSL. Total protein was extracted from Arabidopsis
FIGURE 1 | Effects of 3OC6-HSL on Arabidopsis root meristem zone and root elongation zone. The length of the root meristem zone (A), the number of cortical cells in the meristem zone (B), the length of the root elongation zone (D) and the number of the cortical cells in elongation zone (E) were measured on the wild type seedlings grown on vertically oriented plates containing with or without 1 μM 3OC6-HSL after 7 days of cultivation. The length of cortical cell in the root meristem zone (C) and in the root elongation zone (F) were calculated by dividing the root meristem or elongation zone length by the number of the cortical cells in the root meristem or elongation zone. Data shown are the average of three independent experiments giving the similar statistically significant results. Each experiment included at least 20 seedlings. Entries with p-values < 0.05 shown with asterisk.

roots exposed to 1 μM 3OC6-HSL at different time points and the overall concentration of CaM protein was determined by western blot and ELISA using specific polyclonal antibody against CaM. Western analysis showed that the concentration of CaM protein in roots increased after treatment with 1 μM 3OC6-HSL and reached the maximum at 6 h post treatment (Figure 4A). Similarly, ELISA result indicated that treatment with 1 μM 3OC6-HSL promoted the accumulation of CaM protein and the CaM concentration reached a maximum 2-fold increase after 6 h of treatment with 3OC6-HSL (Figure 4B). These results provide the evidence for a positive regulation of 3OC6-HSL on CaM protein level.

3OC6-HSL INDUCES THE EXPRESSION OF ALL NINE CaM ISOFORM GENES IN ARABIDOPSIS

The results described above demonstrated that 3OC6-HSL increased the CaM protein accumulation, however the response of CaM genes to 3OC6-HSL treatment remains unknown. Therefore, we next investigated the expression profiles of CaM genes after exposure to 3OC6-HSL using qRT-PCR. The Arabidopsis genome contains nine CaM genes encoding CaM protein isoforms (Zielinski, 2002). Since a high nucleic acid sequence conservation exists among nine AtCaM genes, the primers for each AtCaM gene were generally chosen from the 3′ untranslated regions of individual AtCaM genes in order to ensure the discrimination between mRNA of genes belonging to highly conserved gene family. Total RNA was isolated from Arabidopsis roots treated with 1 μM 3OC6-HSL and used to perform the real-time RT-PCR. The expressions of all nine AtCaM genes were up-regulated significantly by 3OC6-HSL although the extent of induction differed from different CaM isoform genes (Figure 5). Compared to the untreated control, a more than 4-fold increase in transcript of AtCaM1, AtCaM3, AtCaM6, and AtCaM8 was observed after treatment with 3OC6-HSL. The mRNA levels of AtCaM4, AtCaM5, AtCaM7, and AtCaM9 genes were increased up to 3-fold by 3OC6-HSL while only 1.7-fold
increase was observed in the expression of AtCaM2 in response to 3OC6-HSL treatment (Figure 5). In summary, these data indicated that 3OC6-HSL induced the expression of all nine AtCaM genes with variable degree.

**MUTATION OF INDIVIDUAL CaM ISOFORM GENES ABOLISH THE PROMOTION OF PRIMARY ROOT LENGTH BY 3OC6-HSL**

The available pharmacological and molecular evidence showed that CaM might be involved in primary root elongation regulated by bacterial AHL, but direct genetic evidence for the participation of CaM is still missing. To solve this question and pursue the specificity of different AtCaM isoform in participation in AHL signaling in plant cells, we compared the primary root elongation in response to 3OC6-HSL between wild-type and the T-DNA insertional mutant lines of individual AtCaM gene. All mutants for all nine AtCaM genes are confirmed to be homozygous lines by antibiotic resistance screening and RT-PCR (Figure 6D) and exhibit no observed phenotypic differences compared with wild-type plants under normal growth conditions (Figure 6A). As shown in Figure 6, the observed increased root elongation upon 3OC6-HSL exposure in wild-type plant was impaired in all mutation lines of nine AtCaM genes. The length of primary roots in cam1, cam2, cam4, cam5, cam6, cam7, and cam9 after treatment with 3OC6-HSL was similar to that of untreated plants of respective mutants (Figures 6A,B). The primary root in cam3 and cam8 after contact to 3OC6-HSL exhibited even slightly shorter than that of untreated respective mutants, however no significant difference was found between treatment and untreated with 3OC6-HSL after statistical analysis (Figure 6B). Additionally, we measured the effects of 3OC6-HSL on the fresh weight of seedlings of wild-type and the individual AtCaM mutants. An obvious increase in the fresh weight of wild type seedlings was observed after treatment with 3OC6-HSL, whereas the fresh weight of seedlings of all nine AtCaM mutants was similar between treated and untreated plants (Figure 6C). Although the conclusion for role of AtCaM in AHL signaling at genetic level needs to be reinforced by complementary and over-expression analysis, our data provide preliminary evidence that all nine AtCaM genes might be involved in plant response to 3OC6-HSL with respect to primary root elongation.

It has been reported that short-side chain AHLs such as C6-HSL and 3OC6-HSL promoted primary root elongation (von Rad
et al., 2008; Jin et al., 2012; Liu et al., 2012) while long-side chain AHLs (10–16 carbons) such as C10-HSL and C12-HSL inhibited primary root growth but promoted the formation of lateral root and root hair (Ortiz-Castro et al., 2008, 2011). To investigate if the inhibitory effects of C10-HSL and C12-HSL on primary root elongation, we compared the primary root length between wild-type and the T-DNA insertion mutants of nine AtCaM genes after exposure to C10-HSL and C12-HSL. The results indicated that while C10-HSL or C12-HSL inhibited the primary root growth but promoted the lateral root and root hair development of wild type plants at concentration of 50 μM, the responses of the mutants of nine AtCaM genes to C10-HSL or C12-HSL were similar as that of wild type plants with respect to the root architecture (Figure 7). These data suggested that AtCaMs might not participated in the regulation of root growth and development by the long-chain AHLs.

DISCUSSION

As one of the most conserved proteins, CaM is thought to be involved in fundamental cellular processes in both animals and plants. In this study, we provide the pharmacological and molecular genetic evidence for involvement of CaM in plant response to bacterial QS signal, 3OC6-HSL, with respect to primary root elongation. To the best of our knowledge, the connection between CaM and 3OC6-HSL signaling in plant cells has not been reported before.

Plant and bacteria have co-existed for millions of years on the earth and a complex networks consisting of different signaling molecules has been evolved. Accumulating evidence suggest that the mechanisms of plant resistance to pathogens or a plants’ responding to beneficial microbes share some commonalities. On the other hand, bacterial pathogens and symbionts depend substantially on QS to colonize and infect their host plants. AHLs are the most common QS signals among Gram-negative bacteria. Like microbial-associated molecular patterns (MAMPs), AHLs might be the indicative for plants that pathogens are in the surroundings to gather themselves for attack or that mutualistes are about to interact with plant roots. Accordingly, Hartmann and coworkers suggested that AHLs are important modulators for interaction between plant and bacteria (Hartmann et al., 2014). However, the early events in the perception and transduction of AHLs in plant cells remain unclear. Recently, we found that several bacterial AHLs can elicit changes in the levels of intracellular Ca2+, and 10 μM C4-HSL causes a significant, transient increase in the intracellular concentration of free Ca2+ in Arabidopsis root cells (Song et al., 2011). As a mediator protein of Ca2+ signal, CaM is activated by binding Ca2+, inducing a cascade of regulatory events. However, whether CaM participate the

![FIGURE 4](image-url) Effects of 3OC6-HSL on the accumulation of CaM protein in Arabidopsis roots. The plants were grown in hydroponic Hoagland medium with or without 1 μM 3OC6-HSL. Total protein was extracted from roots at different time interval after 3OC6-HSL treatment. (A) The change of CaM protein level detected by western blot. Actin was used as an internal quantification control. (B) The change of CaM protein level detected by ELISA using polyclonal antibody against wheat CaM. The results were normalized to the CaM protein level of untreated samples. Each value was the average of three independent experiments. Entries with p-values < 0.05 shown with asterisk.

![FIGURE 5](image-url) Effects of 3OC6-HSL on the expression of CaM genes in Arabidopsis roots. The plants were grown in hydroponic Hoagland medium with or without 1 μM 3OC6-HSL. Total RNA was extracted from roots at different time interval after 3OC6-HSL treatment and analyzed by quantitative RT-PCR with specific primers for individual AtCaM gene. Values were normalized to the 0 h time point. Data represent the mean from three independent biological replications. Entries with p-values < 0.05 shown with asterisk. Relative expression of genes (ratio) > 1.5 was considered as up-regulation (single asterisk) and ratio < 0.8 was considered as down-regulation (double asterisk).
downstream cascade of Ca\(^{2+}\) signal triggered by bacterial AHLs remain unknown. Pharmacological assay showed that application of CaM antagonist TFP or W-7 into 3OC6-HSL-treated roots of Arabidopsis abolished the stimulatory effect of 3OC6-HSL on primary root elongation, while W-5, an inactive structural analog of W-7, did not affect the effect of 3OC6-HSL (Figure 3).

Control experiments demonstrated that these compounds in the concentration used in this study did not affect the root elongation under normal conditions (Figures 2, 3), indicating that the disappearance of 3OC6-HSL-promoted primary root elongation in presence of TFP or W-7 was due to the inhibited activity of CaM. Then we analyzed the accumulation of CaM protein in plant after treatment with 3OC6-HSL by western blot and ELISA with antibody against CaM protein. It was found that accumulation of the overall CaM protein in Arabidopsis roots increased in response to 3OC6-HSL and reached its maximum 6h after treatment (Figure 4). Thus, the data in this study suggested the involvement of CaM in response of plant to 3OC6-HSL and the role of CaM in 3OC6-HSL-mediated root elongation.

In plant cells, the CaM gene family contains variable members that share a high level of nucleic acid and amino acid sequence identity, and different type of CaM isoforms might has distinct physiological roles. Potato plants have eight CaM genes encoding at least two distinct CaM isoforms, among which the PCM1 gene was up-regulated upon touch and PCM6 during tuberization (Takezawa et al., 1995). Among the five divergent soybean CaM genes, ScaM-4 and ScaM-5 were transcriptionally activated by phytopathogenic bacterial infection or fungal elicitor application, indicating their involvement in defense against pathogen attack (Lee et al., 1995). Yamakawa et al. (2001) isolated 13 CaM genes from tobacco genome and found that levels of individual CaM genes are differentially regulated both transcriptionally and post-transcriptionally when tobacco are exposed to stresses such as pathogen-induced hypersensitive cell death and wounding. The Arabidopsis genome has nine genetic loci encoding CaM protein isoforms. To determine the role of specific AtCaM gene in response of plant to bacterial 3OC6-HSL, the level of transcription for the nine CaM genes in root of wild-type seedling grown on medium with and without 3OC6-HSL was measured using specific primers. We found that the expression of all AtCaM genes were up-regulated after treatment with 3OC6-HSL, although the level of gene expression was variable between different AtCaM members (Figure 5). These data indicated that all these AtCaM isoform genes are 3OC6-HSL -inducible. On the other hand, single AtCaM gene knockout did not exhibit any phenotype difference including primary root growth as compared with the wild-type plants under normal growth conditions (data not shown). These observations together with the data of transcription analysis for nine AtCaM genes after 3OC6-HSL treatment (Continued)
gave the hints for the existence of functional redundancy between different AtCaM members for plant growth.

However, mutation of any single AtCaM gene abolished the stimulatory effect of 3OC6-HSL on primary root elongation and the fresh weight (Figure 6). The presence of other AtCaM gene in one of AtCaM T-DNA insertion mutant seemed not to rescue the increased primary root elongation caused by treatment with 3OC6-HSL. It should be noted that more evidence need to be obtained from the complementary or over-expressing lines of individual AtCaM gene. On the other hand, there exist a possibility that some other genes are perhaps required and play additive or complimentary functions. Nevertheless, the molecular and genetic evidence showed the participation of AtCaM genes in the response of Arabidopsis to bacterial QS signal 3OC6-HSL. At moment, we could not figure out the specificity of individual AtCaM gene in participating the AHL-mediated primary root growth. Liu et al. (2005) found that the expression of nine AtCaM genes (AtCaM 1-9) was differentially regulated by heat-shock (HS) at 37°C. The expression of AtCaM3 and AtCaM7 genes increased, while the expression of AtCaM2, AtCaM5, and AtCaM6 decreased during HS. The levels of AtCaM1, AtCaM4, and AtCaM8 mRNA showed no change during HS. Furthermore, molecular and genetic evidence suggest that endogenous AtCaM3 is a key component in HS signaling in Arabidopsis. On the other hand, Al-Quraan et al. (2010) reported that AtCaM1 and AtCaM5 play important roles in response to HS at 42°C. These inconsistent results implicated the complexity of AtCaM participating in the signal transduction of response to the external stimuli.

Previous studies revealed that AHLs exert concentration and acyl-chain length dependent effects on plant primary root growth (Ortiz-Castro et al., 2008, 2011; von Rad et al., 2008; Jin et al., 2012; Liu et al., 2012; Palmer et al., 2014). Generally, AHLs with short-side chain such as C6-HSL and 3OC6-HSL promoted primary root elongation (von Rad et al., 2008; Jin et al., 2012; Liu et al., 2012) while AHLs with long-side chain (10–16 carbons) such as C10-HSL and 3OC12-HSL inhibited primary root length but promoted the formation of laterous root and root hair (Ortiz-Castro et al., 2008, 2011). In addition, even 3OC6-HSL and C6-HSL which exhibit a stimulating effects on primary root growth at concentration below 10 μM showed an inhibitory effects on root length when higher concentration were used (Palmer et al., 2014). In the present study, we found that the prolonged primary root caused by 3OC6-HSL with the tested concentration was the consequence of increased cell division in meristic zone and increased cell elongation in elongation zone (Figure 1). These observation implicated that plants might perceive and respond to different types of AHLs in a different manner. In this study, we provided the evidence that CaM activation might be the early event in 3OC6-HSL signaling in plant cells. However, the genetic evidence shows that CaM may not participates the inhibition of primary root length caused by application of long-chained AHLs such as C10-HSL and C12-HSL (Figure 7). A more thorough understanding of how CaM regulates plant responses to different types of AHL is of great importance. Such investigation is under way.

Our previous work has shown that G-protein-coupled receptor (GPCR) and heterotrimeric G-protein are involved in AHL-mediated primary root elongation of Arabidopsis, and GPCRs might be the candidate receptor for AHL in plant cells (Liu et al., 2012). Moreover, a transient elevation in [Ca²⁺]i concentration could be induced by a number of AHL in plants and the Ca²⁺ influx from the extracellular space into the cells contributed mainly to the increased concentration of cytosolic free Ca²⁺
(Song et al., 2011). Based on our findings herein, we propose a hypothesis that 3OC6-HSL might first be perceived by GPCR and then activate Gq. Gq activation is closely followed by an increase in Ca^{2+} through opening of Ca^{2+} channels in the plasma membrane. The elevated level of cytoplasmic Ca^{2+} then directly activates CaM and promotes the expression and accumulation of CaM. Activated CaM triggers the downstream cascades that ultimately lead to the specific cellular reaction. Further studies are needed to identify the downstream target proteins of activated CaM and to find how CaM activates its target proteins.

**AUTHOR CONTRIBUTIONS**

Qian Zhao and Chao Zhang contributed equally to this work and performed the pharmacological analysis and molecular, genetic assay, respectively. Yali Huang, Zhenhua Jia and Haili Li have helped with the acquisition, analysis and interpretation of data for the work. Shuishan Song is responsible for the design of the work.

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**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: http://www.frontiersin.org/journal/10.3389/fpls.2014.00807/abstract

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