Autoimmunity in Arabidopsis acd11 is mediated by epigenetic regulation of an immune receptor

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Introduction

Unlike vertebrates, plants lack a somatic, adaptive immune system and immunological memory [1]. Therefore, plants rely on a large repertoire of pre-existing immune receptors, encoded by hypervariable Resistance (R) genes, which recognize specific pathogens and activate strong defense responses. These responses include the programmed cell death (PCD) of host cells at infection sites to restrict pathogen access in a process called the hypersensitive response (HR). R proteins are triggered by pathogen-specific effector proteins that have evolved to perturb or disrupt host processes to facilitate infection. While some pathogen effectors are recognized extracellularly, the majority are targeted to various intracellular compartments of the plant host and identified there. In most cases, R proteins are activated by detecting modifications to host proteins targeted by pathogen effectors. This model, known as the “guard hypothesis” [2,3], has been supported in numerous instances. For example, RIN4, a host protein with key roles in basal defense, is under surveillance by multiple R proteins, and at the same time is the target of multiple pathogen effectors [4]. Most R proteins have been classified as NB-LRRs, named after their central nucleotide-binding (NB) and C-terminal leucine-rich repeat (LRR) domains, although various exceptions exist [5]. The N-terminal domains of NB-LRR R proteins fall into two broad categories: those with homology to Drosophila Toll and mammalian Interleukin-1 Receptor (TIR), and those with predicted coiled-coil (CC) regions [6]. Members of the animal NOD-like receptor (NLR) family exhibit similar domain architecture to plant NB-LRRs, and NLRs are likewise involved in immunity [7,8]. Like NB-LRR proteins, NLRs have several types of amino-termini including protein–protein interaction domains associated with proteins involved in programmed cell death and inflammation. Several autoimmune diseases in humans have been associated with mutations in NLRs [9].

In plants, there are numerous examples of mutants with autoimmune-related phenotypes. These so-called “lesion-mimics” are, in many cases, caused by mutations in genes hypothesized to be negative regulators of the HR [10]. Other examples include point mutations in NB-LRR R proteins [11,12]. Since R proteins have the potential to trigger host PCD, their activity is tightly regulated. R genes are typically constitutively expressed at low levels and some are up-regulated in response to pathogen-derived...
Plants defend themselves against pathogens via immune receptors that trigger responses including the suicide of infected cells to limit pathogen growth. The accelerated cell death 11 (acd11) knockout mutant of the model plant Arabidopsis thaliana kills itself in the absence of invading pathogens. By screening for secondary mutations that rescue acd11, we discovered two LAZARUS (LAZ) genes required for death. The first, LAZ2, encodes an enzyme that methylates histones, the major protein component of chromatin. This particular histone modification is generally involved in epigenetic remodeling of chromatin to a more permissive state for transcription of associated DNA. We show that expression of the second gene, LAZ5, is dependent on LAZ2 activity, suggesting that LAZ5 is a direct target of LAZ2. LAZ5 is a member of an immune receptor class involved in detection of specific pathogens and subsequent cell death. We propose that acd11, and other suicidal mutants, result from autoimmunity triggered by immune receptors controlled by chromosomal modifications. Interestingly, we found that defects in LAZ2 result in enhanced susceptibility to bacterial pathogens, suggesting that it controls other genes involved in innate immunity.

Results

lac2 suppresses cell death in acd11

To isolate genes required for cell death in acd11, Landsberg erecta (Ler) ecotype acd11-1 plants harboring the NahG transgene were mutagenized with ethyl-methanesulfonate (EMS), diepoxybutane (DEB) or γ-irradiation. ~200 suppressors of acd11 were subsequently identified as plants that survived following BTH treatment. Genetic analyses of 43 such suppressors grouped them into 12 recessive and 2 dominant loci referred to as lazarus (lac) mutants, after the biblical resurrection. One of the lac mutants found in the suppressor screen, lac2, abolished cell death in response to BTH in the acd11 NahG background, and exhibited similar levels of cellular ion leakage as wild type (Fig. 1, A and B). lac2-1 acd11-1 NahG plants also exhibited abnormal development (e.g. early flowering, increased shoot branching) that, along with acd11 suppression, was inherited recessively (data not shown). Two other lac2 alleles with similar morphology, lac2-2 and lac2-3, were confirmed by complementation tests (Fig. S1A). Global transcript profiles of lac2-1 acd11-1 NahG, Ler wild-type, NahG, and acd11-1 NahG plants were acquired by hybridizing total mRNA, isolated before and 72 h after BTH treatment, to Affymetrix ATH1 GeneChip arrays. lac2-1 exhibited dramatic suppression of the top 500 most significantly regulated genes in acd11-1 after 72 h BTH (Fig. S2A) In addition, a strong negative Pearson correlation of r = −0.87 was obtained for global expression fold change between lac2-1 acd11-1 and acd11-1, indicating that gene expression in acd11-1 was strongly affected by the lac2-1 mutation (Fig. S2B).

The LAZ2 locus was identified using a map-based approach. Briefly, Ler lac2-1 acd11 NahG was crossed to Columbia ecotype (Col-0) acd11 NahG to generate a segregating F2 mapping population after BTH treatment. Ecotype-specific linkage markers were used to map lac2-1 to a ~150 kb region at the bottom of chromosome 1 (Fig. S3). Candidate genes were selected and sequenced based on annotated mutant phenotypes at The Arabidopsis Information Resource (TAIR; http://www.arabidopsis.org), revealing an irradiation-induced 28-bp deletion in the third exon of the gene Atlg77300 (Fig. 2A). This locus was also sequenced in lac2-2 acd11-1 NahG, revealing an EMS-induced G to A transition converting tryptophan 1536 to a premature stop.

LAZ2 encodes the histone methyltransferase SDG8

Sequence analysis revealed that LAZ2 encodes the histone lysine methyltransferase (HMT) SET (Su(var)3-9, E(z) and Trithorax-conserved) DOMAIN GROUP 8 (SDG8), otherwise known as EARLY FLOWERING IN SHORT DAYS (EFS) and CAROTENOID CHLOROPLAST REGULATORY 1 (CCR1) [21,22]. The mutation in lac2-1 causes a frame-shift just upstream of the sequence encoding the conserved SET associated cysteine-rich domains, while that in lac2-2 introduces a stop codon upstream of a motif conserved within the RB1 subunits of RNA polymerase II [23]. SDG8 is homologous to yeast SET2, which is associated with methylations at histone 3 lysine 36 (H3K36). Another yeast HMT, SET1, modifies H3K4. Both H3K4 and H3K36 methylations are typically associated with active transcription [24]. While Arabidopsis has 43 annotated SDG proteins, SDG8 groups with H3K36-specific HMTs in fungi and animals along with 4 other Arabidopsis proteins [25]. During transcription in
yeast, SET1 and SET2 are recruited to active chromatin by the RNA polymerase II-associated PAF1 complex, where they promote gene expression by facilitating chromatin opening, thus enhancing transcription initiation and elongation, respectively [26]. A similar mechanism seems to be conserved in Arabidopsis based on studies of sdg mutants. SDG8 was first identified as a gene that controlled flowering time via its activity on the transcription of the key floral repressor FLOWERING LOCUS C (FLC), an epigenetically regulated MADS box transcription factor (TF) [27,28]. Expression of the FLC paralog MADS AFFECTING FLOWERING 1 (MAF1) is also dependent on SDG8, which is required for di- and trimethylation of H3K36 [25]. In addition to flowering time, SDG8 regulates carotenoid composition and shoot branching via modification of chromatin at specific loci [22,29]. Our microarray expression analysis revealed that MAF1 and CRTISO, both recently confirmed as direct targets of SDG8 [22,25], exhibited very low expression levels in the absence of LAZ2 (Fig. S4). Deficient expression of these and similar genes likely contributes to the developmental phenotypes observed in laz2 [22]. A number of acd11 suppressors found in the same screen as laz2 were dominant. One of these, laz5 Dominant 1 (laz5-D1), was mapped to a region close to this R gene (Fig. S5). Sequencing of At5g44870 in laz5-D1 revealed a G to A transition at the splice donor site (+1 position) of intron 4 likely resulting in deletion of exon 5 (Fig. 3B). To confirm that this mutation resulted in suppression of acd11, two allelic dominant suppressors, laz5-D2 and laz5-D3, were sequenced: both had lesions in At5g44870 (below), hereafter referred to as LAZ5.

LAZ5 encodes a TIR-class NB-LRR of unknown pathogen specificity with sequence similarity to RPS4 (Fig. S6), an R protein conferring resistance to Pseudomonas syringae expressing the effector AvrRPS4 [30]. The DEB-induced laz5-D2 mutation is a T to A transversion changing isoleucine 287 to asparagine (I287N). This mutation is within the P-loop motif of the NB domain essential for coordination of bound nucleoside triphosphates [5]. The EMS-induced point mutation in laz5-D3 (G811E) lies in the LRR domain, which provides pathogen recognition specificity and has been implicated in R protein activation [31]. Accelerated cell death in acd11-1 was suppressed by laz5-D1 and laz5-D2 (Fig. 3C), and laz5-D alleles suppressed acd11 cell death irrespective of BTH induction or the presence of NahG (Fig. 3D). Furthermore, over-expression of laz5-D2 or laz5-D3 (35S::laz5-D2 or 35S::laz5-D3) suppressed acd11 death after induction, confirming that dominant negative mutations in LAZ5 are responsible for suppression of the acd11-dependent autoimmune response (Fig. S7).

Transgenic plants over-expressing R-genes can exhibit spontaneous cell death and/or constitutive defense responses [32]. In agreement with these observations and the phenotype associated with deletion of ACD11, over-expression of wild-type LAZ5

Figure 1. laz2 suppresses cell death in acd11. A, 21-day-old acd11-1 NahG and laz2-1 acd11-1 NahG plants 1 week after treatment with 100 μM BTH. Size bar = 0.5 cm. B, Ion leakage cell death assay of leaf discs from 5-week-old Ler wild-type (WT), acd11-1 NahG and laz2-1 acd11-1 NahG plants after BTH treatment. Means ± s.d. were calculated from 4 discs per treatment with 4 replicates within an experiment. doi:10.1371/journal.ppat.1001137.g001
(35SLAZ5) in the Col-0 background resulted in 30 out of 38 transgenic plants exhibiting acd11-like cell death which did not survive to set seed (Fig. S8). Since LAZ5 transcription is likely dependent on SDG8 HKMT activity, and the suppression of acd11 by laz2/sdg8 is recessive, we predicted that a loss-of-function mutation in LAZ5 would suppress acd11 in a recessive manner. As expected, a null T-DNA insertion mutant of At5g44870 (SALK_087262; here termed laz5-1) suppressed acd11-2 cell death recessively in plants without NahG (Fig. 4A). A second T-DNA insertion mutant allele of LAZ5 (SAIL_874-D10) also suppressed cell death in acd11-2 (data not shown). Expression of LAZ5 was assayed by real-time PCR in wild-type, laz5-1, and sdg8-2 plants 24 hours after syringe inoculation with the virulent bacterial pathogen Pseudomonas syringae tomato (P.s.t.) DC3000 or with 10 mM MgCl2 (mock control). While pathogen treatment induced LAZ5 expression in wild type, transcript levels in sdg8-2 were comparable to that in the laz5-1 null mutant (Fig. S9A). This confirms the microarray expression data shown in Fig. 3A. The apparent lack of LAZ5 expression in sdg8-2 was seen in several independent experiments with plants at different stages and/or treated with other pathogen strains (data not shown). Moreover, ACD11 expression was unaffected in laz5-1 and sdg8-2 (Fig. S9B), and transcript accumulation of several TIR-NB-LRR-encoding genes homologous to LAZ5 was seemingly unaffected in 3-week old sdg8-2 plants compared to wild-type control with the possible exception of At5g45230 (Fig. S10).

An important question is whether LAZ5 is the relevant target of SDG8 required for acd11 cell death. To help answer this question, we transformed laz2-1 acd11-1 NahG plants with a genomic construct of LAZ5 under control of a constitutive promoter and monitored cell death by ion leakage after BTH treatment compared to relevant controls (Fig. S11). LAZ5 over-expression restored cell death in leaf discs between 3 and 8 days after induction, indicating that lack of LAZ5 expression in sdg8 is a major cause of the suppression of acd11 cell death. However, it cannot be excluded that other targets of SDG8 histone methyltransferase activity also contribute to BTH-induced cell death in acd11.

SDG8 directly modifies chromatin at the LAZ5 locus

To test whether laz2 directly affects histone methylation at the LAZ5 locus, chromatin immunoprecipitation (ChIP) was conducted using antibodies against specifically modified histones. In laz2-1 acd11-1 NahG, trimethylated (me3) H3K36 levels were reduced in chromatin associated with the 5' coding regions of MAF1 (control) and LAZ5, when compared to the acd11-1 NahG control (Fig. 4B). Enrichment of H3K36me3 in LAZ5 chromatin was not influenced by BTH treatment or acd11 homozygosity (Fig. S12A). This suggests that activation of cell death in acd11 does not result in hyper-trimethylation at H3K36, but rather that this histone modification is required for proper LAZ5 expression. There was no effect of genotype on levels of total H3 (Fig. 4C). H3K36me3 is not a general mark for genes up-regulated in acd11, such as FMO1 [18], since we found no enrichment at FMO1 chromatin 72 h after BTH induction (Fig. S12B, C). Moreover, absence of LAZ2/SDG8 had no effect on H3K36me3 levels at the constitutively expressed ACTIN locus (Fig. 4C) or the MAP KINASE KINASE 4 (MKK4) locus (Fig. S12D).

To elucidate H3K36 methylation status irrespective of acd11 and NahG, we also conducted ChIP assays on sdg8-2 single mutant

Figure 2. LAZ2 encodes a histone lysine methyltransferase that is essential for cell death in acd11. A, SDG8 gene with locations of the laz2-1 deletion, laz2-2 mutation, and sdg8-2 T-DNA insertion (SALK_026642). Boxes and lines represent exons and introns, respectively. 1 kb = 1000 base pairs. B, Col-0 WT, sdg8-2, acd11-2 and sdg8-2 acd11-2 plants 4 weeks after germination.

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and Col-0 wild-type seedlings. It was previously shown that loss of SDG8 resulted in both a decrease in global H3K36me3 levels and a coincident increase in global monomethylated (me1) H3K36, a mark associated with transcriptional repression in Arabidopsis [25]. In wild-type plants, MAF1 and LAZ5 chromatin was enriched for H3K36me3, whereas the level of H3K36me3 was diminished in sdg8-2 (Fig. 4D). Conversely, H3K36me1 levels at these loci were higher in sdg8-2 and reduced in wild type. Treatment of seedlings for 3 hours with an HR-inducing bacterial pathogen had no effect on the methylation status of H3K36 (data not shown). Also, H3 trimethylation of LAZ5 chromatin at other lysine residues (K4, K9, K27), was not affected by loss of SDG8 (Fig. S12E).

**Figure 3.** acd11 autoimmunity requires LAZ5, a TIR-NB-LRR immune receptor. A, Expression of AtSG44870 (LAZ5) in Ler WT, acd11-1 NahG and laz2-1 acd11-1 NahG before and 72 h after treatment with 100 μM BTH relative to Ler WT at time 0 (log2 scale). Microarrays were performed on duplicates or triplicates of independent RNA preparations from aerial parts of 4-week-old plants before and 72 h after BTH treatment. LAZ5 is significantly differentially expressed between the genotypes (p = 7e-7) and over time (p = 0.005) as determined by Two-Way ANOVA. B, LAZ5 with locations of mutations in 3 laz5-D alleles and the 2 T-DNA insertions laz5-1 (SALK_087262) and laz5-2 (SAIL_ B74-D10). Boxes and lines represent exons and introns. Domains encoded by exons are marked TIR (A), NB (B), LRR (C), C-terminal extension (D). Asterisk marks the region amplified for ChIP and short arrows represent flanking primers. 500 bp = base pairs. C, Ion leakage death assay of leaf discs from 5-week-old Ler WT, acd11-1 NahG, laz5-D1 acd11-1 NahG, and laz5-D2 acd11-1 NahG after treatment with 100 μM BTH. Means ± s.d. were calculated from 4 discs per treatment with 4 replicates in an experiment. D, Ler WT, acd11-1 and laz5-D1 acd11-1 plants 3 weeks after germination.

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SDG8 is required for pathogen resistance in Arabidopsis

To determine whether SDG8 and/or LAZ5 are required for basal resistance to virulent pathogens, leaves of 4-week old sdg8-2, laz5-1, wild-type and an allele of enhanced disease susceptibility 1 (eds1-2 introgressed into Col-0) mutants were syringe-inoculated with P. stutzeri DC3000 and growth was assayed after 4 days. Bacteria grew to ~9-fold higher titers in sdg8-2 than in wild-type or laz5-1, while titers in eds1 were yet another order of magnitude higher (Fig. 5A). Growth of another strain of bacterial pathogen, Pseudomonas syringae maculicola (P.s.m.) ES4326, was tested on sdg8-2, laz5-1, wild-type and eds1 with similar results (Fig. 5B). We did not observe elevated bacterial growth in sdg8-2 when we used P. stutzeri DC3000 expressing HR-inducing AvrRpm1, AvrRpt2, AvrRps4 or AvrPphB and growth was assayed after 3 or 4 days. Bacterial titers were ~15-fold higher in sdg8-2 than in wild-type or laz5-1 for P. stutzeri expressing AvrRpm1 (Fig. 5C). This suggested that RPM1-mediated resistance is defective in sdg8-2. To confirm this, growth of P. mela ES4326 expressing AvrB was assessed after 3 days: AvrB is also recognized by RPM1, and resistance to this avirulent pathogen was affected in sdg8-2 to a similar level as P. stutzeri with AvrRpm1 (Fig. 5D). In both cases, bacterial titers were comparable to the rpm1-3 null mutant [34]. Defects in SDG8 had a consistent, yet statistically insignificant effect on growth of P. stutzeri DC3000 expressing AvrPphB, (Fig. S13B) resistance to which is dependent on the R gene RPS5 [35]. In addition, sdg8-2 did not affect RPS2- or RPS4-mediated resistance to AvrRpt2 [36,37] (Fig. 5E) and AvrRps4 [30] (Fig. 5F). Corroborating the pathogen growth assay, transcript levels of RPM1 and RPS5 were low or absent in 4-week old sdg8-2 compared to wild-type, whereas expression of RPS2 and RPS4 in sdg8-2 was similar to that in wild-type (Fig. S13C). Defects in LAZ5 did not have a detectable effect on transcript accumulation of RPM1, RPS5, RPS2 or RPS4 (data not shown). As with LAZ5, we conducted ChIP assays at the RPM1 locus in untreated seedling tissue from laz2-1 acd11-1 NahG versus acd11-1 NahG (in Ler) and sdg8-2 versus wild-type (in Col-0). We observed lower H3K36me3 and higher H3K36me1 levels at RPM1 chromatin in the absence of functional LAZ5 and/or SDG8, indicating that RPM1 is an example of another R gene that is regulated by histone methylation (Fig. S14). These results indicate that SDG8 targets a subset of R genes and other genes involved in more general aspects of basal defense.

Figure 4. SDG8 regulates LAZ5 and modifies chromatin at the LAZ5 locus. A. Representative 3-week-old Col-0 WT, acd11-2 (in Col-0) and laz5-1 acd11-2 plants. B. ChIP analysis of MAF1 and LAZ5 with 1 µg anti-H3K36me3 antibody (IP) or no Ab (mock) expressed as proportion of input material in eluates after IP with appropriate Ab (% input), mean ± s.d. (n = 3). Tissue was from 3-week-old acd11-1 NahG and laz2-1 acd11-1 NahG seedlings (Ler background) before and 72 h after treatment with 100 µM BTH. The experiment was repeated twice with similar results. C. Total H3 levels at MAF1, LAZ5 and ACT7 chromatin, and H3K36me3 levels at ACT7, are not affected by laz2-1 as determined by ChIP analysis with 1 µg anti-H3K36me3 antibody or 1 µg anti-H3 (total) antibody, presented as EtBR-stained PCR product (34 cycles). D. ChIP of MAF1 and LAZ5 with 1 µg anti-H3K36me3 antibody (IP) or no Ab (mock) expressed as % input, mean ± s.d. (n = 3). Tissue was from 3-week-old homozygous sdg8-2 and Col-0 WT seedlings. The experiment was repeated twice with similar results.

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Autoimmunity Mediated by an HKMT and NB-LRR

A. P. syringae DC3000

B. P. syringae ES4326

C. P. syringae + AvrRpm1

D. P. syringae + AvrB

E. P. syringae + AvrRpt2

F. P. syringae + AvrRps4

G. RPM1

RPS2

RPS4

Relative expression

Col-0 WT  sgdβ-2

Col-0 WT  sgdβ-2

Col-0 WT  sgdβ-2
Figure 5. SDG8 is required for plant innate immunity. A. Growth of virulent P.s.t. DC3000 in Col-0 WT, sgd8-2, laz5-1 and eds1 0 and 4 days after infiltration with bacteria at OD600 = 0.0001. B. Growth of virulent P.s.m. ES4326 in 3-week old Col-0 WT, sgd8-2, laz5-1 and eds1 plants 0 and 4 days after infiltration with bacteria at OD600 = 0.0001. C. Growth of avirulent P.s.t. DC3000 expressing AvrRpml in Col-0 WT, sgd8-2, laz5-1 and rpm1-3 plants 0 and 3 days after infiltration with bacteria at OD600 = 0.001. D. Growth of avirulent P.s.t. ES4326 expressing Avr8 in Col-0 WT, sgd8-2, laz5-1 and rpm1-3 plants 0 and 3 days after infiltration with bacteria at OD600 = 0.001. E. Growth of avirulent P.s.t. DC3000 expressing AvrRpt2 in Col-0 WT, sgd8-2, laz5-1 and ndr1 0 and 3 days after infiltration with bacteria at OD600 = 0.001. F. Growth of avirulent P.s.t. DC3000 expressing AvrRps4 in Col-0 WT, sgd8-2, laz5-1 and eds1 plants 0 and 4 days after infiltration with bacteria at OD600 = 0.00005. Log-transformed values are means ± s.d. (n = 6). Asterisks indicate statistical significance (p < 0.0001) determined by unpaired Student’s t-test. The experiments were repeated twice with similar results. cfu = colony forming units. G. Transcript accumulation of RPM1, RP52, and RPS4 in 3-week-old Col-0 wild-type and sgd8-2 plants, as determined by qRT-PCR. Data is normalized to ACT1 (ACT1) and presented as relative expression compared to Col-0 = 1.0, mean ± s.d. (n = 3).

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Discussion

Chromatin remodeling has emerged as a complex regulator of transcription and an epigenetic mechanism to maintain lasting changes in gene activity states. Dynamic post-translational modifications of various residues of histones tails, including methylation, phosphorylation, acetylation, and ubiquitination, play important roles in both promoting and repressing gene expression by recruiting histone binding proteins and chromatin remodeling enzymes [38]. The combinatorial nature of histone modifications results in a complex “histone code” that adds an important level of control to fine-tune gene-specific responses to broader transcriptional inputs [39]. Changes in chromatin state may therefore modulate gene expression in a context-dependent manner to maintain a flexible response to pathogen attack. In plants, this process has been proposed as a mechanism for priming SA-responsive loci during systemic acquired resistance to pathogens [40].

So far, relatively few studies directly addressed epigenetic processes related to chromatin modification to plant innate immunity and/or PCD. Defects in HISTONE DEACETYLASE 19 (HDAC19) and HISTONE MONOUBIQUITINATION 1 (HUB1) increase susceptibility to necrotrophic fungal pathogens in Arabidopsis [41,42]. Furthermore, defects in genes involved in histone variant replacement, and the variant H2AZ itself, result in increased resistance to virulent bacterial pathogens, some spontaneous cell death, and up-regulation of defense genes [43]. More commonly, the “memory” of chromatin remodeling activity is observed as increased levels of open chromatin marks (H3Ac, H3K4me2, etc) at the promoters of many SA-responsive genes, such as PATHOGENESIS-RELATED 1 (PR-1) and WRKY TFs [40,44,45]. The clearest example of immune response at the level of chromatin comes from Alvarez-Venegas and colleagues, who showed that the HKMT ARABIDOPSIS TRITHORAX-RELATED-ED PROTEIN 5 (ATXR5, also known as SDG15) and ATXR6/SDG34 are H3K27-specific monomethyltransferases essential for transcriptional repression in heterochromatin [53]. Further studies should examine if other predicted H3K36-specific HKMT’s, namely SDG4, SDG7, SDG24 and SDG26, have any role in H3K36 monomethylation, trimethylation and/or antagonistic control of expression of LAZ5 and other genes with roles in immunity or are required for cell death in add11. Moreover, further work is required to determine the mechanisms by which SDG8-dependent changes in H3 methylation regulates the expression of specific genes.

A clue to the function of LAZ5 activation comes from the isolation of our screen of dominant alleles. This indicates that the mutant form (laz5-D) of the R protein likely interferes with activity of the wild-type copy since plants heterozygous for the laz5 null mutation do not suppress add11, indicating haplosufficiency of LAZ5. Dominant negative activity has been described for mutations in the R gene N from tobacco, and indeed for a point mutation (G216E) in the P-loop motif of N [54]. N was later found to oligomerize in the presence of a Tobacco mosaic virus elictor, likely through interaction of TIR domains [55]. This oligomerization was an early event in pathogen perception and was independent of mutations that have an effect on HR induction. Therefore, it is possible that laz5-D mutants form inactive oligomers with wild-type LAZ5 and/or accessory proteins. An example of this scenario from animal innate immunity comes from NOD2, an NLR involved in recognition of bacterial cell wall components: an endogenously truncated form, NOD2-S, interacts with other genes with roles in immunity and are required for cell death in add11. Moreover, further work is required to determine the mechanisms by which SDG8-dependent changes in H3 methylation regulates the expression of specific genes.
mutants were placed into complementation groups, and even fewer were mapped. Therefore, a recessive laz5 knockout allele may exist among our unmapped suppressors.

In this study we have identified the chromatin modifying enzyme SDG8, and its specific target LAZ5, as regulators of autoimmune cell death in acd11. Furthermore, sdg8 mutants exhibit enhanced susceptibility to virulent and avirulent pathogens, whereas laz5 mutants do not, suggesting that other targets of SDG8 are important for general resistance. We also show that transcription of a subset of R genes, including LAZ5 and RPM1, is likely to be directly or indirectly dependent on LAZ5 activity. One scenario that may account for the enhanced susceptibility of sdg8 mutants to virulent pathogens could be the consequence of SDG8 action on multiple NB-LRR loci. If the suite of effectors delivered by Pseudomonas triggers a weak R gene response, in sdg8 a subset of these do not accumulate and thus are no longer available to signal for defense against the invading pathogen. Intriguingly, SDG8 is not expressed until 8 days after germination [28], a stage preceding the initiation of cell death in acd11. SDG8 may therefore developmentally regulate targets such as LAZ5, and may exemplify a key difference in the programmed defenses required during seed maturation and the inducible defenses used during plant growth.

Lesion mimic mutants such as acd11 are useful tools in the genetic dissection of innate immunity in plants [10]. Whereas several of these mutants have putative roles in ceramide signaling or synthesis [59,60] or auto-activate R proteins [11], the majority of lesion mimic mutants represent proteins with no straightforward connection to PCD. Milder autoimmunity, associated with constitutive activation of defense responses and dwarf morphology without coincident HR, can similarly be the result of point mutations in immune receptors (Zhang et al., 2005), or deletion of signaling intermediates such as MAP kinases [61]. Knockout mutants that eliminate host guardees mimic the effects of pathogen effectors, and have been found to exhibit R-gene-dependent lethality [62]. Therefore, it is possible that many lesion mimic/ autoimmune mutants may correspond to gene functions that are guarded by NB-LRRs. If so, the diverse functions of these genes may be “red herrings” not directly related to PCD but only implicated in this process due to their targeting by pathogen effectors. Such may be the case for acd11, although we have been unable to detect any interaction between full-length or truncated LAZ5 and ACD11 in yeast or in planta (data not shown). Previously, we reported the identification of ACD11-interacting proteins [63], which we are testing for interaction with LAZ5. Two predictions about wild-type products of autoimmune mutants emerge from this model. First, suppressor screens should identify R genes. Second, pathogen effectors should target them either directly or indirectly via interacting partners or products of their activities. We currently have no evidence that ACD11 is targeted by pathogen effectors, or that ACD11 contributes to disease resistance in the absence of LAZ5. While future work may strengthen this hypothesis, an alternative model is that ACD11 is involved in negatively regulating SA-dependent expression of LAZ5 (or a subset of R genes) perhaps via some lipid signal.

Materials and Methods

Plant material and growth conditions

Arabidopsis plants were grown on soil or MS-agar plates at 21°C with an 8 h or 12 h photoperiod. sdg8-2 (SALK_065642) and laz5-1 (SALK_007262) T-DNA insertion lines, both previously described as null mutants [23,64], were generated by SIGnAL [65] and obtained from the Nottingham Arabidopsis Stock Centre (NASC; Nottingham, UK). Homozygous genotyping primers were 5’- TAAAGAGGGTCGTGATCATG-3’ with 5’-CAGTGTTCCA- GTAAACGTCG-3’ for sdg8-2 and 5’-TATGTTTTCCAGATGCCAG- 3’ with 5’-ATCGATGTTCAACGTGAG-3’ for laz5-1. Sequences of primers used to detect acd11-1, acd11-2, and NahG are available upon request.

Suppressor screen

Three lots of 920–950 mg Ler acd11-1 NahG seeds were incubated for 4 hr in either 0.74% (w/v) EMS (Sigma-Aldrich, St Louis, MO, USA) prepared in 0.1M sodium phosphate buffer, pH 5, with 5% DMSO, or 10 nM DEB (Sigma-Aldrich) in water, followed by rinsing. γ-irradiation of 300 mg acd11-1 NahG seeds was performed at the Risø Reference Laboratory (Denmark) with 500 Gy from a Cobalt-80 source. M1 plants were grown in families of 125 individuals, 3500 M2 plants per family were screened for BTH-resistant suppressors. ~3 million M2 plants from 845 M1 pools or ~100.000 M1 plants were scored. Putative mutants were genotyped to be homozygous for acd11-1 by PCR.

Ion leakage assay

Conductivity assays were conducted essentially as previously described [66].

Microarray hybridization

Total RNA was isolated from three independent biological replicates of relevant genotypes at 0 and 72 hr after BTH treatment. RNA was labeled and amplified according to the MessageAmp Biotin-enhanced kit (Ambion) protocol and hybridized to 51 ATH1 GeneChips after Affymetrix protocols.

Chromatin immunoprecipitation and real-time PCR

ChIP antibodies purchased from Abcam (Cambridge, UK) included anti-H3 (ab1791), anti-H3K36me1 (ab9048), anti- H3K36me3 (ab9050) and anti-H3K27me3 (ab6002). ChIP antibodies against H3K4m3 (pAb-056-050) and H3K9me3 (pAb-003-050) were purchased from Diagenode (Liège, Belgium). Quantitative PCR primers for ChIP analysis were LAZ5 5’-GAATGTCGTGGCAAGTTATTCCATG-3’ with 5’- GAAGATGTCGGACGACTTGGATTC-3’. FMO1: 5’-CTCTAGATTCCTCTACTATG- 3’ with 5’-CTCTATTTTTGGGCATGGAAG-3’. MIF1: 5’-CGCCCTTTTGAGATTTGGAAGC-3’ with 5’-GGGGATTTCCAGAGAATCTG-3’. ACTIN: 5’-GGGAACATCGTCTCTAGGG-3’ with 5’-ACCCAGATAGACAGACAC-3’. ChIP was performed essentially as described [67], using 1μg of each antibody. Real-time PCR to quantify the immunoprecipitated DNA was performed using Brilliant II SYBR Green qPCR kit (Stratagene), and reactions were run on an iCycler IQ (Bio-Rad, Hercules, CA, USA). In all cases, ChIP values were calculated using the Delta-Delta Ct (ΔΔCt) algorithm to determine relative gene expression utilizing the ‘percent input method’. Briefly, signals obtained from the ChIP were divided by signals obtained from an input sample representing the amount of chromatin used in the ChIP. The % ‘input’ value shows what proportion of this starting material is found in the eluate after IP with appropriate Ab.

For expression analyses, RNA was extracted from relevant genotypes using the Qiagen RNeasy RNA extraction kit followed by DNase treatment as per the manufacturer’s instructions. Equal amounts of RNA were subjected to one-step real-time PCR using data not shown.

Chromatin immunoprecipitation and real-time PCR
Cloning and generation of transgenic plants

3.9-kb fragments of laz2-D alleles were amplified from genomic DNA (laz5-D1 acl11-2 NahG, laz5-D2 acl11-2 NahG, laz5-D3 acl11-2 NahG) and cloned into modified pCAMBIA-3300 as described [68], using a uracil-excision based cloning technique (USER, New England Biolabs). Cloning primers were 5′-gctttaaATGGCG-GCATCTTCCGAAATACC-3′ and 5′-ggtnaaATTACAAATAA-CGAGTATAAATTG-3′. A 3.9-kb fragment of LAZ5 was amplified from genomic DNA (wild type Ler), cloned into pENTR/D-TOPO (Invitrogen) and transferred to Gateway-compatible constitutive expression vectors pGBWB502 or pGBW521 [69] by LR recombination reaction (Invitrogen). Cloning primers were 5′-CACCATGCGCAATCCCTCGGAAATACC-3′ and 5′-TTAGGCTAATCCGGTGATT-3′. The final constructs were verified by sequencing, electroporated into Agrobacterium tumefaciens strain GV3101 and used to transform acl11-1 NahG or wild type plants by floral dip method [70]. Transgenic plants were selected on soil with glufosinate (35S/laz5-D alleles) or on MS-agar plate with 20mg/L hygromycin B followed by transplanting to soil (35S/LAZ5).

Supporting Information

Figure S1 laz2 alleles and sg8 share morphological phenotypes, such as early flowering. A, 16-day-old Ler acl11-1 NahG plants homozygous for 3 different laz2 alleles. B, 21-day-old Col-0 WT plants homozygous for sg8-2. Found at: doi:10.1371/journal.ppat.1001137.s001 (3.77 MB TIF)

Figure S2 Transcriptome analysis of laz2-1 suppression of the BTH-induced response in acl11-1 A. The effect of laz2-1 on 355 significantly over-expressed genes among the top 500 differentially expressed genes in response to BTH treatment in acl11-1 NahG plants. B, Scatterplot of global expression fold change comparison between acl11-1 NahG versus NahG (y-axis) and laz2-1 acl11-1 NahG versus acl11-1 NahG (x-axis) 72 h after BTH induction. Found at: doi:10.1371/journal.ppat.1001137.s002 (0.41 MB TIF)

Figure S3 Ecotype-specific markers used to map the LAZ2 locus to ~120 kb on the bottom of chromosome 1. Left is centromeric, right is telomeric. Relative positions of markers are indicated, as are numbers of recombinants remaining at each marker position. Figure shows a rough (~1 megabase) and fine (~150 kb) map of the laz2-1 locus and detail of genomic region between final recombinants, with associated genes and BAC clones. A star marks the LAZ2 gene with the defect determined by sequencing. Found at: doi:10.1371/journal.ppat.1001137.s003 (0.19 MB TIF)

Figure S4 Expression of A) CRTISO (At1g06820) and B) MAF1 (At1g77070) in Ler WT, NahG, acl11-1 NahG and laz2-1 acl11-1 NahG before and 72 h after treatment with 100 μM BTH relative to WT at time point 0 (log2 scale). Found at: doi:10.1371/journal.ppat.1001137.s004 (0.12 MB TIF)

Figure S5 Ecotype-specific markers were used to map the LAZ5 locus to ~80 kb on the bottom of chromosome 5. Left is centromeric, right is telomeric. Relative positions of mapping markers and numbers of recombinants are indicated. Figure shows a map of the laz5-D1 locus and the genomic region between final recombinants, with associated genes. Asterisk marks the LAZ5 gene with the defect determined by sequencing. Found at: doi:10.1371/journal.ppat.1001137.s005 (0.24 MB TIF)

Figure S6 Alignment of LAZ5 and the five most similar Arabidopsis TIR-NB-LRR R proteins, as determined by The Functional and Comparative Genomics of Disease Resistance Gene Homologs Project (http://niblrrs.ucdavis.edu/TN_TNL_phylogeny.html). Sequences include RPS4 (At3g15290), CSA1 (At1g17880), At4g36150, At5g43200, and At5g45230. Mutated residues in laz5-D2 and laz5-D3 are highlighted. Asterisks indicate amino acids predicted to be absent due to the splice site mutation in laz5-D1. Found at: doi:10.1371/journal.ppat.1001137.s006 (1.35 MB TIF)

Figure S7 Over-expression of dominant negative laz5-D alleles suppresses acl11. Figure shows acl11-2 NahG (in Col-0) control and representative transgenic lines of acl11-2 NahG stably transformed with A) 35S/laz5-D2 or B) 35S/laz5-D3, 10 d after treatment with 100 μM BTH. Found at: doi:10.1371/journal.ppat.1001137.s007 (1.70 MB TIF)

Figure S8 Over-expression of the wild-type LAZ5 R gene results in cell death. Figure shows Col-0 wild-type control and two representative transgenic lines of Col-0 stably transformed with a construct over-expressing genomic LAZ5 (35S/LAZ5). Found at: doi:10.1371/journal.ppat.1001137.s008 (2.43 MB TIF)

Figure S9 Expression of A) LAZ5 and B) ACL11 in 3-week-old Col-0 wild-type, laz5-1 and sg8-2 mutant plants 24 h after infiltration with P.s.t. DC3000 at OD600 = 0.01 or 10 mM MgCl2 mock control, as determined by qRT-PCR. Data is normalized to ACTIN1 (ACT1) and presented as relative expression (fold) compared to Col-0 mock = 1.0 (dashed line), mean ± s.d. (n = 3). Found at: doi:10.1371/journal.ppat.1001137.s009 (0.15 MB TIF)

Figure S10 Transcript accumulation of LAZ5 homologs in 3-week-old Col-0 wild-type and sg8-2 plants, as determined by qRT-PCR. Data is normalized to ACT1 and presented as relative expression compared to Col-0, mean ± s.d. (n = 3). Found at: doi:10.1371/journal.ppat.1001137.s010 (0.23 MB TIF)

Figure S11 Ion leakage cell death assay of leaf discs from 3-week-old WT, laz2-1 acl11-1 NahG, acl11-1 NahG and laz2-1 acl11-1 NahG over-expressing LAZ5 plants after BTH treatment. The former were selected segregating T2 plants from a transgenic line of genomic LAZ5 in expression vector pGBW521, and confirmed by qRT-PCR. Data is presented as fold change in conductivity (μS cm⁻¹) relative to initial value at Day 3. Means ± s.d. were calculated from 6 discs per treatment with 4 replicates within an experiment. Found at: doi:10.1371/journal.ppat.1001137.s011 (0.13 MB TIF)

Figure S12 A, H3K36me3 at LAZ5 chromatin is independent of acl11. ChIP analysis of LAZ5 with 1 μg anti-H3K36me3 antibody (IP) or no Ab (mock) expressed as % input. Tissue was from 3-week-old NahG, acl11-1 NahG and laz5-1 acl11-1 NahG seedlings (Ler background) before and 24 h after treatment with 100 μM BTH. B, H3K36me3 is not a general mark for genes up-regulated in acl11. Expression of FMO1 (At1g19259) in Ler WT, acl11-1 NahG and laz2-1 acl11-1 NahG before and 72 h after treatment with 100 μM BTH relative to wild-type at time point 0 (log2 scale). C, ChIP analysis of FMO1 with 1 μg anti-H3K36me3 antibody

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Data is normalized to and 4 days after infiltration with bacteria at OD600 = 0.001. Log-H3K36-me3 antibody or 1 forming units. were repeated once or twice with similar results. cfu = colony 6
sbg8-2
Col-0 wild-type and expressing bacteria at OD600 = 0.001. Ta3 appropriate antibody. In parallel, ChIP samples were used as templates for PCR at the transcriptionally repressed transposon Tas3 locus for comparison. Data is presented as EbR-stained PCR product (34 cycles).

Figure S13 A. Growth of non-pathogenic P.s.t. DC3000 hrcC-mutant in Col-0 WT and sdg8-2 0 and 3 days after infiltration with bacteria at OD600 = 0.001. B. Growth of avirulent P.s.t. DC3000 expressing AvrPphB in Col-0 WT, sdg8-2, laz-5-1 and nfrd1 plants 0 and 4 days after infiltration with bacteria at OD600 = 0.001. Log-transformed values are means ± s.d. (n = 6). The experiments were repeated once or twice with similar results. cfu = colony forming units. C. Transcript accumulation of RPS5 in 3-week-old Col-0 wild-type and sdg8-2 plants, as determined by qRT-PCR. Data is normalized to ACTIN1 (ACT1) and presented as relative expression compared to Col-0 = 1.0, mean ± s.d. (n = 3).

Figure S14 A. ChIP analysis of RPM1 with 1 μg anti-H3K36-me3 antibody (IP) or no Ab (mock) expressed as proportion of input material in the eluate after IP with appropriate Ab (% input), mean ± s.d. (n = 3). Tissue was from 3-week-old nfrd1-1 NhaD and nfrd1-1 NhaG seedlings (Leu background) 72 h after treatment with 100 μM BTH. The experiment was repeated twice with similar results. B. ChIP of RPM1 with 1 μg anti-H3K36-me1 antibody (me1), 1 μg anti-H3K36-me3 antibody (me3), or no Ab (mock) expressed as % input, mean ± s.d. (n = 3). Tissue was from 3-week-old homozygous sdg8-2 and Col-0 WT seedlings. The experiment was repeated with similar results.

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Author Contributions

Conceived and designed the experiments: KP ST FGM BKF PB MP JM. Performed the experiments: KP ST FGM BKF PB DH. Analyzed the data: KP ST FGM BKF PB MP JM. Contributed reagents/materials/analysis tools: HBN. Wrote the paper: KP. Supervised the project: JM.

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