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OPEN

Induction of distinct plant cell death programs by secreted proteins from the wheat pathogen *Zymoseptoria tritici*

Thomas Welch^{1,2}, Carlos Bayon³, Jason J. Rudd³, Kostya Kanyuka⁴ & Graeme J. Kettles^{1,2}✉

Cell death processes in eukaryotes shape normal development and responses to the environment. For plant–microbe interactions, initiation of host cell death plays an important role in determining disease outcomes. Cell death pathways are frequently initiated following detection of pathogen-derived molecules which can lead to resistance or susceptibility to disease depending on pathogen lifestyle. We previously identified several small secreted proteins (SSPs) from the wheat-infecting fungus *Zymoseptoria tritici* that induce rapid cell death in *Nicotiana benthamiana* following Agrobacterium-mediated delivery and expression (agroinfiltration). Here we investigated whether the execution of host cells was mechanistically similar in response to different *Z. tritici* SSPs. Using RNA sequencing, we found that transient expression of four *Z. tritici* SSPs led to massive transcriptional reprogramming within 48 h of agroinfiltration. We observed that distinct host gene expression profiles were induced dependent on whether cell death occurs in a cell surface immune receptor-dependent or -independent manner. These gene expression profiles involved differential transcriptional networks mediated by WRKY, NAC and MYB transcription factors. In addition, differential expression of genes belonging to different classes of receptor-like proteins and receptor-like kinases was observed. These data suggest that different *Z. tritici* SSPs trigger differential transcriptional reprogramming in plant cells.

Multicellular organisms frequently sacrifice individual cells during specific developmental stages or in response to environmental cues. Forms of programmed cell death (PCD) such as apoptosis and autophagy are a normal part of growth and development and also contribute towards the recycling of nutrients. In flowering plants, initiation of PCD plays important roles in processes as diverse as temperature stress, hypoxia, organ development and response to biotic stimuli^{1–3}.

In plant–pathogen interactions, cell death is an essential part of the plant immune system². In interactions with biotrophic pathogens, the active triggering of host cell death termed the hypersensitive response (HR) is often associated with disease resistance⁴. HR is considered an orderly form of PCD, characterised by DNA laddering, organelle fragmentation and cell shrinkage^{5–7}. Induction of HR is assumed to both deny a nutrient supply and spatially restrict invading pathogens⁸. Activation of HR is mediated by plant disease resistance (R) proteins directly recognising secreted pathogen virulence proteins (effectors) or through recognition of effector action on other host proteins^{9–11}. In contrast to HR, necrosis or other uncontrolled forms of cell death are often beneficial to pathogens. Necrotic tissue is characterised by the rupture of plasma membrane and release of cytoplasm to the extracellular spaces. Necrotrophic pathogens in particular benefit from the release of nutrients during necrosis. The importance of control of cell death is illustrated by the variety of pathogen-produced molecules which interfere with these processes. Biotrophic or hemibiotrophic pathogens frequently secrete effectors that are able to suppress activation of immune stimulation that may lead to HR¹². In contrast, necrotrophic pathogens produce necrotrophic effectors that actively trigger host cell death pathways¹³. The lifestyle of each pathogen determines whether induction of cell death has a beneficial or detrimental outcome to the host.

The ascomycete fungus *Zymoseptoria tritici* (*Z. tritici*) causes Septoria tritici blotch (STB) disease of wheat (*Triticum aestivum*) and is a major threat to wheat productivity globally¹⁴. *Z. tritici* is hemibiotrophic, with

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infection typically being symptomless for 10–14 days, before a rapid transition to necrotrophic phase of the life cycle¹⁵. This is initially characterised by leaf chlorosis, followed by the appearance of necrotic lesions in infected areas and sometimes even death of infected leaves. There is considerable transcriptional reprogramming both in host plants and in the fungal cells during infection^{16–18}. Wheat responses are characterised by downregulation of defence-related genes during the early symptomless phase, followed by upregulation of many of the same genes during the transition to necrotrophy^{16,17}. In the fungus, there is upregulation of numerous secreted proteins that are likely to function as effectors and in genes associated with production of secondary metabolites^{16,17}.

In previous work, we identified > 100 *Z. tritici* small secreted proteins (SSPs) that were upregulated during the switch from symptomless to necrotrophic growth¹⁶. These were classed as candidate effectors that might be involved in the induction of cell death during this transition. We used the model plant *Nicotiana benthamiana* to identify a number of SSPs with ability to induce macroscopic cell death in leaves^{19,20}. We found that 13 SSPs induced cell death, and that for 12 of these, initiation of cell death required protein localisation to the apoplastic space. Further, for a smaller group of SSPs (Zt9, Zt11, Zt12) we showed that cell death required the Brassinosteroid Insensitive 1 (BRI1)-Associated Receptor Kinase 1 (BAK1) and Suppressor of BIR1-1 (SOBIR1) receptor-like kinases (RLKs). Both BAK1 and SOBIR1 are important co-receptors for the initiation of intracellular signalling following perception of extracellular ligands. These ligands are frequently microbe-associated molecular patterns (MAMPs) or apoplastic effectors. This indicated that initiation of cell death in response to the *Z. tritici* SSPs occurs at the cell-surface and is likely dependent on recognition by currently unidentified cell-surface immune receptors. In contrast, Zt6 was identified as a ribonuclease toxin that initiates cell death independent of BAK1/SOBIR1²⁰. Moreover, Zt6 induced cell death irrespectively of whether it was secreted to the apoplast or localised to the cytoplasm. Zt6 was demonstrated to have RNase activity against rRNA and display a broad toxicity against monocot and dicot plants, yeast and bacteria, though not to *Z. tritici* itself²⁰.

Based on previous results, we hypothesised that *Z. tritici* effectors may trigger different immune pathways in *N. benthamiana* that ultimately lead to macroscopically similar cell death phenotypes. To test this hypothesis, we used a transcriptomic approach (RNA sequencing, RNA-seq) to investigate early host responses to transient expression of a group of previously described *Z. tritici* SSPs that induce cell death in either a BAK1/SOBIR1-dependent or -independent manner.

Results

RNAseq overview. We aimed to determine the changes in the *N. benthamiana* transcriptome that occur preceding cell death driven by non-host recognition of three (Zt9, Zt11, and Zt12) *Z. tritici* SSPs, and a secreted phytotoxic RNase (Zt6) in comparison to a green fluorescent protein (GFP) control. GFP, Zt6 and SSPs were transiently expressed in *N. benthamiana* leaves using agroexpression and samples were collected at 24- and 48-h post-inoculation (hpi), i.e. prior to the HR becoming visible by eye. RNA was extracted from treated leaves to produce 30 RNA-seq libraries (five treatments x three biological replicates x two timepoints) for sequencing by paired-end sequencing on the Illumina HiSeq 2000 platform. The libraries contained 23.95–35.42 million raw reads. Subsequent quality filtering reduced the number of reads in each library by 37.4% to 51.15%. Of the remaining reads, 92.8% to 98.9% were successfully mapped to the reference *N. benthamiana* genome²¹ (Table 1).

Principle component analysis (PCA) of the overall gene-expression profile showed that replicates of each treatment clustered tightly, as well as revealing minimal difference between the GFP and SSP treatments at 24hpi (Fig. 1A). This pattern changed by 48hpi, with SSP treatments showing clear separation compared to the GFP control, although with minimal difference among themselves. In contrast, Zt6 expression induced a different gene-expression profile compared to both the GFP and SSP treatments at 24hpi. This was further exaggerated by the 48hpi timepoint (Fig. 1A).

To examine these different responses, differential expression analysis was conducted using DESeq2 Bioconductor package²², with FDR less than 0.05, to identify differentially expressed genes (DEGs) between the SSP and Zt6 treatments compared to the GFP control (Fig. 1B). At 24hpi across all treatments, more DEGs were upregulated than downregulated. At 48hpi, the ratio of upregulated and downregulated DEGs was similar, although the total number of DEGs greatly increased. The increase in number of DEGs was most noticeable for the Zt9 treatment, which had a much smaller number of DEGs at 24hpi than the other treatments (Fig. 1B).

The profound difference between the Zt6 and SSP treatments observed by PCA (Fig. 1A), was reflected in the number of DEGs shared between treatments (Fig. 2). For example, the number of upregulated genes common to all three SSP treatments at 48 h (1294) was far higher than that of any other group of three treatments at 48hpi that contained Zt6 (732, 131, 127 DEGs respectively, Fig. 2A). A similar pattern was observed for downregulated genes, although one particular Zt6-containing group of treatments (Zt11/Zt9/Zt6 vs GFP) at 48 hpi shared a much larger number of DEGs (1069) than other groupings, and almost as much as the SSP treatments group (Zt9/Zt11/Zt12 vs GFP, 1104 DEGs, Fig. 1B). Noticeably, at 48 hpi the Zt11 treatment shared far more DEGs with the Zt6 treatment (482 and 808 upregulated and downregulated respectively) than either of the other two SSP treatments, but at 24hpi was much more similar to the Zt12 treatment (Fig. 1).

Gene ontology enrichment analysis. In order to examine whether differentially expressed genes (DEGs) were involved in specific developmental processes, we performed a GO-enrichment analysis of up- and down-regulated genes for the Zt6 (Tables 2, 3) and SSP treatments (Tables 4, 5). For brevity, only the top 20 GO terms (i.e. those with the largest number of DEGs averaged across treatments) enriched in the SSP treatments are listed here.

Most significantly enriched GO terms were present across several treatments, and most were enriched only at the 48hpi time point. Among upregulated DEGs, only four GO terms were enriched in more than one treatment at 24hpi, whilst none were enriched among downregulated DEGs. Out of all significantly enriched GO

| Treatment | Replicate | No. raw reads | No. reads post-filtering | Mapped reads | % alignment rate |
|--------------------------------|-----------|---------------|--------------------------|--------------|------------------|
| Green fluorescent protein 24 h | 1 | 29,632,800 | 15,043,938 | 14,602,207 | 98.59 |
| Green fluorescent protein 24 h | 2 | 28,034,184 | 16,374,738 | 15,431,491 | 95.73 |
| Green fluorescent protein 24 h | 3 | 27,135,887 | 13,275,593 | 12,882,194 | 98.56 |
| Green fluorescent protein 48 h | 1 | 30,478,092 | 18,756,098 | 17,567,294 | 95.26 |
| Green fluorescent protein 48 h | 2 | 30,961,071 | 16,482,238 | 15,930,691 | 98.34 |
| Green fluorescent protein 48 h | 3 | 23,947,546 | 14,644,113 | 13,453,249 | 93.53 |
| Zt6 24 h | 1 | 27,366,516 | 16,894,171 | 16,337,539 | 98.38 |
| Zt6 24 h | 2 | 29,777,551 | 18,459,699 | 17,046,698 | 93.79 |
| Zt6 24 h | 3 | 32,521,194 | 20,013,222 | 19,366,353 | 98.34 |
| Zt6 48 h | 1 | 26,459,619 | 15,042,329 | 13,746,564 | 92.83 |
| Zt6 48 h | 2 | 28,336,919 | 17,480,929 | 16,911,429 | 98.40 |
| Zt6 48 h | 3 | 31,071,683 | 18,719,718 | 17,363,364 | 94.21 |
| Zt9 24 h | 1 | 29,426,692 | 15,565,703 | 15,122,756 | 98.76 |
| Zt9 24 h | 2 | 27,278,801 | 15,849,440 | 14,386,160 | 92.20 |
| Zt9 24 h | 3 | 32,763,879 | 16,106,357 | 15,677,521 | 98.83 |
| Zt9 48 h | 1 | 33,367,108 | 20,400,605 | 18,581,986 | 92.58 |
| Zt9 48 h | 2 | 27,783,195 | 14,402,672 | 13,992,594 | 98.68 |
| Zt9 48 h | 3 | 24,128,224 | 14,765,579 | 13,204,618 | 90.88 |
| Zt11 24 h | 1 | 25,960,835 | 13,004,520 | 12,668,292 | 98.89 |
| Zt11 24 h | 2 | 28,971,906 | 17,874,511 | 17,309,244 | 98.52 |
| Zt11 24 h | 3 | 24,640,400 | 12,036,157 | 11,725,777 | 98.91 |
| Zt11 48 h | 1 | 25,904,899 | 14,658,337 | 14,215,966 | 98.54 |
| Zt11 48 h | 2 | 20,601,920 | 10,108,513 | 9,836,307 | 98.90 |
| Zt11 48 h | 3 | 30,912,455 | 18,370,079 | 17,786,924 | 98.32 |
| Zt12 24 h | 1 | 27,944,558 | 16,863,219 | 15,503,985 | 93.32 |
| Zt12 24 h | 2 | 26,596,304 | 15,083,099 | 14,507,692 | 98.15 |
| Zt12 24 h | 3 | 32,596,510 | 20,162,208 | 18,479,220 | 93.10 |
| Zt12 48 h | 1 | 30,851,089 | 18,800,164 | 18,208,649 | 98.46 |
| Zt12 48 h | 2 | 35,424,590 | 22,184,733 | 20,792,722 | 95.27 |
| Zt12 48 h | 3 | 32,486,965 | 19,870,839 | 19,252,930 | 98.51 |

Table 1. Overview of the RNA-seq data collected for each treatment.

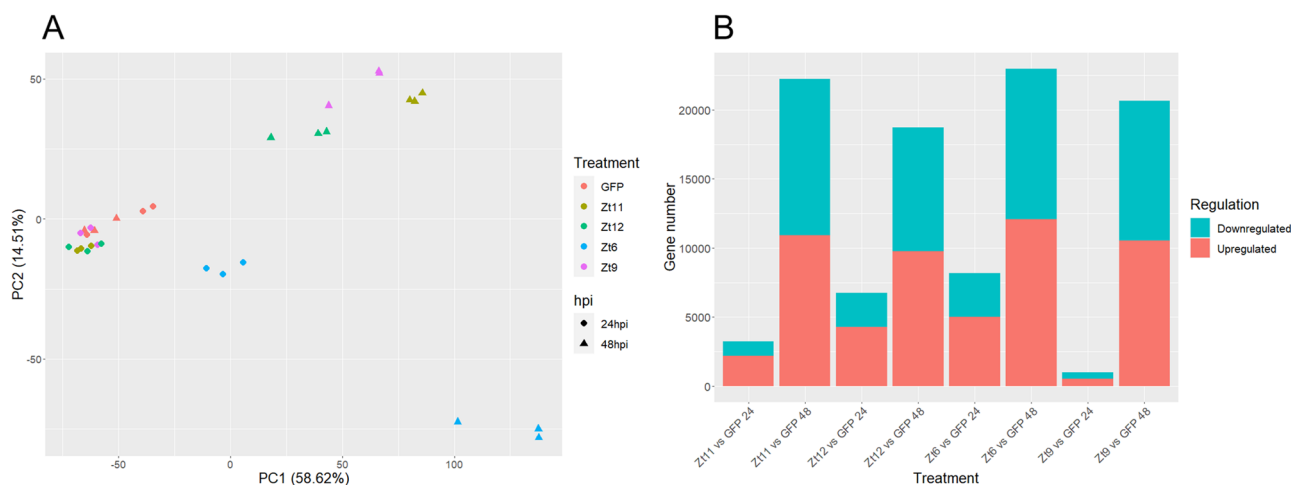


Figure 1. (A) PCA plot of RNA-seq data. SSP treatments and timepoints indicated. (B) Total numbers of upregulated (red) and downregulated (blue) DEGs.

terms (70 in downregulated DEGs and 113 in upregulated DEGs) 37 were enriched only at the 24hpi time point of only one treatment. Noticeably, 20 of these were enriched only in the Zt12 treatment, while eight, seven, and two were exclusive to the Zt11, Zt9, and Zt6 treatments respectively. Only one of the eight GO terms exclusive to Zt11 (sulfate reduction) was enriched in the downregulated DEGs.

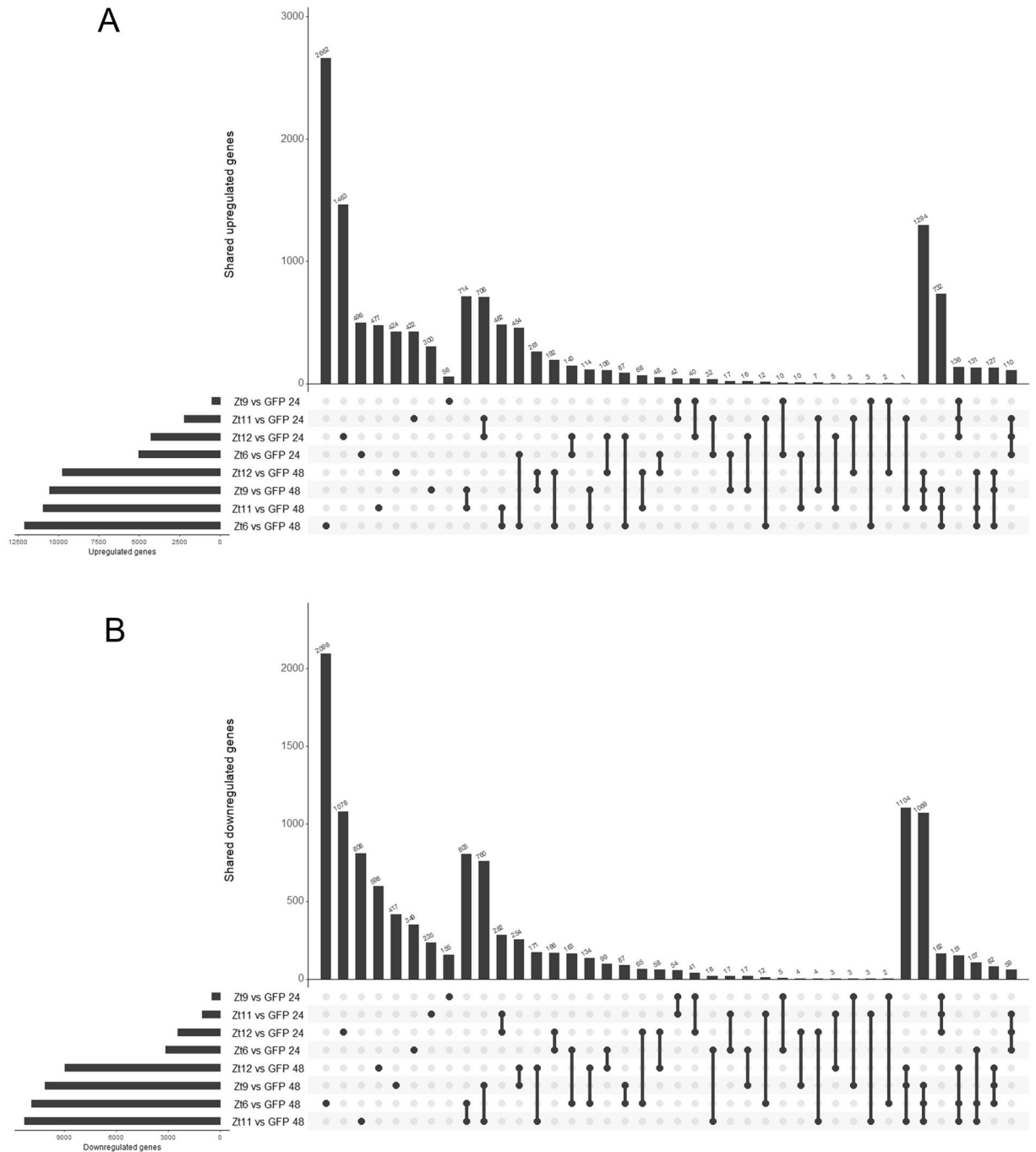


Figure 2. Plots showing total numbers of significantly upregulated (A) and significantly downregulated (B) DEGs. Numbers of DEGs shared among one, two, and three treatment combinations are represented by bar size in each main plot, with the individual treatments that make up each combination represented in the bead and line chart below. Total numbers of DEGs in each individual treatment are represented by bar size on the lower left of each plot. Plots generated using UpsetR⁶³.

The contrasting transcriptional response to SSP and Zt6 treatments revealed by PCA (Fig. 1A) was also observed in terms of enriched GO categories. Among the upregulated DEGs, only seven GO terms were enriched in all treatments at 48hpi (structural constituent of ribosome, translation initiation factor activity, intracellular, ribosome, translation, metabolic process, and ribosome biogenesis) (Tables 2, 4). Most other GO terms were enriched in either one or all SSP treatments, or they were exclusive of the Zt6 treatment. There were only two terms (hydrolase activity, and catalytic activity) that were common to Zt6 and at least one other treatment. This pattern was similar in downregulated DEGs, where only nine GO terms were enriched across all treatments (carbohydrate metabolic process, extrinsic component of membrane, fructose 1,6-bisphosphate 1-phosphatase activity, membrane, photosynthesis, light harvesting, photosystem I reaction centre, photosystem II, and photosystem II oxygen evolving complex) (Tables 3, 5) also in this case at 48hpi. Strikingly, these nine common GO

| Category | Ontology | GO term | Zt6 24hpi | Zt6 48hpi |
|------------|----------|---|-------------|-------------|
| GO:0005622 | CC | Intracellular | – | 1.61E–16 |
| GO:0016772 | MF | Transferase activity, transferring phosphorus-containing groups | 1.47E–17 | 0.012755501 |
| GO:0006468 | BP | Protein phosphorylation | 1.05E–17 | 0.005818144 |
| GO:0004672 | MF | Protein kinase activity | 1.24E–17 | 0.005818144 |
| GO:0003824 | MF | Catalytic activity | – | 0.012632623 |
| GO:0005524 | MF | ATP binding | 1.11E–05 | – |
| GO:0003735 | MF | Structural constituent of Ribosome | – | 1.85E–22 |
| GO:0006412 | BP | Translation | – | 1.85E–22 |
| GO:0006355 | BP | Regulation of transcription, DNA-templated | 6.74E–09 | 2.65E–05 |
| GO:0008152 | BP | Metabolic process | – | 0.005084615 |
| GO:0005840 | CC | Ribosome | – | 1.57E–21 |
| GO:0004674 | MF | Protein serine/threonine kinase activity | 3.08E–18 | 2.34E–06 |
| GO:0005515 | MF | Protein binding | 0.001049327 | – |
| GO:0003700 | MF | DNA-binding Transcription factor activity | 8.46E–20 | 4.24E–07 |
| GO:0043565 | MF | Sequence-specific DNA binding | 5.52E–11 | 8.73E–05 |
| GO:0016787 | MF | Hydrolase activity | – | 0.012631703 |
| GO:0050660 | MF | Flavin adenine dinucleotide binding | – | 0.011249318 |
| GO:0003743 | MF | Translation initiation factor activity | – | 0.026946629 |
| GO:0006096 | BP | Glycolytic process | – | 0.001674608 |
| GO:0004842 | MF | Ubiquitin–protein transferase activity | 0.000193821 | – |

Table 2. Top 20 GO terms enriched among Zt6 treatment in upregulated DEGs and their BH-FDR adjusted *P* values.

| Category | Ontology | GO term | Zt6 24hpi | Zt6 48hpi |
|------------|----------|--|-----------|-------------|
| GO:0055114 | BP | Oxidation–reduction process | – | 4.85E–05 |
| GO:0016020 | CC | Membrane | – | 3.38E–06 |
| GO:0003824 | MF | Catalytic activity | – | 0.000329751 |
| GO:0008152 | BP | Metabolic process | – | 4.85E–07 |
| GO:0016491 | MF | Oxidoreductase activity | – | 1.29E–05 |
| GO:0005975 | BP | Carbohydrate metabolic process | – | 2.66E–05 |
| GO:0006508 | BP | Proteolysis | – | 0.003108672 |
| GO:0005737 | CC | Cytoplasm | – | 0.019343795 |
| GO:0015979 | BP | Photosynthesis | – | 2.84E–08 |
| GO:0005509 | MF | Calcium ion binding | – | 0.047329562 |
| GO:0045454 | BP | Cell redox homeostasis | – | 0.019313876 |
| GO:0009765 | BP | Photosynthesis, light harvesting | – | 8.51E–10 |
| GO:0009523 | CC | Photosystem II | – | 3.38E–06 |
| GO:0006096 | BP | Glycolytic process | – | 0.017098235 |
| GO:0004222 | MF | Metalloendopeptidase activity | – | 0.030923014 |
| GO:0009654 | CC | Photosystem II oxygen evolving complex | – | 2.38E–07 |
| GO:0019898 | CC | Extrinsic component of membrane | – | 3.38E–06 |
| GO:0004427 | MF | Inorganic diphosphatase activity | – | 0.000591418 |
| GO:0009538 | CC | Photosystem I reaction center | – | 0.00010569 |
| GO:0042132 | MF | Fructose 1,6-bisphosphate 1-phosphatase activity | – | 0.008522977 |

Table 3. Top 20 GO terms enriched among Zt6 treatment in downregulated DEGs and their BH-FDR adjusted *P* values.

terms included five (of a total six) photosynthesis-related GO terms. Of those remaining, only five were shared between the Zt6 treatment and at least one SSP treatment. Furthermore, whilst four photosynthesis related GO terms were also significantly enriched among upregulated DEGs, three were exclusive to the Zt11 treatment at 24hpi, and one was significantly enriched in both Zt11 and Zt12 treatments, also only at 24hpi.

The response to Zt6 expression was characterised by an overrepresentation of up-regulated genes involved in protein phosphorylation and kinase activity (Table 2). This included the GO terms transferase activity (of

| Category | Ontology | GO term | Zt9 24hpi | Zt11 24hpi | Zt12 24hpi | Zt9 48hpi | Zt11 48hpi | Zt12 48hpi |
|------------|----------|--|-----------|------------|------------|-----------|------------|-------------|
| GO:0000166 | MF | Nucleotide binding | – | – | – | – | 0.007614 | – |
| GO:0016020 | CC | Membrane | – | – | 0.019551 | – | – | – |
| GO:0003723 | MF | RNA binding | – | – | – | 0.01373 | 0.001801 | 0.003530677 |
| GO:0005737 | CC | Cytoplasm | – | – | – | 4.80E–05 | 0.005605 | 3.39E–08 |
| GO:0006886 | BP | Intracellular protein transport | – | – | – | 0.025869 | – | 0.009031875 |
| GO:0005525 | MF | GTP binding | – | – | – | 0.01373 | 0.033079 | – |
| GO:0006457 | BP | Protein folding | – | – | – | 0.000534 | 0.021678 | – |
| GO:0015031 | BP | Protein transport | – | – | – | 0.001189 | 0.020858 | 0.009555824 |
| GO:0003924 | MF | GTPase activity | – | – | 0.017039 | 0.002129 | – | 0.012699485 |
| GO:0005215 | MF | Transporter activity | – | – | 0.049319 | – | – | – |
| GO:0006511 | BP | Ubiquitin-dependent protein catabolic process | – | – | – | 0.040493 | 0.033365 | – |
| GO:0006184 | BP | Obsolete GTP catabolic process | – | – | 0.001484 | 0.030019 | – | – |
| GO:0015035 | MF | Protein disulfide oxidoreductase activity | – | – | 4.77E–07 | – | – | – |
| GO:0004298 | MF | Threonine-type endopeptidase activity | – | – | – | 3.81E–11 | 1.62E–07 | 4.28E–05 |
| GO:0005839 | CC | Proteasome core complex | – | – | – | 3.81E–11 | 1.62E–07 | 4.28E–05 |
| GO:0051603 | BP | Proteolysis involved in cellular protein catabolic process | – | – | 3.81E–11 | 1.62E–07 | 4.28E–05 | – |
| GO:0051536 | MF | Iron-sulfur cluster binding | – | – | – | – | – | 0.025876055 |
| GO:0006913 | BP | Nucleocytoplasmic transport | – | – | – | 0.044736 | – | – |
| GO:0004175 | MF | Endopeptidase activity | – | – | – | 7.37E–11 | 5.20E–06 | 5.71E–05 |
| GO:0015935 | CC | Small ribosomal subunit | – | – | – | – | – | 0.000673305 |

Table 4. Top 20 GO terms enriched among SSP treatments in upregulated DEGs and their BH-FDR adjusted *P* values.

phosphorous-containing groups), protein phosphorylation, protein kinase activity, and protein serine/threonine kinase activity. By contrast, these signaling-related GO terms were enriched amongst downregulated genes for the SSP treatments (Table 5).

GO categories overrepresented among upregulated genes in the SSP treatments were those related to cellular protein catabolic processes. This included threonine type endopeptidase activity, endopeptidase activity, proteasome complex, and proteolysis involved in cellular catabolic process, as well as categories involved in other cellular protein metabolic processes such as protein folding (Table 4).

Over-represented among genes downregulated in the SSP treatments were those involved in diverse processes such as response to hormone and signal transduction, binding and activity of microtubules, lipid and fatty acid metabolism, DNA replication, and protein complex genes (e.g. MCM complex and kinesin complex) (Table 5). Closer investigation showed that downregulation of histone protein genes was the sole driver of enrichment of the GO term nucleosome. GTPase activity and GTP-catabolic process were the only GO terms enriched among both up and downregulated genes of the SSP treatments.

Differential expression of immune receptor-like genes. We previously demonstrated the requirement for the cell-surface co-receptors NbBAK1 and NbSOBIR1 for full induction of cell death by the SSP effector group¹⁹. These co-receptors are not required for Zt6-induced cell death²⁰. It is therefore likely that cell death induced by the SSP treatments is a form of immune receptor-mediated programmed cell death. In contrast, Zt6-induced cell death is likely to be more similar to necrosis. Expression of receptors is often upregulated in response to the presence of their ligand^{23–25}. We therefore assessed whether there were differential expression patterns of cell surface and cytoplasmic receptor gene families commonly associated with pathogen-associated molecular pattern (PAMP) and/ or effector recognition. Lists of these gene family members used in this assessment were obtained by filtering of the *N. benthamiana* genome annotation (GFF3) file according to their description in the note field (Supplementary material 1).

There were clear differences in transcriptional profiles across receptor families induced by Zt6 and SSP treatments. At 24hpi, transcriptional changes among wall-associated kinases (WAKs), receptor like kinases (RLKs), leucine-rich repeat receptor like-kinases (LRR-RLKs), and lectin-receptor kinases (LecRLKs) showed a clear bias toward upregulation in the Zt6 treatment (Fig. 3). For WAKs and RLKs this bias was reversed in the

| Category | Ontology | GO Term | Zt9 24hpi | Zt11 24hpi | Zt12 24hpi | Zt9 48hpi | Zt11 48hpi | Zt12 48hpi |
|------------|----------|---|-----------|------------|------------|-----------|------------|------------|
| GO:0005524 | MF | ATP binding | - | - | - | 0.003562 | 0.000859 | 0.001602 |
| GO:0016772 | MF | Transferase activity, transferring phosphorus-containing groups | - | - | - | 0.00188 | 0.001457 | 0.000473 |
| GO:0006468 | BP | Protein phosphorylation | - | - | - | 0.00051 | 2.08E-05 | 0.000285 |
| GO:0004672 | MF | Protein kinase activity | - | - | - | 0.000472 | 2.05E-05 | 0.000285 |
| GO:0004674 | MF | Protein serine/threonine kinase activity | - | - | - | 0.046399 | 0.008818 | - |
| GO:0005622 | CC | Intracellular | - | - | - | 0.039391 | - | - |
| GO:0004553 | MF | Hydrolase activity, hydrolyzing O-glycosyl compounds | - | - | - | 0.016349 | 0.031375 | 0.000686 |
| GO:0003676 | MF | Nucleic acid binding | - | - | 0.013303 | - | - | - |
| GO:0006629 | BP | Lipid metabolic process | - | - | - | 0.014781 | 0.00668 | - |
| GO:0007165 | BP | Signal transduction | - | - | - | 0.04235 | - | - |
| GO:0008017 | MF | Microtubule binding | - | - | - | 0.025374 | 0.006388 | 3.70E-06 |
| GO:0003777 | MF | Microtubule motor activity | - | - | - | - | - | 0.002007 |
| GO:0007018 | BP | Microtubule-based movement | - | - | - | - | - | 0.002007 |
| GO:0003924 | MF | GTPase activity | - | - | - | 0.025071 | - | 0.043564 |
| GO:0005871 | CC | Kinesin complex | - | - | - | - | 0.042549 | 0.001638 |
| GO:0006184 | BP | Obsolete GTP catabolic process | - | - | - | 0.008125 | 0.016356 | 0.001431 |
| GO:0046982 | MF | Protein heterodimerization activity | - | - | - | 2.45E-07 | 7.58E-09 | 0.000252 |
| GO:0030246 | MF | Carbohydrate binding | - | - | - | 0.025642 | - | - |
| GO:0000786 | CC | Nucleosome | - | - | - | 5.28E-11 | 6.23E-08 | 8.46E-05 |
| GO:0006633 | BP | Fatty acid biosynthetic process | - | - | - | - | - | 0.042652 |

Table 5. Top 20 GO terms enriched among SSP treatments in downregulated DEGs and their BH-FDR adjusted *P*-values.

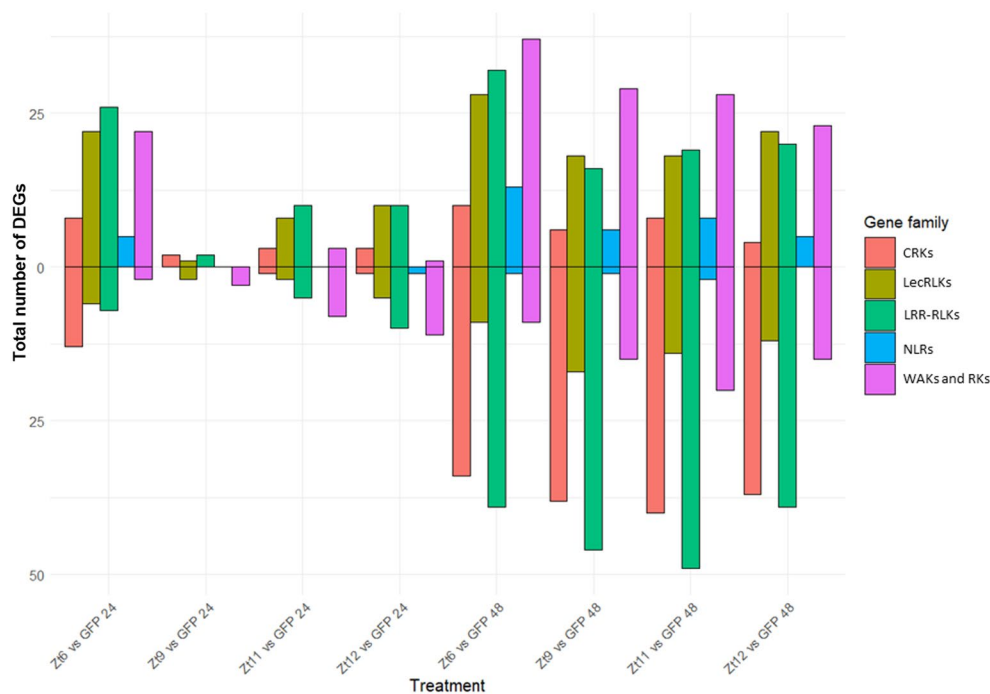
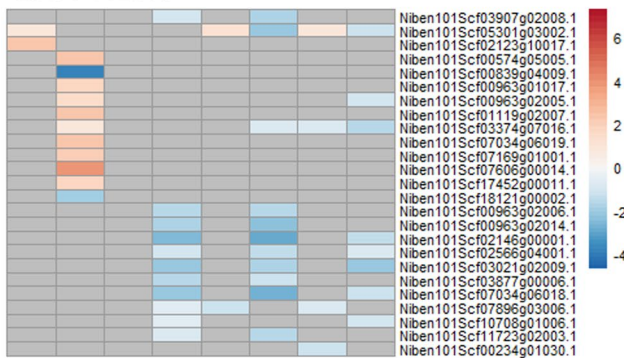
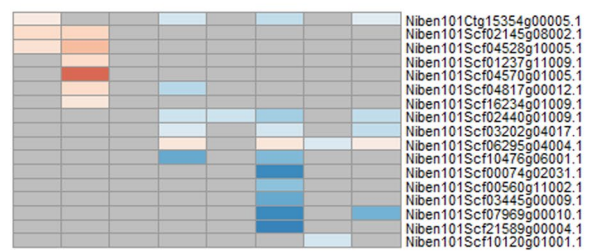


Figure 3. Total numbers of DEGs from each of five pathogen response associated gene families in each treatment. CRKs, LecRLKs, LRR-RLKs, NLRs, WAKs and RKs. Number of upregulated DEGs are represented by bar size above the x-axis, number of downregulated DEGs are represented by bar size below the x-axis.

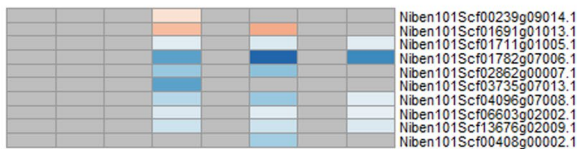
LRR-RLKs



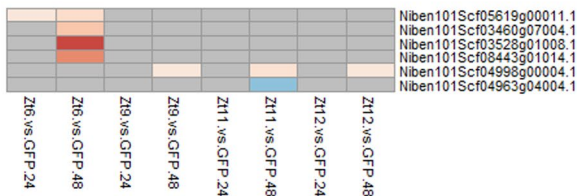
WAKs and Receptor Kinases



CRKs



NBS-LRRs



LecRLKs

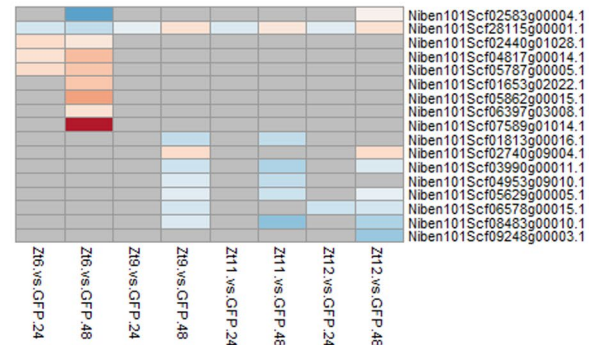


Figure 4. Expression profile of defence associated genes with marked difference in expression change between Zt6 and Zt9, Zt11, and Zt12.

SSP treatments, and for LRR-RLKs there was a smaller number of genes upregulated in the SSP treatments in comparison to Zt6. For LecRLKs, a similar bias towards upregulation existed between the Zt6, Zt11, and Zt12 treatments, although this was less prominent for Zt11 and Zt12. At 48hpi, LRR-RLKs in the SSP treatments showed a bias towards downregulation, this was less prominent in the Zt6 treatment. Interestingly, at 48hpi cysteine-rich receptor-like kinases (CRKs) showed a strong bias toward downregulation across all treatments. Genes annotated as nucleotide-binding site leucine-rich repeats (NLRs) showed bias toward upregulation at 48hpi across all treatments, though this was a comparatively small number of genes relative to the total *N. benthamiana* NLR gene complement.

Given the differences between the transcriptomes of the Zt6 and SSP treatments, we next identified and investigated specific genes within these five receptor gene families, based on whether they showed a marked difference in expression change between the Zt6 and SSP treatments. For the purposes of this investigation, we defined a “marked difference” as a \log_2 fold change of an absolute value of at least 1.0 in one or all of the SSP treatments which was either not present or reversed in the Zt6 treatment, or a \log_2 fold change of an absolute value of at least 1.0 in the Zt6 treatment which was either not present or reversed across all the SSP treatments. For all six receptor gene families, few expression changes were apparent at 24hpi in comparison to the GFP control. However, at 48hpi a clear difference in expression pattern is visible between the Zt6 and SSP treatments. Zt6 treatment specifically induced expression of 11 out of a total 149 LRR-RLKs, 4 out of a total 65 NLRs, 7 out of a total 107 WAKs/RLKs and 7 out of a total 54 LecRLKs in the current *N. benthamiana* genome annotation (Fig. 4). Zt6 also induced expression of 34 out of a total 795 RLPKs in the current *N. benthamiana* genome annotation (Fig. 5). Only two LRR-RLKs, two LecRLKs and 7 RLPKs were downregulated in response to Zt6 treatment (Fig. 4). Interestingly, no CRKs showed any transcriptional response to Zt6 (Fig. 4). Very few receptor genes were induced by the SSP treatments, although notable exceptions included one NLR gene induced by all three SSP treatments at 48hpi (Fig. 4), and two RLPK genes relatively strongly induced only by the Zt12 treatment at 48hpi (Fig. 5). Most receptor family genes that were transcriptionally responsive to the SSP treatments were downregulated. These included a notable over-abundance of WAKs and RLKs that were strongly downregulated in response only to the Zt11 treatment at 48hpi. Together, this suggests that there is reprogramming of receptor gene expression following exposure to SSPs, and that this differs between individual SSPs.

Differential expression of transcription factors, senescence- and programmed cell death-associated genes. There are differences in how developmental, pathogen-associated, and stress-induced PCD is executed. These distinct but partially overlapping pathways share some common components, although no

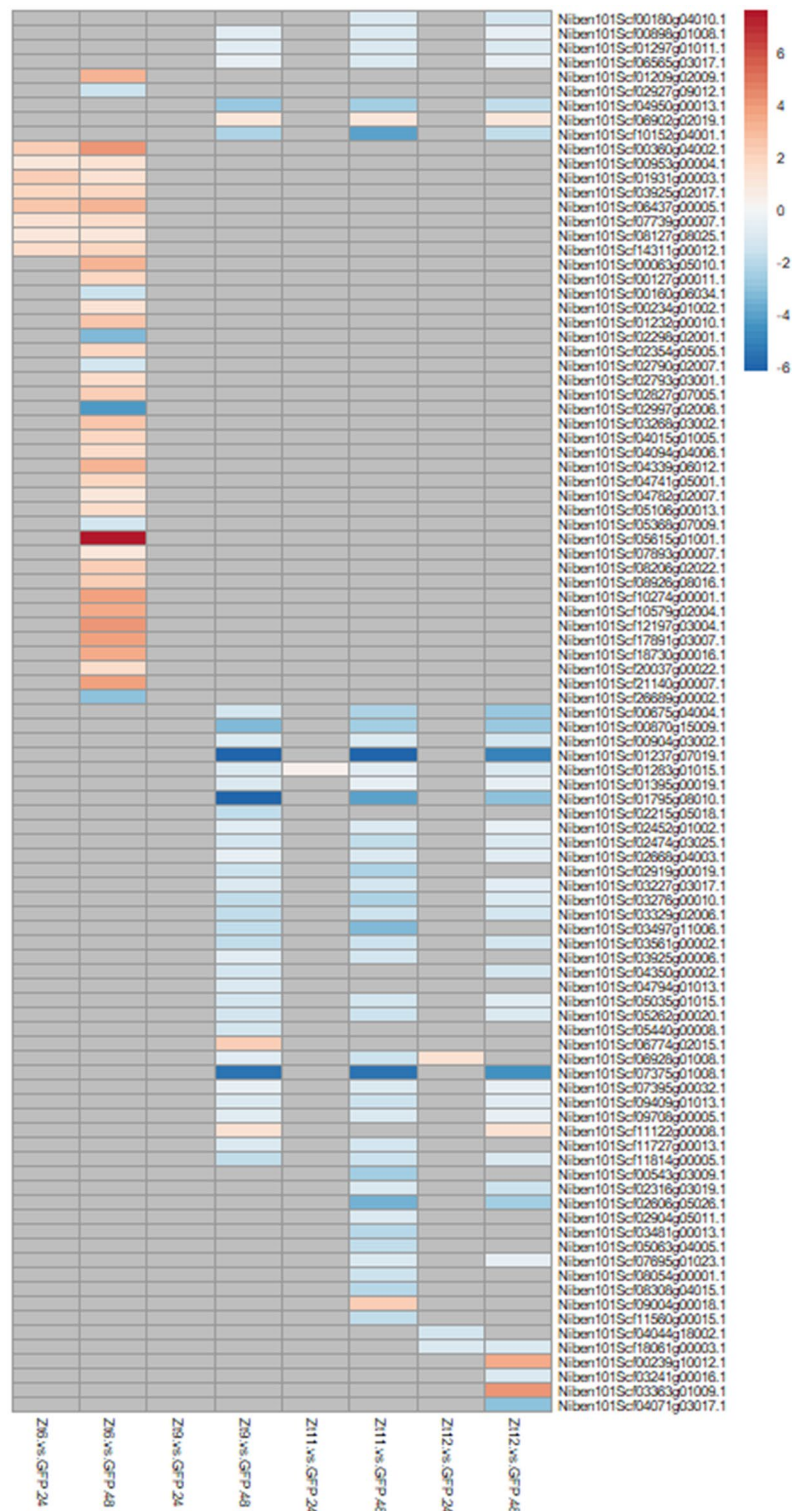
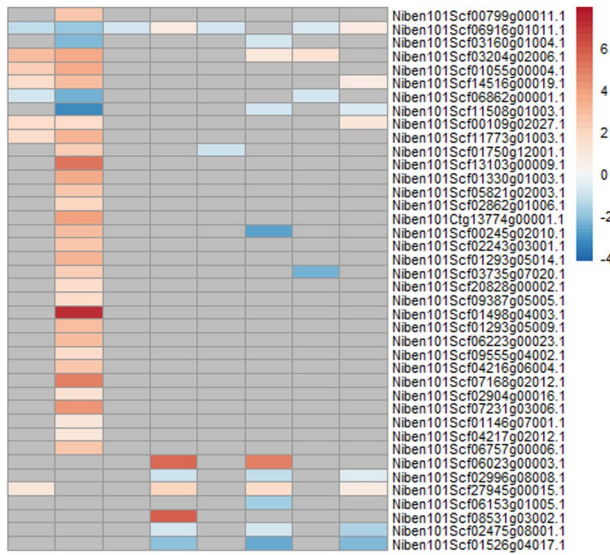


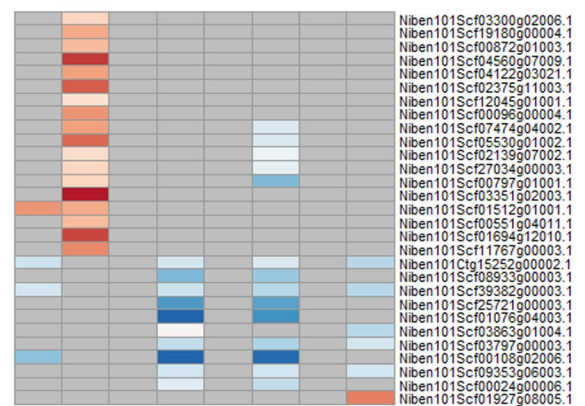
Figure 5. Expression profile of RLPKs genes with marked difference in expression change between Zt6 and Zt9, Zt11, and Zt12.

pathway is fully characterised^{26–28}. Our GO enrichment analysis showed that Zt6 and SSP treatments appear to be inducing different types of transcriptional reprogramming, and that Zt6-induced cell death is distinct from ordered receptor-mediated PCD²⁶. We therefore investigated whether gene families commonly associated with transcriptional reprogramming related to senescence and PCD displayed differential expression patterns between the Zt6 and SSP effector treatments. We first evaluated the expression of the NAC, WRKY, TCP and MYB transcription factor (TF) families (Fig. 6), which are widely reported as regulators of leaf senescence^{29–32}.

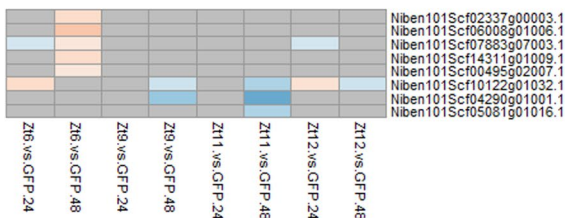
NAC transcription factors



MYB transcription factors



TCP transcription factors



WRKY transcription factors

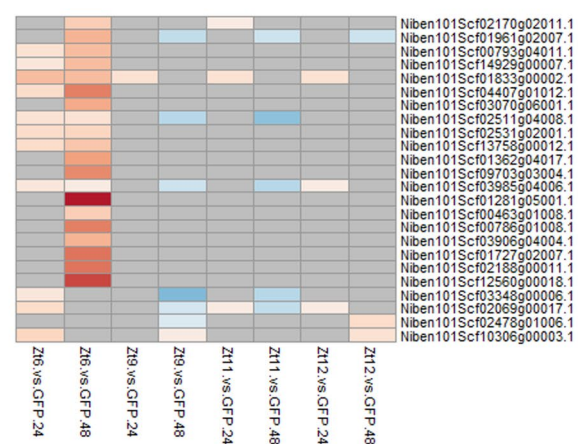


Figure 6. Expression profile of senescence regulating transcription factor genes with marked difference in expression change between Zt6 and Zt9, Zt11, and Zt12.

At 24hpi there were few differential expression changes across any of the treatments. However, by 48hpi there were clear differences in expression patterns between Zt6 and SSP treatments. Overall, Zt6 treatment induced expression of 29 out of a total 241 NACs, 20 out of a total 151 WRKYs, 5 out of a total 61 TCPs and 18 out of a total 242 MYBs in the current *N. benthamiana* genome annotation. Only four TFs (all NAC family) were downregulated in response to Zt6 at this timepoint. In contrast, few TFs of any family were induced by the SSP treatments. Indeed, for WRKY, MYB and TCP families the majority of differentially expressed TFs were downregulated in response to effector treatment. There was minimal overlap between the individual genes transcriptionally induced by Zt6, and those transcriptionally repressed by the SSP treatments.

Several other non-TF gene families have previously been linked to senescence and cell death in plants. Metacaspases, zinc-finger domain containing proteins and HR-inducing proteins have all been shown to regulate PCD or HR^{33,34}. The senescence-associated genes *SAG1* (*Arabidopsis thaliana*), *SAG102* (*Medicago truncatula*), *phytoalexin-deficient 4* (*PAD4/SAG101*) and *harpin-induced gene 1* (*HIN1*), are markers of senescence and HR respectively^{35–37}. In addition, *STAY-GREEN* (*SGR*) genes operate downstream of NAC TF regulation of senescence in the catabolism of chlorophyll³⁸. We investigated expression of these genes across all treatments and found that expression of genes encoding metacaspases, SAG, HR-inducing and SGR proteins between Zt6 and SSP treatments was similar (Fig. S1). However, a zinc-finger containing protein encoding gene was strongly induced by Zt6 treatment at 24- and 48hpi but not by the SSP treatments.

Discussion

In this investigation, we aimed to understand the mechanisms of cell death induced by several SSPs of *Z. tritici* in a non-host plant *N. benthamiana*. To do this, we made use of the Agrobacterium-mediated transient expression system originally used to identify the cell death-inducing activity of these proteins. Our results indicate that expression of this group of proteins initiate massive transcriptional reprogramming of plant cells prior to onset of macroscopic cell death. Furthermore, the cytotoxic secreted ribonuclease Zt6 induces a transcriptional response distinct from other SSPs.

The transcriptional responses to SSP treatments were broadly similar, and clearly distinct from that induced by Zt6 expression. PCA (Fig. 1A) illustrates clustering of SSP treatments at 24hpi and obvious separation from both Zt6 and GFP control treatments. This separation is further exaggerated by 48hpi. The total number of genes that are differentially expressed is greater in Zt6 treated leaves at 24hpi in comparison to SSP treatments. This indicates that transcriptional reprogramming begins earlier for Zt6 compared to SSP treatments. This is consistent with the earlier onset of macroscopic cell death induced by Zt6 in comparison with the SSPs²⁰. This may reflect that Zt6 induces cell death due to its enzymatic RNase activity targeting rRNA²⁰, whereas the SSPs presumably induce immune receptor-mediated cell death¹⁹.

Given the dependency of SSP induced cell death on BAK1, broad similarity of transcriptional response to all three SSP treatments is not surprising. BAK1 functions as a co-receptor for various RLKs, including some whose ligands are PAMPs or secreted effectors; it is a convergence point of multiple pathogen-triggered physiological pathways that lead to PCD³⁹. However, BAK1 dependent PCD has been shown to be highly ligand specific and could still proceed via a variety of mechanisms, dependent upon how its PAMP/effector co-receptor role disrupts its normal functioning and modifies how it subsequently interacts with other RLKs (including but not limited to BIR1 and SOBIR1), which also have important roles in regulation of PCD³⁹.

A notable difference in response to the Zt6 and SSP treatments was differential regulation of genes involved in microtubule activity, movement and binding. Downregulation of genes in these categories was enriched in SSP treatments at 48hpi (Table 5), but not in Zt6 at either timepoint (Table 3). Depolymerisation of the microtubule network has been associated with PCD⁴⁰, in particular in HR reactions in *A. thaliana*⁴¹ and soybean⁴². Microtubule reorganisation is also associated with developmental PCD processes such as self-incompatibility⁴³. That genes facilitating maintenance of a normal microtubule network are downregulated in SSP treatments is a characteristic of an orderly form of PCD. The absence of this downregulation during Zt6 expression is consistent with a form of cell death relying less on cytoskeleton arrangement. In contrast, GO term analysis revealed enrichment of terms associated with ribosome, structural constituent of the ribosome, translation and translation initiation factor activity in genes upregulated by Zt6 at 48hpi (Table 2). This suggests Zt6 treatment induced a significant stress on ribosome function and on protein translation in general. These terms are not enriched in either up- or downregulated groups for the SSP treatments at either timepoint (Tables 4, 5). An upregulation of genes related to ribosome structure and function could be indicative of cells experiencing ribosomal stress and therefore perturbations in protein synthesis. This would be anticipated in cells expressing Zt6 which has previously been shown to cleave plant rRNA in a semi-specific manner²⁰. These expression signatures are a likely response to compensate for reduced ribosome functionality in cells expressing Zt6.

The GO terms protein kinase activity and protein serine/threonine kinase activity were strongly induced by Zt6 treatment at the 24hpi timepoint (Table 2). In contrast, these groups were unchanged at 24hpi and subsequently downregulated at 48hpi in the SSP treatment group (Table 5). The activity of several serine/threonine kinase proteins is important for control of apoptosis and autophagy in animal systems⁴⁴. However, the role of this protein class in direct activation of plant cell death is less clear. Many transmembrane receptor kinases and intracellular kinases play important roles in ligand perception and signal transduction during pathogen interaction. We were specifically interested in expression patterns of genes belonging to the RLP, RLK and WAK receptor families. These classes of receptors are well known to be involved in recognition of PAMPs or apoplastic effectors. Indeed, the only two cloned R genes against *Z. tritici* are *Stb6* and *Stb16q*, encoding a WAK and a CRK respectively^{24,45}. Expression of immune receptors is often upregulated in response to pathogens^{23–25}. Here, we found a number of genes annotated as receptor-like kinases, WAKs, LRR-RLKs, CRKs and LecRLKs were transcriptionally responsive to SSP expression (Fig. 4). Expression patterns were similar between SSP effector treatments in comparison to Zt6. Most differentially expressed receptors in these classes were downregulated upon SSP expression, but upregulated in response to Zt6. A small number of genes annotated as NLRs were induced by Zt6 expression only. A larger number of genes annotated as RLPKs were differentially expressed upon SSP treatment. Again, there is clear differentiation between expression patterns induced by SSPs in comparison to Zt6. Nearly all RLPKs differentially expressed in the Zt6 group were upregulated, whereas nearly all RLPKs that changed in response to SSPs were downregulated. Our data therefore provides a small number of candidate cell surface immune receptors for recognition of *Z. tritici* SSPs for validation in a follow-on study.

The GO terms proteasome core complex and proteolysis involved in cellular protein catabolic process were enriched amongst upregulated genes for all three of the SSP group treatments at 48hpi. In contrast, proteolysis was enriched amongst downregulated genes at 48hpi for the Zt6 treatment. The contribution of proteasome function to PCD has previously been investigated. Silencing of components of the 26S proteasome leads to build-up of polyubiquitinated proteins and induction of PCD⁴⁶, suggesting proteasome function negatively regulates PCD. In contrast, Hatsugai and colleagues identified a mechanism involving PBA1 linking proteasome function with promotion of PCD⁴⁷. These contrasting results suggest that role of the proteasome in cell death may be highly complex.

Given the differences in expression profile of many receptor gene groups between Zt6 and SSP treatments, we focussed on TF family expression patterns. Overall, Zt6 expression led to induction of many NAC, MYB and WRKY TFs (Fig. 6). In contrast, the majority of these genes were either unresponsive or repressed by

SSP expression. This pattern was also true for TCP TFs, although the number of genes that were differentially expressed were much lower. NAC TFs play important roles in senescence and in both biotic and abiotic stress responses. Expression of the NAC transcription factor gene *Niben101Scf01498g04003* was striking as it was strongly upregulated in the Zt6 48 h treatment exclusively (Fig. 6A). This gene shows a high level of homology to *A. thaliana* *ANAC032* (E -value = $8e-08$). *ANAC032* regulates senescence through modulation of *AtNYE1*, the so-called *STAY-GREEN* gene involved in ability to catabolize chlorophyll³⁸. In *N. benthamiana* there are six *STAY-GREEN* genes, however, only two were responsive to treatment and this pattern was similar across Zt6 and SSP treatments. This suggests that while transcriptional promotion of senescence at the level of TF genes appears starkly different between Zt6 and SSP treatments, there may yet be some downstream regulatory convergence.

Other pathways by which upregulation of *Niben101Scf01498g04003* may promote senescence are alluded to by its other closest homologs in *A. thaliana*. The second of these was *ANAC047* (AKA. *SPEEDY HYPONASTIC GROWTH*) (E -value = $1e-07$). *ANAC047* is upregulated during and may promote leaf senescence via regulation of *ACO5* (1-aminocyclopropane-1-carboxylic acid oxidase 5), an enzyme involved in ethylene biosynthesis^{48,49}. In our results, a number of genes annotated as *ACO5* and *ACO5* homologs exhibit both up and downregulation in the Zt6 treatments. Alternatively, the 10th closest homolog of *Niben101Scf01498g04003* is *ANAC082* (E -value = $2e-05$) which plays a role in the sensing of nucleolar stress⁵⁰. This is due to the presence of an upstream open reading frame (uORF) in *ANAC082* mRNA. Previously studied uORFs act as negative regulators of the main ORF due to ribosome stalling on the mRNA. Upregulation of *ANAC082* expression could therefore be indicative of ribosome instability induced by Zt6 expression. Given the known interaction between Zt6 and rRNA, this could provide a mechanism for Zt6-specific patterns of transcriptional reprogramming and a role for this TF as a master regulator of downstream gene expression.

A number of MYB TF genes were also highly upregulated by Zt6 but not by other treatments (Fig. 6B). Of these, *Niben101Scf04560g07009* and *Niben101Scf01694g12010* were most strongly induced. These genes are orthologous to *AtMYB36/MYB68* and *AtMYB119* respectively. These TFs are known to have roles in root cell differentiation⁵¹, root development⁵² and cellular differentiation during female gametogenesis⁵³. However, these TFs are not documented to be involved in induction of cell death or pathogen responses. Similarly, the *N. benthamiana* ortholog of *AtTCP5* was induced by Zt6 treatment alone (Fig. 6C). This TF has roles in floral development and ethylene biosynthesis⁵⁴ but not known to be involved in cell death pathways.

WRKY transcription factors have been identified as important regulators of biotic stress responses⁵⁵. It is possible that increased WRKY activity accounts for the modified expression of defence genes such as RLPs and RLKs. Therefore, the complex interplay between these TF families might explain the transcriptional profile induced by the effector treatments. A number of WRKYs were transcriptionally induced by Zt6 but not by SSP expression (Fig. 6D). Of these, *Niben101Scf01281g05001* and *Niben101Scf12560g00018* were the most strongly upregulated. These TFs are orthologues to *AtWRKY14/35* and *WRKY22* respectively. Constitutively activated *WRKY14* is known to promote cell death in *N. benthamiana*⁵⁶. *WRKY22* has previously been implicated in plant defence responses and loss of *WRKY22* expression compromises effector-induced cell death in *N. benthamiana*⁵⁷. This suggests that WRKY-dependent transcriptional reprogramming may contribute to cell death induced by Zt6.

Taken together our results show that *Z. tritici* secreted ribonuclease Zt6 and three SSPs, trigger cell death in non-host *N. benthamiana* at least partially via gene expression changes in clearly distinct cohorts of genes. Within these cohorts, only a small number of genes that could function as immune receptors were upregulated in response to SSP expression, and therefore provide a manageable set of candidates for further study as potential *Z. tritici* non-host R-genes. Among TF genes in these cohorts, those transcriptionally responsive to Zt6 expression suggest a potential pathway to ribosomal stress induced PCD. This work provides a detailed picture of transcriptional changes that occur in *N. benthamiana* prior to cell death induced by apoplastic recognition of non-host pathogen SSPs, and ribonuclease activity.

Methods

Plants and bacterial strains. All *N. benthamiana* plants were from a seed stock used in our previous investigation under the same growth conditions^{19,20}. The *Agrobacterium tumefaciens* GV3101 (pMP90) strains expressing Zt6, Zt9, Zt11 and Zt12 from pEAQ-HT-DEST3 were described previously^{19,20}.

Generation of RNA samples. Leaves of 5-week old *N. benthamiana* seedlings were syringe infiltrated with *Agrobacterium* suspensions at OD₆₀₀ = 1.2 in Agroinfiltration buffer (10 mM MgCl₂, 10 mM MES, 150 μM acetosyringone, pH 5.6). Six plants were infiltrated per treatment with 30 plants used in total. Leaf sampling was performed at 24 h and 48 h post infiltration. Three infiltrated leaf patches, one each from three individual plants were cut from leaves and pooled to produce each sample. Samples were snap frozen in liquid nitrogen and stored at -80 °C until processing.

RNA extraction. Frozen leaf samples were ground in liquid nitrogen using a mortar and pestle. RNA was extracted using a Trizol/Chloroform procedure as described previously⁵⁸. DNase digest was performed using RQ1 DNase (Promega) and RNA recovered by ethanol precipitation. RNA quality and purity was measured using Qubit and Nanodrop. RNA-seq was performed by BGI on the Illumina HiSeq2500 platform.

Quality control and alignment. Quality of raw reads was manually assessed using FastQC software v0.119⁵⁹. Filtering of raw reads was then performed using PRINSEQ-lite software v0.20.4 to a minimum Phred-quality score of 26⁶⁰. Version 1.0.1 of the *N. benthamiana* reference genome was downloaded from the Sol Genomics Network ftp site (ftp://ftp.solgenomics.net/genomes/Nicotiana_benthamiana). Index of the reference genome was built using the build function in HISAT2 v2.1.0, filtered paired-end reads were aligned to

the reference genome using the same software⁶¹. Following alignment, resulting .BAM files for each treatment were checked for uniformity of gene body coverage and sufficient reads per kilo base per million mapped reads (RPKM) saturation using RSeQC v2.6.4⁶².

Read counting and differential expression analysis. Reads mapped to each gene were counted using HTSeq v0.11.0. Differential expression analysis was conducted between each treatment and the control treatment (GFP) at the appropriate time point using the DESeq2 R package v1.32.0, with a Benjamini-Hochberg – false discovery rate (BH-FDR) corrected *P*-value of 0.05²².

Gene ontology enrichment analysis. Significantly differentially expressed genes (DEGs) were divided into those that were up-regulated and those that were down-regulated at each time point in each treatment. Gene ontology enrichment analysis was conducted on each of these two groups of DEGs using the Goseq R package with BH-FDR *P*-value adjustment and gene length bias correction, GO terms with adjusted *P*-values less than 0.05 were considered to be significantly enriched in that group²².

Ethics approval. All handling of plants, microorganisms and associated samples in this study was performed under biosafety regulations in place at Rothamsted Research. All work was carried out under the Department for Environment, Food and Rural Affairs (Defra) plant health licenses Nos. 101941/197343/8 and 101948/198285/4.

Data availability

All RNA-seq raw sequencing data used in this study were deposited into the NCBI SRA under BioProject accession number PRJNA858969.

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Author contributions

Agroinfiltrations were performed by G.K. and RNA extracted and purified by C.B. Computational analysis of RNAseq data was done by T.W., G.K., J.R. and K.K. conceived the project and T.W., G.K., J.R. and K.K. wrote the manuscript. This work was funded by Rothamsted Research Fellowship Grant (to J.R. and K.K.) and a JABBS Foundation startup package to G.K.

Competing interests

The authors declare no competing interests.

Additional information

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