The Impact of Water Deficiency on Leaf Cuticle Lipids of Arabidopsis

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Arabidopsis (Arabidopsis thaliana) plants subjected to water deficit, sodium chloride (NaCl), or abscisic acid treatments were shown to exhibit a significant increase in the amount of leaf cuticular lipids. These stress treatments led to increases in cuticular wax amount per unit area of 52% to 80%, due primarily to 29% to 98% increases in wax alkanes. Of these treatments, only water deficit increased the total cutin monomer amount (by 65%), whereas both water deficit and NaCl altered the proportional amounts of cutin monomers. Abscisic acid had little effect on cutin composition. Water deficit, but not NaCl, increased leaf cuticle thickness (by 49%). Electron micrographs revealed that both water-deprived and NaCl-treated plants had elevated osmium accumulation in their cuticles. The abundance of cuticle-associated gene transcripts in leaves was altered by all treatments, including those performed in both pot-grown and in vitro conditions. Notably, the abundance of the ECERIFERUM1 gene transcript, predicted to function in alkane synthesis, was highly induced by all treatments, results consistent with the elevated alkane amounts observed in all treatments. Further, this induction of cuticle lipids was associated with reduced cuticle permeability and may be important for plant acclimation to subsequent water-limited conditions. Taken together, these results show that Arabidopsis provides an excellent model system to study the role of the cuticle in plant response to drought and related stresses, and its associated genetic and cellular regulation.
indicating that properties of the cutin polyester or specific monomers likewise play an important role in establishing the water-barrier properties of the cuticle membrane. Although many hypotheses have been set forth to explain the function of cuticle lipid composition and structure in plant drought tolerance, rigorous experimentation that confirm these hypotheses have yet to be reported.

Recent studies establish that many plants respond to water deficit stress through increased cuticular wax deposition (Shepherd and Wynne Griffiths, 2006; Kosma and Jenks, 2007). For example, plants like tree tobacco (Nicotiana glauca), sesame (Sesamum indicum), soybean (Glycine max), and rose (Rosa × hybrida) possessed more leaf wax per unit area after short periods of water deficit (Jenks et al., 2001; Cameron et al., 2006; Kim et al., 2007a, 2007b). In some studies, stress-induced increases in wax amount were associated with major reductions in leaf water-loss rates (Williams et al., 1999, 2000; Cameron et al., 2006). It is still unclear however, which stress-induced changes in the cuticle are most critical for reducing foliar water loss and drought acclimation. Water deficiency also stimulates production of the phytohormone abscisic acid (ABA), which in turn leads to rapid responses like stomatal closure (Bartels and Sunkar, 2005) and the induction of abiotic stress-responsive genes (Shinozaki and Yamaguchi-Shinozaki, 2007). To date it is unclear whether ABA is involved in abiotic stress-induced cuticle lipid accumulation. Data mining of gene expression databases has recently implicated several cuticle-associated genes as ABA responsive (Kosma and Yamaguchi-Shinozaki, 2007). To assess more leaf wax per unit area after short periods of water deficit-treated plants had a relative water content (RWC) of approximately 60%, the 150 mM-treated plants had a RWC of approximately 79%, the ABA-treated plants RWC was approximately 93%, whereas control plants had a RWC of approximately 95%.

All treatments resulted in a significant increase in total wax amount per unit leaf area (Fig. 1), with the most striking increase observed on NaCl-treated and water deficit-treated plants, which had 80% (t(6) = 7.62, P = 0.0068) and 75% (t(6) = 6.04, P = 0.0009) more wax than control plants, respectively. ABA treatment also led to a significant increase in total wax amount, with 32% (t(4) = 6.38, P = 0.0031) and 54% (t(4) = 9.61, P = 0.0007) increases observed on plants that were sprayed with 10 μM or 100 μM ABA, respectively. The leaves used for wax extraction from water deficit- and NaCl-treated plants had a smaller surface area compared to control plants (31% and 42% smaller, respectively), whereas ABA-treated plants did not have smaller leaves. In addition, we observed significant shifts in the proportion of specific wax constituents. As such, the changes in wax profiles resulting from these stress treatments were not due to leaf area effects alone, but most likely involve wax metabolic responses.

All treatments shifted the proportional distribution of wax classes. In general, alkanes accounted for the observed large increases in total wax amount, with 98% (t(6) = 7.98, P = 0.0002) and 93% (t(6) = 6.79, P = 0.0005) increases observed on NaCl-treated and water deficit-treated plants, respectively, and 29% (t(4) = 4.27, P = 0.0129) and 69% (t(6) = 9.61, P = 0.0007) increases on 10 μM and 100 μM ABA-treated plants, respectively (Fig. 1). In general, the increased leaf alkane content was attributable, primarily, to increases in the very-long-chain (C_{29}, C_{31}, and C_{33}) constituents (Fig. 1). Minor but significant increases were observed in other wax class amounts such as fatty acids (t(6) = 2.98, P = 0.0247) and aldehydes (t(6) = 4.67, P = 0.0034) on NaCl-treated plants, ketones (t(6) = 9.88; P < 0.0001) on water deficit-treated plants, and aldehydes (10 μM ABA, t(4) = 3.52, P = 0.0245; 100 μM ABA, t(4) = 4.15, P = 0.0142) on ABA-treated plants. Nonetheless, in all cases, alkanes represented the largest absolute increase of any given wax class.

Impact of Water Deficiency on Arabidopsis Cuticle

RESULTS

Impact of Water Deficit, Sodium Chloride, and ABA on Cuticular Wax

The wax composition of rosette leaves from plants subjected to 150 mM sodium chloride (NaCl), water deficit, or 10 μM or 100 μM ABA treatments was measured to assess the influence of water deficit and associated treatments on wax composition. Waxes of water deficit-treated plants were sampled approximately 8 d after water was withheld. NaCl and ABA treatments were applied three times over an 8 d period. Control plants were subirrigated with water every 3 d, with waxes sampled on the same day as corresponding stress treatments. At the end of the stress treatment period, water deficit-treated plants had a relative water content (RWC) of approximately 60%, the 150 mM-treated plants had a RWC of approximately 79%, the ABA-treated plants RWC was approximately 93%, whereas control plants had a RWC of approximately 95%.

Impact of Water Deficit, NaCl, and ABA on Cutin Monomer Composition

Cutin monomers were sampled from water deficit-, NaCl-, and ABA-treated plants using the same treatments as described above for wax sampling. Water deficit led to a 65% (t(5) = 3.28, P = 0.022) increase in total cutin per unit leaf area, largely manifested as a 67% increase (t(5) = 2.57, P = 0.0498) in C_{18:2} dioic acids (Fig. 2). Additionally, water deficit significantly increased C_{16:0} dioic acid (57%; t_{(5)} = 6.51; P = 0.0013), C_{18:1} dioic acid (52%; t_{(5)} = 5.28; P = 0.0033), and C_{16:0} (9,10,16-dihydroxy acid amounts (104%; t_{(5)} = 3.94, P = 0.0277). The 150 mM NaCl treatment did not result in an increase in...
in the total amount of cutin monomers (Fig. 2). However, salt-treated plants did have significantly higher amounts of C_{16:0} dioic acids ($t(5) = 7.35$, $P = 0.0007$), C_{16:0} (9)-10,16-dihydroxy acids ($t(5) = 4.46$, $P = 0.0007$), and C_{18:0} 18-hydroxy acids ($t(5) = 3.03$, $P = 0.0289$). ABA treatments did not significantly increase the amounts of total measured cutin or any individual constituent (Fig. 2).

**Impact of Water Deficit and NaCl on Cuticle Ultrastructure**

Transmission electron microscopy (TEM) analysis revealed that water deficit-treated plants had thicker cuticles than unstressed control plants (Fig. 3). On average, controls from separate experiments of water deficit-treated and NaCl-treated plants had leaf cuticles $55.3 \pm 14.8$ nm and $55.7 \pm 8.1$ nm thick, respectively. Water deficit-treated and 150 mM NaCl-treated plants had leaf cuticles $82.3 \pm 10.6$ nm and $66.0 \pm 12.1$ nm thick, respectively, representing 49% and 18% increases in cuticle thickness, respectively. Water deficit-treated and salt-treated plants also exhibited higher accumulations of osmium in the cuticle, especially the cuticular layer, as revealed by the darker appearance of treated cuticles in the micrographs (Fig. 3).
Impact of Water Deficit, NaCl, and ABA on the Expression of Genes Associated with Cuticle Production

Transcript profiling from leaves of pot-grown plants showed that several cuticle genes (selected based on compositional analysis) were quite responsive to water deficit, salt, and ABA (Fig. 4). **ECERIFERUM1** (**CER1**), **CER5**, and **MYB41** exhibited significant increases in transcript abundance in water deficit-treated plants, whereas **CER4**, **LACSI1**, **LACSI2**, **ATT1**, and **MYB30** showed significantly lower expression (Fig. 4). In salt-exposed plants, only **KCS6/CER6** was induced 24 h after the potting medium was subirrigated with 150 mM NaCl; whereas **MYB41** had notably lower

![Figure 2.](image)

**Figure 2.** Cutin monomer profiles of Arabidopsis rosette leaves from plants subjected to water deficit (A), 150 mM NaCl (B), 10 μM ABA (C), or 100 μM ABA (D). ABA-treated plants and their respective controls were greenhouse grown. Methanol was used to dissolve ABA, which was diluted to treatment concentrations with distilled, deionized water. Control plants for ABA treatments were sprayed with distilled, deionized water containing an equivalent amount of methanol. NaCl-treated, water deficit-treated, and their respective control plants were grown in a growth room. Bars represent mean values ± SD (n = 3–4, the experiments were repeated once with similar results). Asterisks denote significant differences (*, P < 0.05; **, P < 0.01) as determined by Student’s t tests or Satterthwaite t tests.

![Figure 3.](image)

**Figure 3.** Leaf cuticle membrane (between arrowheads) from leaf number four of control plants (A and C), water deficit-treated (B), 150 mM NaCl-treated (D) plants. The cuticle is divided into an outermost, electron-translucent layer, the cuticle proper, and an innermost, electron-dense, darker-staining layer, the cuticular layer, that is most evident in leaves of stressed plants (B and D). CW, Cell wall; scale bars = 100 nm.
ABA treatment caused (24 h after treatment) an over 2-fold increase in CER1 transcript abundance, but also increased the abundance of CER3/WAX2, CER5, and LACS2 transcripts by about 2-fold (with the LACS1 expression level falling just below the 2-fold increase).

As a comparison to studies using pot-grown plants, cuticle-associated gene expression studies were also performed on plants grown in vitro, and exposed to analogous water deficit, NaCl, and ABA treatments as in the pot-grown plants (Fig. 5). For in vitro water deficit treatments, petri dish lids were removed, exposing plants to the dry atmosphere; RNA extraction was performed 6 and 24 h later. The 6 h water deficit caused an approximately 2-fold or higher increase in 22 of the 24 cuticle-associated gene transcripts, except GPAT8 and ACR4, whereas the 24 h treatment increased, by approximately 2-fold or higher, transcript abundance of 14 of the 24 genes, including KCS6, KCR, CER1, CER4, CER5, WBC11, CER7, LACS2, LACS3, ATT1, LCR, HTH, SHN3, and MYB41 (Fig. 5). CER1 was increased most by the in vitro water deficit treatment. The NaCl treatments caused a approximately 2-fold increase in KCS6, KCR, CER1, CER3, SHN2, SHN3, and MYB41 (Fig. 5). The ABA treatment caused an approximately 2-fold increase in KCS6, CER1, CER3, WBC11, ATT1, SHN2, SHN3, and MYB41 (Fig. 5). NaCl treatment reduced abundance of LACS1 and LCR transcripts, whereas ABA treatment led to lower abundance of GPAT8 and MYB30 transcripts.

Association of Cuticle Composition with Cuticle Permeability and Plant Drought Acclimation

To assess the impact of stress-associated changes in cuticle properties on determinants of plant water status, rosette water-loss and chlorophyll-leaching assays were employed on plants that had received the same treatment schemes as those outlined for wax/cutin sampling. All treatments resulted in a reduced rate of both leaf water loss (Fig. 6) and leaf chlorophyll leaching (Fig. 7). Although the differences were very small, plants subirrigated with 150 mM NaCl could be shown to exhibit the same small reduction in leaf water-loss rate curve in two replicate experiments, suggesting that these differences were significant. It must be considered that supraoptimal levels of salt inflict both hyperosmotic and hyperionic stress on plants (Hasegawa et al., 2000), and that NaCl has also been shown to inhibit dark-induced stomatal closure (Jarvis and Mansfield, 1980). As such, incompletely closed stomata may have influenced the leaf water loss rates of NaCl-treated plants reported here.

We also sought to determine if the cuticle changes induced by exposure to water deficit would improve plant tolerance to subsequent water-limited environments (i.e. whether treatments provide drought acclimation). It was observed that plants subjected to water deficit were better able to withstand subsequent water deprivation, compared to previously nonstressed plants, as evidenced by delayed wilting and the maintenance of a higher RWC (Fig. 8).

DISCUSSION

We report that Arabidopsis plants exposed to a water deficit treatment (wherein irrigation was withheld for approximately 8 d) exhibited a significant increase in total leaf cuticle wax amount of approximately 75% relative to nontreated plants, a value similar to those reported for other plants exposed to water deficit, including both dicotyledonous and graminaceous species (Seiler, 1985; Jefferson et al., 1989; Bondada et al., 1996; Jenks et al., 2001; Samdur et al., 2003; Shepherd and Wynne Griffiths, 2006; Kim et al., 2007a, 2007b; Kosma and Jenks, 2007). The highest reported induction of wax by water deficit occurs on the leaves of tree tobacco, wherein waxes increased by over 150% after exposure to multiple drying events (Cameron et al., 2006). Whether such high wax amounts could be obtained with Arabidopsis using repeated drying cycles is uncertain. Notwithstanding, the wax induction we observed on Arabidopsis is comparable to some of the most responsive plants reported to date. The most notable change in the wax constituent profile of water deficit-treated Arabidopsis leaves is the dramatic increase in alkane constituents, which explains essentially all of the observed increase in the total leaf wax amount. As such, it appears that alkane synthesis is key to this stress response. Besides waxes, we report that water deficit also increases the...
amount of leaf cutin monomers on Arabidopsis by 65%. Previous studies have not examined cutin monomer responses to water deficits, and so, to our knowledge, this report becomes the first to implicate cutin induction as a drought-responsive adaptation. Unlike wax induction, water deficit increased the amounts of nearly all leaf cutin monomers, indicating that more total cutin, rather than more of any specific cutin constituent, may be of greater importance in this water deficit stress response.

Previous studies in rose (Williams et al., 1999, 2000; Jenks et al., 2001) and tree tobacco (Cameron et al., 2006) provide some evidence that water deficit-induced increases in cuticle wax produce a less water-permeable cuticle, an adaptation that may limit transpiration and delay the onset of cellular dehydration stress during prolonged climatic drought (Kosma and Jenks, 2007). Studies presented here demonstrate that water deficit alters both wax and cutin in Arabidopsis, and produces plants with less permeable cuticles. These induced cuticle changes were associated with delayed wilting in Arabidopsis and the maintenance of normal water status during water deprivation treatments. Nonetheless, a contributory role for osmotic adjustment, stomatal regulation, and other physiological changes in acclimated Arabidopsis, rose, and tree tobacco cannot be ruled out. Moreover, whether plants maintain reduced cuticle permeability for an extended period of time after recovery from water deficit, or whether permeability returns to normal nonstress levels at some time after the stress period, has yet to be explored in Arabidopsis or any plant species. As such, further studies are needed to dissect the contribution of cuticle induction relative to other acclimation responses during the acquisition of stress tolerance. As demonstrated here, Arabidopsis can be a preeminent model for these studies.

The distribution, size, and number of wax crystalline domains within the cuticle are thought to define tortuous paths for the diffusion of water, and thereby serve as major determinants of cuticle permeability (Riederer and Schreiber, 1995; Kerstiens, 1996; Buchholz, 2006; Burghardt and Riederer, 2006; Schreiber, 2006; Kosma and Jenks, 2007). A strong negative

Figure 5. Stress profiling of cuticle gene transcripts from in vitro stress treatments. Modulation of cuticle gene expression was determined in 15-d-old Arabidopsis plants subjected to different stress conditions including: water deficit (A), 150 mM NaCl (B), or 10 μM ABA (C). Water deficit treatment consisted of removal of petri dish lids and exposure of plantlets to dry air for 6 or 24 h. All other treatments lasted 24 h. The gene expression level was determined by quantitative RT-PCR analysis. Results are presented as differential relative transcript abundance. The data represent the means ± so of three replicates. KCS6 = CER6, ECR = CER10.
relationship exists between polymer crystallinity and membrane/coating permeability to water (Klute and Franklin, 1958; Lasoski and Cobbs, 1959; Sangaj and Malshe, 2004). Wax alkanes, primary alcohols, and aldehydes are known to confer greater resistance to water movement in artificial membranes than fatty acids and triterpenoids, and among these, alkanes may form the most impermeable crystalline regions (Grncarevic and Radler, 1967). Whether the increased alkane amounts we observe on Arabidopsis following water deficit contribute to the formation of less permeable crystalline structures in Arabidopsis leaf cuticles awaits further study using spectroscopic, calorimetric, atomic force, and NMR approaches. The Arabidopsis

Figure 6. Water-loss rates (expressed as a percentage of initial water-saturated weight) of isolated rosettes from whole plants deprived of water (water deficit; A), subjected to 150 mM NaCl (B), or 10 μM ABA (C). Points represent mean values ± se (n = 5, the experiments were repeated once with similar results).

Figure 7. Chlorophyll-extraction rates (expressed as a percentage of total chlorophyll extracted after 24 h) of rosettes from plants deprived of water (water deficit; A), subjected to 150 mM NaCl (B), or 10 μM ABA (C). Points represent mean values ± se (n = 4, the experiments were repeated once with similar results).
species itself, along with the large collection of diverse cuticle mutants now available (Kosma and Jenks, 2007), may provide an excellent model system to explore these relationships.

As another issue that requires further study, it is yet unclear what role cutin plays in cuticle permeability. Numerous cutin monomer-deficient mutants in Arabidopsis have been reported that show large elevations in cuticle permeability, and from these studies and others (Kosma and Jenks, 2007), it has been postulated that the cutin matrix provides a kind of framework within which waxes are deposited and crystallized. Potentially, mutations that destroy the cutin framework severely disrupt wax crystallization, and subsequently the cuticle’s permeability barrier function. Our results, which show a significant increase in total leaf cutin monomers and an increase in the thickness of both the cuticle proper and cuticular layer of the Arabidopsis cuticle after water deficit, lead us to speculate that drought acclimation involves synthesis of a larger cutin framework that supports more concentrated intracuticular packing of wax crystalline regions. As another consideration, essentially all cutin monomers possess carboxyl or hydroxyl functional groups capable of participating in ester or hydrogen bonds. In synthetic polymers, increased cross-linking between monomers is thought to reduce permeability by decreasing segment mobility and the availability of hydrogen-bonding sites within the polymer (Sangaj and Malshe, 2004). Increased cross-linking of the cutin polymer may prohibit hydrogen bonding of water molecules to unlinked, oxygenated cutin functional groups and in this way slow the diffusion of water. Whether the induced cuticle changes we report here influence the structure of the cutin matrix or internal cross-linking in a way that alters water diffusion pathways in water deficit-stressed Arabidopsis is completely unknown.

We implicate the three genes CER1, CER5, and MYB41 as important water deficit-inducible cuticle-associated genes, which is consistent with their proposed roles in alkane synthesis (Aarts et al., 1995; Jenks et al., 1995), wax export (Pighin et al., 2004), and regulation of wax synthesis (Cominelli et al., 2008), respectively. Interestingly, five cuticle-associated transcripts (CER4, LACS1, LACS2, ATT1, and MYB30) were less abundant in plants experiencing water deficit, whereas only three were elevated by water deficit. The fact that CER4 was not more abundant seems logical, as the 1-alcohols are thought to be synthesized by the CER4 protein and were reduced, as a proportion of total waxes, by water deficit treatment. However, both LACS1 and LACS2 genes (Lü et al., 2009) and the ATT1 gene (Xiao et al., 2004), thought to encode important

Figure 8. Water deprivation acclimation experiment. Approximately 15-d-old plants were exposed to 14 d of water deprivation (dwd) until reaching a RWC of 49% alongside well-watered control plants. Both groups of plants were then rewatered and allowed to recover for 20 h. Plants that recovered from water deficit (acclimated) and control plants (nonacclimated) were then subjected to further water deprivation for up to 15 d post recovery (dpr). Water deficit-acclimated plants were resistant to further water deprivation and were able to maintain a higher RWC compared to nonacclimated plants. ND, Not determined.
genes like genes are responsive to ABA and cuticle-associated genes, have suggested that several cuticle-associated genes are responsive to ABA and cuticle-associated genes like CER6 and ATT1 are reported to contain ABA-responsive element-like elements in their promoters (Hooker et al., 2002; Duan and Schuler, 2005). Whereas we did not observe elevated cutin with ABA treatment, we did record elevated waxes on leaves treated with low and high levels of ABA, with a preferential increase in alkanes as was observed for water deficit. The CER1 transcript was increased most by our ABA treatments. Taken together, these studies suggest that ABA may be necessary for the activation of several cuticle genes, particularly CER1, during water deficit. Comprehensive analysis of gene expression and cuticle composition responses in ABA-associated mutants (Finkelstein et al., 2002; Nambara and Marion-Poll, 2005) may yield valuable information on the requirement of ABA for cuticle stress responses.

Few studies have examined NaCl or other salts in plant cuticle induction. Nevertheless, Suaeda maritima shows a 60% increase in leaf cuticle membrane thickness after salt treatment, and this was associated with a 35% reduction in transpiration rate (Hajibagheri et al., 1983). Although we did not observe a significant increase in cuticle membrane thickness or total cutin monomer amount of salt-treated Arabidopsis, we did observe increased osmium accumulation in cuticles from micrographs of salt-stressed leaves, a decrease in cuticle permeability, increased wax, and, consistent with observations from public databases (Kosma and Jenks, 2007), significant induction of the gene KCS6/CER6 by salt treatments. The KCS6/CER6 gene is thought to function in wax synthesis, and as such, may have contributed to the altered cuticle properties we observed on our salt-treated plants. Whether other treatments at different time points would reveal the involvement of additional genes requires further study. Moreover, salt stress has both an osmotic and ionic component, and so it might be hypothesized that the observed cuticle changes contribute to water conservation by plants growing in salt-saturated soils and/or the shedding of salt spray from foliage via improved cuticle barrier properties. Further study is now clearly warranted.

A higher proportion of genes were induced with in vitro-grown Arabidopsis plants than were induced in pot-grown plants exposed to comparable water deficit treatments, especially at the 6 h time point following exposure to dry air (including those reported by Joubès et al., 2008). This is likely due to the fact that in vitro plantlets have quite poorly developed, diminutive, and often discontinuous cuticles (Wetzstein and Sommer, 1982; Wardle et al., 1983; Hazrika, 2003), creating a need to rapidly synthesize a whole and normally functioning cuticle more adequately suited to the lower humidity levels of ex vitro conditions. By comparison, pot-grown plants with a fully formed cuticle apparently require the induction of only certain genes when water becomes more limited, such as CER1, that target synthesis of a less permeable or drought-acclimated cuticle. Just as for many other treatments, CER1 was induced more than other genes in the in vitro plants exposed to dry air, strengthening our previous stated hypothesis that CER1 is a very important drought-responsive gene.

Collectively, our results demonstrate that Arabidopsis plants respond to water deficit treatment by increasing the deposition of both leaf cuticular waxes and cutin monomers, and increasing the thickness and osmiophilicity of the cuticle membrane. Likewise, NaCl and ABA induce leaf waxes, but in contrast to water deficit have little effect on cutin monomers. The larger increase in waxes versus cutin monomers on treated leaves, particularly the predominant increase in wax alkanes, fits well with previous models that implicate waxes as critical determinants of cuticle permeability (Buchholz, 2006; Burghardt and Riederer, 2006; Kosma and Jenks, 2007). Results here also reveal that stress-induced accumulation of cuticle lipids involves both accumulation and reduction of specific gene transcripts, with CER1 being conspicuously elevated in most treatments. Although these findings are consistent with a hypothesis that cuticle induction provides an adaptive mechanism for plant acclimation to drought, research is still needed to elucidate the cellular and molecular-genetic mechanisms that underlie cuticle-associated stress responses.

**MATERIALS AND METHODS**

**Plant Growth Conditions and Water Deficit, NaCl, and ABA Treatments for Wax and Cutin Chemistry, TEM Analysis, Epidermal Permeability, Water Deprivation Acclimation, and Pot-Grown Gene Expression Experiments**

Arabidopsis (Arabidopsis thaliana; ecotype Columbia-0) seeds were stratified for 3 to 4 d at 4°C, and plants were grown in 3-inch pots of Promix PGX soilless media (Premier Horticulture) at a density of four to six plants per pot. Growth conditions consisted of a growth room at 21°C to 22°C with 30% to 60% relative humidity, a 16/8-h light/dark cycle, and a light intensity of 125 to 150 μmol m⁻² s⁻¹ or a greenhouse during the months of September to February with an average temperature of 22°C, an average relative humidity of 68%, and ambient sunlight supplemented with a combination of high-pressure sodium and metal halide lamps contributing approximately 100 μmol m⁻² s⁻¹ at plant height on a 16/8-h light/dark cycle. Plants were fertilized once per week for the 2 weeks of growth prior to stress treatments with 1,000 mg per liter 15-5-15 of Miracle Gro Excel Cal-Mag (The Scotts Co.) at a pH range of 5.7 to 6.0 with alkalinity reduction achieved via 93% sulfuric acid (Ulrich Chemical) at 0.08 mL per liter.

Fifteen to 16-d-old plants were used for all stress treatments. NaCl stress was imposed by subirrigation with a 150 mS NaCl (Mallinckrodt Baker)
solution, allowing 20 to 30 min for absorption, three times over a 7 d period. Control plants for NaCl-treated plants were watered with greenhouse tap water each time NaCl was applied. For water deficit treatments pots were deprived of water until wilting of lower leaves was observed (typically 7–9 d; RWC approximately 60%). ABA treatment consisted of spraying with 10 μM or 100 μM ABA (Sigma-Aldrich) solution three times over a 7 d period. ABA was dissolved in methanol and treatment solutions prepared by dilution, with distilled, deionized water, to the appropriate volume. Control plants for the ABA treatments were sprayed with distilled, deionized water containing equivalent amounts of methanol. For NaCl and ABA treatments, all biochemical and physiological analyses were performed on the 8th d after initial treatment. For gene expression analysis, rosette leaves were harvested 24 h after 150 mM NaCl and 10 μM ABA application; rosette leaves from water-deprived plants were harvested when leaves showed minor wilting (RWC approximately 60%). Leaves from untreated control plants were harvested at the same time as leaves from treated plants.

For water deprivation acclimation studies, plants were grown in a growth chamber at 22°C to 25°C, under a light/dark 8/16-h photoperiod of white light at 185 to 210 μmol m⁻² s⁻¹. Fifteen to 16-old plants were deprived of water until nearly all plants wilted (typically 14 d) and had reached a RWC of 49%. Control plants grown alongside water deficit-treated plants were watered as needed with tap water. Once water deficit-treated plants had wilted, the soil of both control plants and water deficit-treated plants was saturated with tap water and allowed to recover for 20 h. After 20 h, excess water was poured off both groups of plants. Control (nonacclimated) and water-deprived (acclimated) plants were then subsequently deprived of water until the majority of control plants had wilted (typically 11 d) and had reached a RWC of 56%.

RWC Measurement

Fresh weights of entire rosettes, removed from their roots, were collected on a microbalance. Rosettes were then submerged in distilled, deionized water for 12 h, blotted dry, and saturated fresh weights were collected. Rosettes were then dried in an oven at 80°C until a constant mass was achieved at which point dry weights were collected. Rosette RWC was calculated as described by Barrs and Weatherley (1962). Each replicate consisted of an entire rosette from a single plant. Each replicate was taken from a separate pot. Three to six replicates were used for all analyses.

Cuticular Wax Analysis

Rosette leaf wax composition was determined as described by Chen et al. (2003) with slight modification. Leaves were submersed in hexane for 30 s followed by a short 1 s rinse. Wax extracts were evaporated under N₂ gas and derivatized by heating at 100°C for 15 min in N₂/O₂-bis(trimethylsilyl)trifluoroacetamide (BSTFA; Supelco). Silylated samples were analyzed by gas chromatography/mass spectrometry (GC-MS) with a Hewlett-Packard 5890 series II GC equipped with a flame ionization detector (FID) and a 12-m, 0.2-mm id HP-1 capillary column with helium as the carrier gas. The GC oven was programmed with an initial temperature of 80°C and increased at 15°C min⁻¹ to 200°C, then increased at 2°C min⁻¹ to 280°C. Injector and detector temperatures were set at 320°C. Quantification was based on uncorrected FID peak areas relative to internal standard methyl heptadecanoate peak area. Areas of rosette leaves were determined by ImageJ software (http://rsb.info.nih.gov/ij/) using digital images of flattened leaves.

TEM

Leaf cuticle ultrastructure was analyzed by TEM according to methods employed by Chen et al. (2003). Briefly, leaf samples were collected from the middle of the blade between the midvein and the margin of leaf 4. Samples were dehydrated in alcohol for 1 h at 70%, 80%, and 95% (v/v) glutaraldehyde and 2% (v/v) formaldehyde in 0.05 M phosphate buffer, pH 6.8 (PB; Karnovsky, 1965). Two 40 s microwave treatments (model 3450; Ted Pella) under a vacuum, using the low-power setting, with a 3-min period between the two exposures was used to improve fixation. After fixation, samples were washed with PB and postfixed in 1 h in 2% (v/v) osmium tetroxide in 0.05 M PB. One 40 s microwave treatment under vacuum as described above was applied during osmium fixation. Tissues were then washed with PB and dehydrated through a gradient series of ethanol, infiltrated with Spurr’s embedding medium (Electron Microscopy Sciences), and polymerized for 48 h at 60°C. Ultrathin sections (80–100 nm) were prepared from Spurr’s resin-embedded samples and mounted on carbon-coated Formvar-100 mesh copper grids (Electron Microscopy Sciences). Sections were air-dried, stained, and then stained with 2% (v/v) lead citrate for 3 min and counterstained with 4% aqueous uranyl acetate for 5 min, dehydrated in ethanol and embedded in LR white resin. Sections were observed with a Philips EM 400 electron microscope (FEI Co.). Cuticle thickness measurements were made on digitized micrograph images using ImageJ software (http://rsb.info.nih.gov/ij/). Measurements were made from the base of the darker osmium-stained cuticular layer to the outermost edge of the lighter-stained cuticle proper. Three replicates were used for each treatment and the appropriate control. Each replicate was taken from a separate plant from a separate pot. Each replicate consisted of 10 measurements made on two to three sections from the same leaf sample.

Measurement of Epidermal Permeability

To quantify excised rosette water loss, plants were dark acclimated for 3 h prior to measurement. Whole rosettes were excised (from roots) and placed immediately in water (in the dark) and soaked for 60 min to equilibrate water contents. Rosettes removed from soaking were shaken gently and blotted dry to remove excess water, with weights determined gravimetrically every 20 min using a microbalance. Data were expressed as a percentage of the initial water-saturated fresh weight.

Epidermal permeability was also assessed using chlorophyll efflux. Plants were watered and allowed to rehydrate during a 3-h dark-acclimated period prior to measurement. Entire rosettes were collected and immersed in an equal volumes of 80% ethanol in glass scintillation vials. Vials were covered with a aluminum foil and agitated gently on a shaker platform. Aliquots of 1 mL were removed every 20 min and 24 h after initial immersion. The amount of chlorophyll extracted into the solution was quantified using a UV-2102 PC spectrophotometer (UNICCO) and calculated from UV light absorption at 647 and 664 nm as described by Lolle et al. (1998). Data were expressed as a percentage of the total chlorophyll extracted after 24 h in 88% ethanol.

Cutin Monomer Analysis

Leaf cutin monomer content was analyzed based on methods described by Lü et al. (2009). Ground, delipidized, dried leaf tissues were used for all reactions. Depolymerization reactions consisted of 6 mL of 3% methanolic hydrochloride (Supelco) and 0.45 mL (7%, v/v) methyl acetate (Sigma-Aldrich) at 60°C. Methyl heptadecanoate was used as an internal standard. After 16 h, reactions were allowed to cool to room temperature and terminated by the addition of 6 mL of saturated, aqueous NaCl. Methyl ester monomers were extracted, twice, with 10 mL dichloromethane (Bonaventure et al., 2004). After washing the organic phase three times with 0.9% (w/v) aqueous NaCl, dichloromethane extracts were dried with 2,2-dimethoxypropane (Sigma-Aldrich), and evaporated under nitrogen gas. Monomers were derivatized in pyridine and BSTFA (1:1, v/v) for 15 min at 100°C. Excess pyridine/BSTFA was removed with nitrogen gas, and samples were dissolved in heptane:toluene (1:1, v/v) prior to analysis with a Hewlett-Packard 5890 series II GC equipped with a flame ionization detector (FID) and 12 m, 0.2 mm id HP-1 capillary column with helium as the carrier gas. The GC oven was programmed with an initial temperature of 80°C and increased at 15°C min⁻¹ to 200°C, then increased at 2°C min⁻¹ to 280°C. Injector and detector temperatures were set at 320°C. Quantification was based on uncorrected FID peak areas relative to internal standard methyl heptadecanoate peak area. Areas of rosette leaves were determined by ImageJ software (http://rsb.info.nih.gov/ij/) using digital images of flattened leaves.

Plant Material and Growth Conditions for in Vitro Gene Expression Studies

Arabidopsis (ecotype Columbia-0) was used in all experiments. Seeds were either plated on Murashige and Skoog (MS) medium supplemented with 0.7% agar, 2.5 mM MES-KOH, pH 5.7 and plants were grown under long-day conditions (16 h of light, 8 h of darkness) at 22°C, or placed in liquid 0.5× MS

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medium supplemented with 2% Glc, 20 mM MES-KOH, pH 5.7 and grown under long-day conditions at 22°C on a rotary shaker (120 rpm). The 15-d-old seedlings grown into 0.5x MS medium were osmotically stressed by transferring the plants into new 0.5x MS medium containing 150 mM NaCl or 10 |m| ABA for 24 h. For water deficit treatment, the plates containing the seedlings were opened for 6 or 24 h. After 6 or 24 h of treatment, plants were used for RNA extraction.

RNA and cDNA Preparation

RNA from Arabidopsis tissues with the RNeasy plant mini kit (Qiagen). Purified RNA was treated with DNase I using the DNA-free kit (Ambion). First-strand cDNA was prepared from 1 μg of total RNA with the Superscript RT II kit (Invitrogen) and oligo(dT)18 according to the manufacturer’s instructions. A 0.66-μL aliquot of the total reaction volume (20 μL) was used as a template in real-time reverse transcription (RT)-mediated PCR amplification.

Real-time (Quantitative) RT-PCR Conditions and Analysis

The PCR amplification was performed with gene-specific primers listed in Supplemental Table S1. PCR efficiency ranged from 95% to 105%. All samples were assayed in triplicate wells. Real-time PCR was performed on an iCycler (Bio-Rad). Samples were amplified in a 25-μL reaction containing 1X SYBR Green Master Mix (Bio-Rad) and 300 nM of each primer. The thermal profile consisted of 1 cycle at 95°C for 3 min 30 s followed by 40 cycles at 95°C for 30 s and at 58°C for 30 s. For each run, data acquisition and analysis was done using the iCycler iQ software (version 3.0a, Bio-Rad). The transcript abundance in treated samples relative to untreated samples was determined using a comparative cycle threshold method. The relative abundance of ACT2 mRNAs in each sample was determined and used to normalize for differences of total RNA amount according to the method described by Vandesompele et al. (2002).

Statistical Analysis

Wax and cutin data were analyzed using SAS 9.1.3 software (SAS Institute Inc.). Student’s t tests were used to analyze data when the assumptions of normality and homoscedasticity were met. Welch-Satterthwaite t tests were used to analyze data when assumptions of normality and homoscedasticity were not met.

Sequence data from this article can be found in the GenBank/EMBL data libraries under accession numbers At1g49240, At1g02205, At4g24510, At5g57800, At4g33790, At1g51500, At1g7840, At3g0500, At1g68530, At1g67730, At3g55360, At2g47240, At1g49340, At1g64400, At4g80360, At1g01610, At4g00400, At2g49570, At1g72970, At1g64670, At5g1190, At5g25390, At3g28910, At4g28110, and At3g59420.

Supplemental Data

The following materials are available in the online version of this article.

Supplemental Table S1. Primers used for quantitative RT-PCR of select cuticle gene.

ACKNOWLEDGMENTS

We would like to thank Debra Sherman and Chia-Ping Huang of the Purdue University Electron Microscopy Center. Received May 22, 2009; accepted October 6, 2009; published October 9, 2009.

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