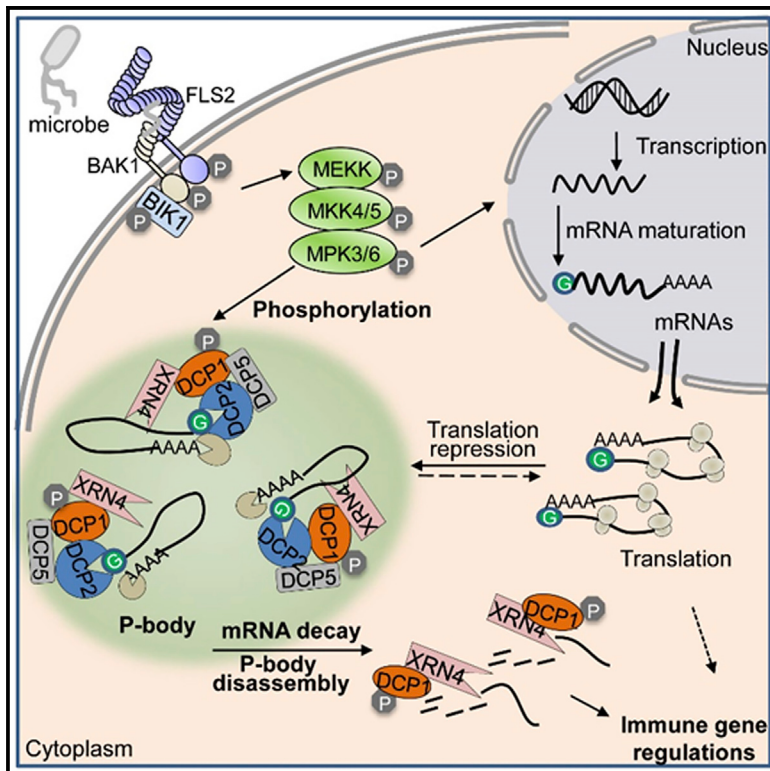


Orchestration of Processing Body Dynamics and mRNA Decay in *Arabidopsis* Immunity

Graphical Abstract



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In Brief

Yu et al. show that P-body core components DCP1 and DCP2 positively regulate plant pattern-triggered immunity. DCP1 phosphorylation by immune-activated MAPKs contributes to P-body disassembly and mRNA decay on a subset of immune-regulated genes, revealing mRNA-decay-mediated posttranscriptional regulation is an integral part of plant immunity.

Highlights

- P-body dynamics is regulated in plant pattern-triggered immunity (PTI)
- P-body components DCP1 and DCP2 positively regulate plant PTI
- Microbe-associated molecular pattern-activated MAP kinases phosphorylate DCP1
- DCP1 phosphorylation contributes to mRNA decay of certain immune-related genes



Orchestration of Processing Body Dynamics and mRNA Decay in *Arabidopsis* Immunity

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SUMMARY

Proper transcriptome reprogramming is critical for hosts to launch an effective defense response upon pathogen attack. How immune-related genes are regulated at the posttranscriptional level remains elusive. We demonstrate here that P-bodies, the non-membranous cytoplasmic ribonucleoprotein foci related to 5'-to-3' mRNA decay, are dynamically modulated in plant immunity triggered by microbe-associated molecular patterns (MAMPs). The DCP1-DCP2 mRNA decapping complex, a hallmark of P-bodies, positively regulates plant MAMP-triggered responses and immunity against pathogenic bacteria. MAMP-activated MAP kinases directly phosphorylate DCP1 at the serine²³⁷ residue, which further stimulates its interaction with XRN4, an exonuclease executing 5'-to-3' degradation of decapped mRNA. Consequently, MAMP treatment potentiates DCP1-dependent mRNA decay on a specific group of MAMP-downregulated genes. Thus, the conserved 5'-to-3' mRNA decay elicited by the MAMP-activated MAP kinase cascade is an integral part of plant immunity. This mechanism ensures a rapid posttranscriptional downregulation of certain immune-related genes that may otherwise negatively impact immunity.

INTRODUCTION

Detecting the presence of microbial components is crucial for hosts to launch an effective defense response against pathogen invasion. Plant pattern-triggered immunity (PTI) is initiated by the recognition of microbe-associated molecular patterns (MAMPs) by cell-surface-resident pattern recognition receptors (PRRs)

(Couto and Zipfel, 2016; Yu et al., 2017). Flagellin-sensing 2 (FLS2) and elongation factor EF-TU receptor (EFR), two PRRs from *Arabidopsis*, recognize flg22 and elf18, the synthetic peptides derived from bacterial MAMP flagellin and EF-Tu, respectively (Gómez-Gómez and Boller, 2000; Zipfel et al., 2006). Both FLS2 and EFR are leucine-rich repeat-receptor-like kinases (LRR-RLKs) and heterodimerize with the shared co-receptor brassinosteroid insensitive 1-associated kinase 1 (BAK1) family LRR-RLKs to form an active PRR complex (Chinchilla et al., 2007; Heese et al., 2007; Sun et al., 2013). Members of the *Botrytis*-induced kinase 1 (BIK1) family receptor-like cytoplasmic kinases (RLCKs) associate with multiple PRR complexes and bifurcate the signaling to downstream components (Kadota et al., 2014; Li et al., 2014c; Lu et al., 2010; Zhang et al., 2010). The PRR complex undergoes layered positive and negative regulations (Böhm et al., 2014; Couto and Zipfel, 2016), and activates two major intracellular signaling modules, namely, mitogen-activated protein kinases (MAPKs) and calcium-dependent protein kinases (CDPKs) that play a primary role in transcriptional control of immune-related gene expression (Asai et al., 2002; Boudsocq et al., 2010).

In addition to the transcriptional regulation, the posttranscriptional control adds another crucial layer of regulation of immune gene expression (Carpenter et al., 2014; Staiger et al., 2013). As an important mechanism in the posttranscriptional regulation, mRNA decay could degrade both aberrant and normal transcripts (Decker and Parker, 2012). Degradation of mRNA is typically initiated with deadenylation, followed by exosome-mediated 3'-to-5' exonucleolytic decay. More often, the deadenylated mRNAs are decapped by the decapping enzyme decapping 2 (DCP2) along with its co-activator DCP1 and other cofactors and then degraded by 5'-to-3' exonuclease exoribonuclease (XRN) (Decker and Parker, 2012; Schoenberg and Maquat, 2012). The decapping machinery, together with associated non-translated mRNAs, are conditionally assembled into RNA Processing bodies (P-bodies). P-bodies, a type of ribonucleoprotein particle (RNP) granules, are microscopically visible and non-membranous cytoplasmic foci consisting of aggregations



of translationally repressed mRNAs bound by various catalytic and regulatory proteins and frequently non-coding RNAs (Mitchell and Parker, 2014). Increasing evidence indicates that P-bodies act as a storage compartment for translationally repressed mRNAs, including in the plant system (Brenques et al., 2005; Hubstenberger et al., 2017; Jang et al., 2019).

P-bodies have been observed in a wide range of organisms, including yeast, metazoans, and plants (Decker and Parker, 2012; Schoenberg and Maquat, 2012; Xu and Chua, 2011). Mutations in DCP1 and DCP2 in *Arabidopsis* cause growth defects, suggesting the importance of P-body core components in regulating gene expression essential for plant development (Xu et al., 2006). In addition, P-body components are suggested to be involved in abiotic stresses, such as drought, cold, and salinity (Perea-Resa et al., 2016; Xu and Chua, 2012). How P-body assembly is initiated and participates in a specific physiological response remain elusive. In this study, we observed rapid P-body disassembly and re-assembly upon MAMP elicitation, indicating the dynamic fate changes of mRNAs stored in P-bodies as part of immune responses. PRR-activated MAPKs directly phosphorylate DCP1 and stimulate DCP1 dissociation from DCP2 and association with exoribonuclease 4 (XRN4). Consistent with the P-body dynamics as an integral part of plant immunity, MAMP treatment potentiates DCP1-dependent mRNA decay on a subset of MAMP-downregulated genes. Such mRNA-decay-mediated posttranscriptional regulation likely underpins the ability of hosts to launch an expeditious and effective immune response by rapid reprogramming of primary immune genes.

RESULTS

Pathogen-Induced P-body Dynamics in *Arabidopsis*

We have deployed a series of random mutagenesis screens for components involved in plant PTI based on the transcriptional response of an immune reporter gene (Feng et al., 2015; Li et al., 2014a). A candidate gene identified from the screen encodes a protein localizing to P-bodies (study in progress). To reveal the potential link between P-bodies and plant immunity, we examined the dynamics of P-body assembly upon PTI elicitation. Interestingly, we observed a significant reduction of DCP1-GFP-labeled P-bodies at 15 and 30 min after flg22 treatment in *Arabidopsis* Col-0 protoplasts. The DCP1-GFP-labeled P-bodies re-appeared at 60 min after flg22 treatment (Figure 1A). The flg22-triggered P-body disassembly was not observed in the *fls2* mutant, indicating that this dynamics is specifically triggered by the flg22 recognition (Figure S1A). Similarly, flg22 treatment for 30 min led to the disassembly of P-bodies labeled with DCP5-GFP (Figure 1B) or XRN4-GFP (Figure 1C). Apparently, the reduction of P-bodies was not due to the reduced expression of DCP1-GFP, DCP5-GFP, or XRN4-GFP proteins (Figure 1D), suggesting that the disappearance of P-bodies was caused by the P-body disassembly. Furthermore, elf18 treatment also induced the disappearance of DCP1-GFP-, DCP5-GFP-, or XRN4-GFP-associated P-bodies (Figure S1B). We further generated transgenic plants expressing *DCP1-GFP* under the control of the 35S promoter. The number of DCP1-GFP-labeled P-bodies in transgenic plants was reduced upon flg22 treatment

within 30 to 60 min (Figure 1E). In addition, infiltration with a non-pathogenic bacterium *Pseudomonas syringae* pv. *tomato* DC3000 (*Pst*) type III secretion mutant, *hrcC*, which carries multiple MAMPs but does not secrete type III effectors, also triggered the disappearance of P-bodies in *DCP1-GFP* transgenic plants (Figure 1F). However, flg22-induced P-body disappearance was not observed with PAT1-GFP-labeled P-bodies, likely due to the enhanced accumulation of PAT1 proteins and/or differential response of different P-body species (e.g., PAT1 versus DCP1/DCP5/XRN4) to flg22 treatment (Figures S1C and S1D) (Roux et al., 2015). Together, the results indicate that the presence of P-bodies is dynamically modulated in response to MAMP perception.

DCP1 and DCP2 Positively Regulate Plant Immunity

The null mutants of *dcp1* or *dcp2* are post-embryonic lethal (Xu et al., 2006). To elucidate the functional relevance of P-body components in plant immunity, we silenced *DCP1* or *DCP2* in wild-type (WT) Col-0 plants with virus-induced gene silencing (VIGS) after germination at the 2-week-old seedling stage. Silencing of *DCP1* or *DCP2* did not cause detectable growth defects (Figures S1E and S1F), which is consistent with the hypothesis that the seedling lethality of *dcp1* and *dcp2* is due to the accumulation of certain proteins whose mRNAs are normally degraded after germination (Xu and Chua, 2012). Significantly, the flg22-induced expression of *FRK1* and *WRKY30*, two PTI marker genes, was reduced in *DCP1*- and *DCP2*-silenced plants compared to plants inoculated with a GFP vector control (Figure 2A). In addition, *DCP1*- or *DCP2*-silenced plants showed compromised flg22-induced callose deposition, a late PTI response, compared to control plants (Figure 2B). Furthermore, both *DCP1*- and *DCP2*-silenced plants were more susceptible to the infection by the virulent bacteria *Pst* (Figure 2C) and *P. syringae* pv. *maculicola* ES4326 (*Psm*) (Figure 2D) than control plants at 2 and 4 days post-inoculation (dpi). The inoculated leaves of *DCP1*- and *DCP2*-silenced plants developed more severe chlorosis than control plants at 4 dpi (Figures 2C and 2D). Thus, DCP1 and DCP2, the integral components of P-bodies, positively contribute to plant immunity.

MAMP-Activated MPK3/MPK6 Phosphorylate DCP1 on Ser²³⁷

Notably, DCP1-GFP proteins were detected as two major polypeptide bands in immunoblots with a shift from low to high molecular weight after flg22 treatment (Figure 1D). The shifted band of DCP1-hemagglutinin (HA) was observed as early as in 2 min, peaking at 15 min, and gradually decreasing at 60 min after flg22 treatment (Figure 3A). The flg22-induced mobility shift of DCP1-GFP proteins was also detected in the 35S::*DCP1-GFP* transgenic plants (Figure S2A). In addition, other MAMPs including bacterial elf18, lipopolysaccharide (LPS) and peptidoglycan (PGN), and fungal chitin also induced mobility shift of DCP1 in both transgenic plants and protoplasts (Figures 3B and S2B). Furthermore, infiltration of *Pst hrcC* in 35S::*DCP1-GFP* transgenic plants induced DCP1-GFP mobility shift (Figure 3C). Compared to individual MAMPs, *Pst hrcC* induced a long-lasting mobility shift of DCP1-GFP, which might be attributed to the

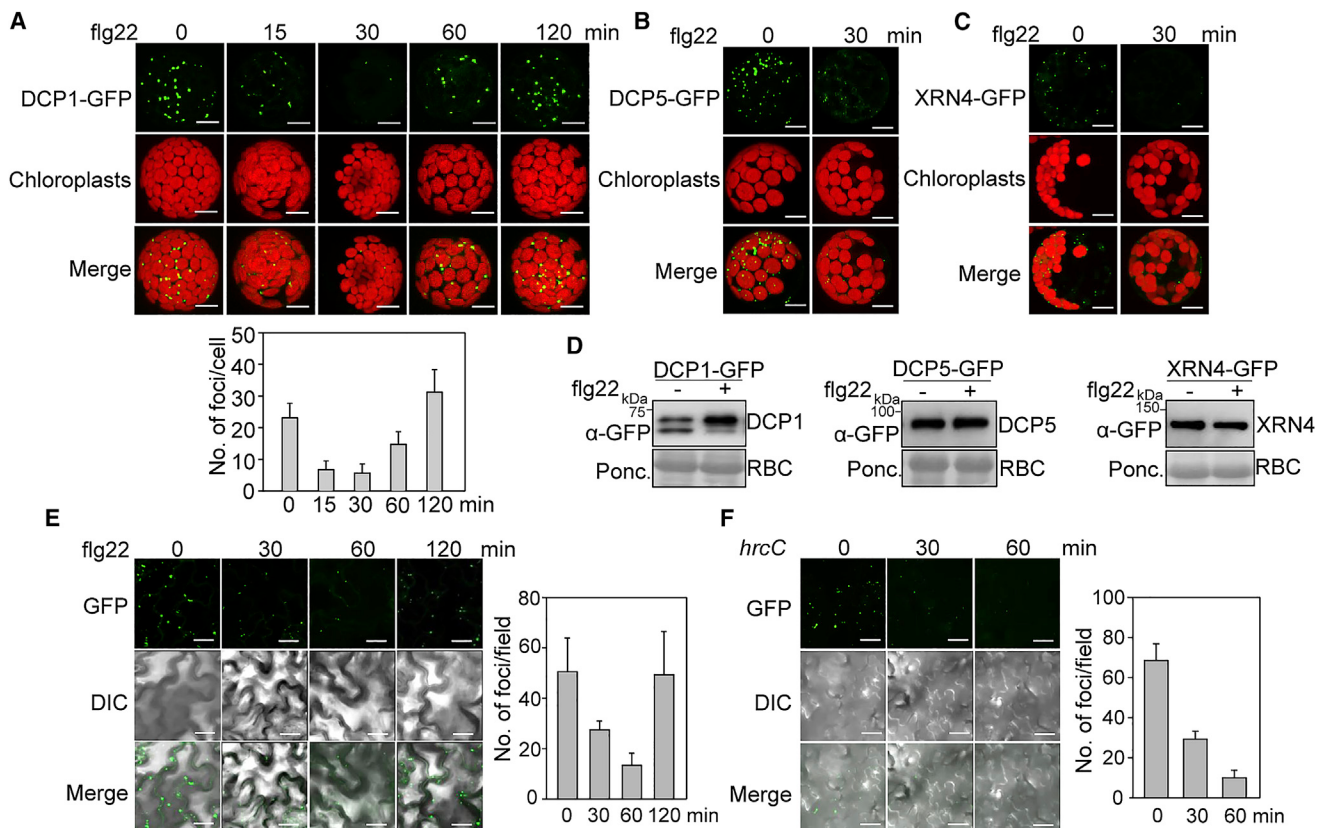


Figure 1. Pathogen-Induced P-Body Dynamics in Arabidopsis

(A) DCP1-GFP-labeled P-body dynamics upon flg22 treatment in *Arabidopsis* protoplasts. Protoplasts from wild-type (WT) Col-0 were expressed with *DCP1-GFP*, and treated with 100 nM flg22 for indicated times. Images were taken with a confocal microscope. Autofluorescence of chloroplasts is shown in the middle panels. Bar, 10 μ m. Quantification of DCP1-GFP-labeled cytoplasmic foci with ImageJ 1.49v software is shown on the bottom.

(B and C) The flg22-triggered P-body disassembly as indicated by DCP5-GFP (B)- or exoribonuclease 4 (XRN4)-GFP (C)-labeled cytoplasmic foci. Protoplasts expressing *DCP5-GFP* or *XRN4-GFP* were treated with 100 nM flg22 for 30 min. Bar, 10 μ m.

(D) Expression of DCP1-GFP, DCP5-GFP, and XRN4-GFP proteins without and with flg22 treatment. Protoplasts expressing *DCP1-GFP*, *DCP5-GFP*, or *XRN4-GFP* were treated without or with 100 nM flg22 for 30 min, and resulting protein extracts were subjected to immunoblotting (IB) with an α -GFP antibody. Protein loading is shown by Ponceau S staining for RuBisCo (RBC).

(E) The flg22-induced P-body dynamics in *35S::DCP1-GFP* transgenic plants. One-week-old seedlings were treated with 100 nM flg22 for indicated times. Bar, 20 μ m. DIC is differential interference contrast microscopy. Quantification of DCP1-GFP-labeled fluorescence foci is shown on the right.

(F) P-body dynamics induced by *Pst hrcC* infection. Leaves of 4-week-old *35S::DCP1-GFP* transgenic plants were hand-inoculated with *Pst hrcC* at OD₆₀₀ = 0.1, and imaged at the indicated time points. Quantification is shown on the right.

Data in (A), (E), and (F) are shown as mean $\pm$ SD (n = 10). The above experiments were repeated three times with similar results.

See also Figure S1.

presence of multiple MAMPs in the nonpathogenic bacterium. The flg22-induced mobility shift of DCP1-HA was largely restored to the unmodified form after the treatment of lambda protein phosphatase (λ PP) or calf intestinal alkaline phosphatase (CIP) (Figure S2C), implying that DCP1 undergoes phosphorylation modification upon flg22 perception. MAPK cascade is a key signaling module in plant PTI responses (Meng and Zhang, 2013; Tena et al., 2011). The flg22-induced phosphorylation of DCP1-HA was blocked by either a MAPK kinase (MEK) inhibitor PD184161 or co-expression of the MAPK-specific phosphatase MKP (Figure S2D). Furthermore, expression of the full-length MEKK1, a top tier kinase in the flg22-activated MAPK cascade, triggered constitutive DCP1-HA phosphorylation

without flg22 treatment (Figure S2D). Together, these data suggest that the flg22-induced MAPK cascade is required for DCP1 phosphorylation.

Sequential phosphorylation of two canonical MAPK cascades upon MAMP perception leads to the activation of MPK3/MPK6 and MPK4, respectively (Bi et al., 2018; Meng and Zhang, 2013; Sun et al., 2018; Tena et al., 2011). The flg22-induced DCP1 phosphorylation was unaffected in the *mpk3*, *mpk6*, or *mpk4* single mutants (Figures S2E and S2F). MPK3 and MPK6 play redundant roles in plant growth and immunity, and the null mutant of *mpk3mpk6* is embryonic lethal (Meng and Zhang, 2013). The flg22-induced DCP1 phosphorylation was largely reduced in the *mpk6/amiR-MPK3* plants in which MPK3 was

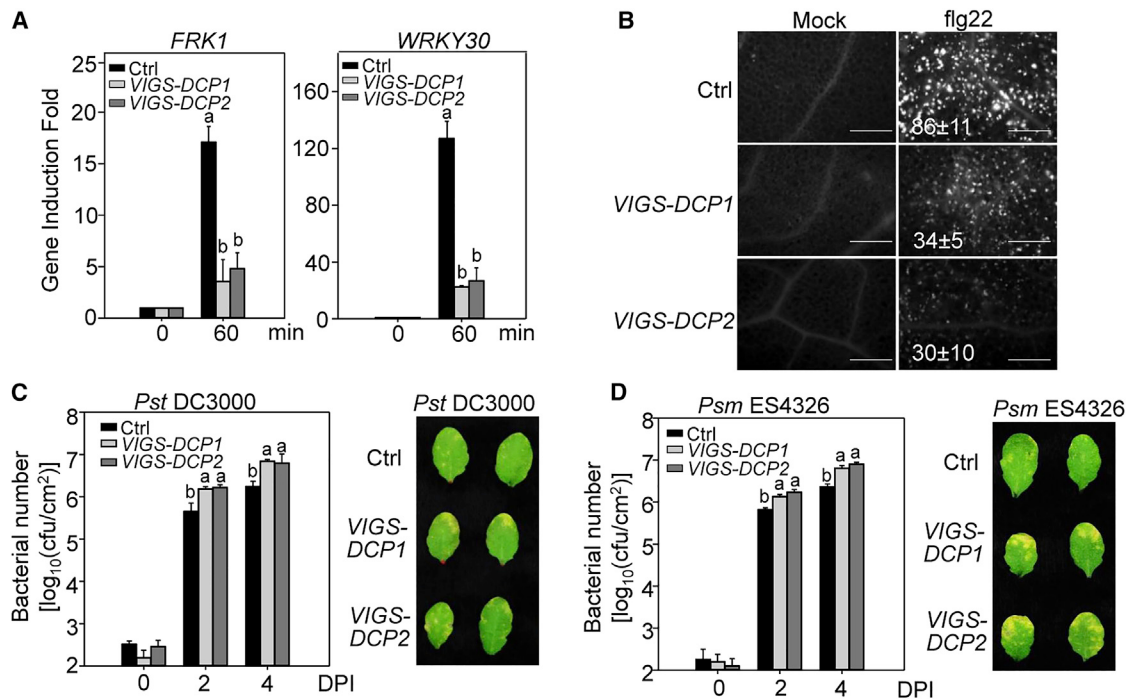


Figure 2. DCP1 and DCP2 Positively Regulate Plant Immunity

(A) Reduced immune gene expression in *DCP1*- and *DCP2*-silenced plants. Leaves from VIGS-silenced *DCP1* and *DCP2* plants at the 4-week-old stage were hand-infiltrated with 100 nM flg22 for 60 min. The VIGS vector containing a GFP fragment was used as control (Ctrl). Data were normalized to the expression of *ACTIN2* in qRT-PCR analysis.

(B) Compromised flg22-triggered callose deposition in *DCP1*- and *DCP2*-silenced plants. Callose deposits were stained with aniline blue 24 h after treatment with 100 nM flg22 or H₂O. The number of callose deposits per mm² is shown as mean ± SD from three biological repeats. Bar, 0.5mm.

(C) Enhanced susceptibility to *Pst* infection in *DCP1*- and *DCP2*-silenced plants. Leaves from 4-week-old plants were hand-inoculated with *Pst* at OD₆₀₀ = 5 × 10⁻⁴, and bacterial counting was performed at 0, 2, and 4 days post-inoculation (dpi) (left). Pictures of inoculated leaves were taken at 4 dpi (right).

(D) Enhanced susceptibility to *Psm* infection in *DCP1*- and *DCP2*-silenced plants.

Data in (A)–(D) are shown as mean ± SD from three independent repeats. Different letters indicate significant differences according to one-way ANOVA test ($p < 0.01$). The above experiments were repeated three times with similar results.

silenced by an artificial microRNA (*amiR-MPK3*) in the *mpk6* background (Figure 3D), suggesting that flg22-induced DCP1 phosphorylation depends on MPK3 and MPK6. We also observed that DCP1-HA was co-immunoprecipitated with MPK3-FLAG and MPK6-FLAG when co-expressed in *Nicotiana benthamiana* (Figure 3E). Apparently, flg22 treatment did not affect DCP1 association with MPK3 or MPK6 (Figure 3E). It has been reported that MPK6 phosphorylates DCP1 at the serine 237 (Ser²³⁷) residue in response to dehydration stress (Xu and Chua, 2012). In line with this observation, we found that DCP1^{S237A}, the phospho-inactive mutant, displayed no mobility shift upon flg22 treatment, whereas DCP1^{S237D}, the phospho-mimetic mutant, showed a constitutive mobility shift in the absence of flg22 treatment (Figure 3F). In addition, flg22-activated MPK3 or MPK6 strongly phosphorylated DCP1 proteins fused to maltose-binding protein (MBP) *in vitro* but to a less extent toward MBP-DCP1^{S237A} proteins (Figure 3G). Both MBP-DCP1 and MBP-DCP1^{S237A} could pull down MPK3 and MPK6 proteins (Figure 3H), suggesting a direct interaction between DCP1 and MPK3 or MPK6. These results indicate that Ser²³⁷ is a major DCP1 phosphorylation site by flg22-activated MPK3 and MPK6.

MAMP-Induced DCP1 Association Dynamics with DCP2 and XRN4

DCP1 functions as a co-activator for DCP2, the major decapping enzyme, to regulate its catalytic activity. In addition, DCP1 interacts with itself and other P-body components, such as the translation repressor DCP5 (Borja et al., 2011; She et al., 2008; Tritschler et al., 2009; Xu and Chua, 2009; Xu et al., 2006). We found here that DCP1 interacted with itself and DCP5 in a flg22-independent manner in *Arabidopsis* (Figures S3A and S3B). DCP1 also co-immunoprecipitated with DCP2 when co-expressed in *Arabidopsis* protoplasts (Figure 4A) or *N. benthamiana* (Figure S3C). Interestingly, flg22 treatment reduced DCP1-DCP2 interaction (Figures 4A and S3C). In line with the observation that flg22 treatment induced DCP1 phosphorylation at Ser²³⁷, DCP1^{S237A} exhibited a stronger association with DCP2 than DCP1^{S237D} in *Arabidopsis* (Figure 4B) and *N. benthamiana* (Figure S3C). An *in vitro* pull-down assay indicated that MBP-DCP1 proteins directly interacted with DCP2 proteins fused with glutathione S-transferase (GST). Similar to *in vivo* results, DCP2 interacted with DCP1^{S237A} stronger than that with DCP1^{S237D} (Figure 4C). The data indicate that DCP1 interacts with DCP2 at the steady state and disassociates from

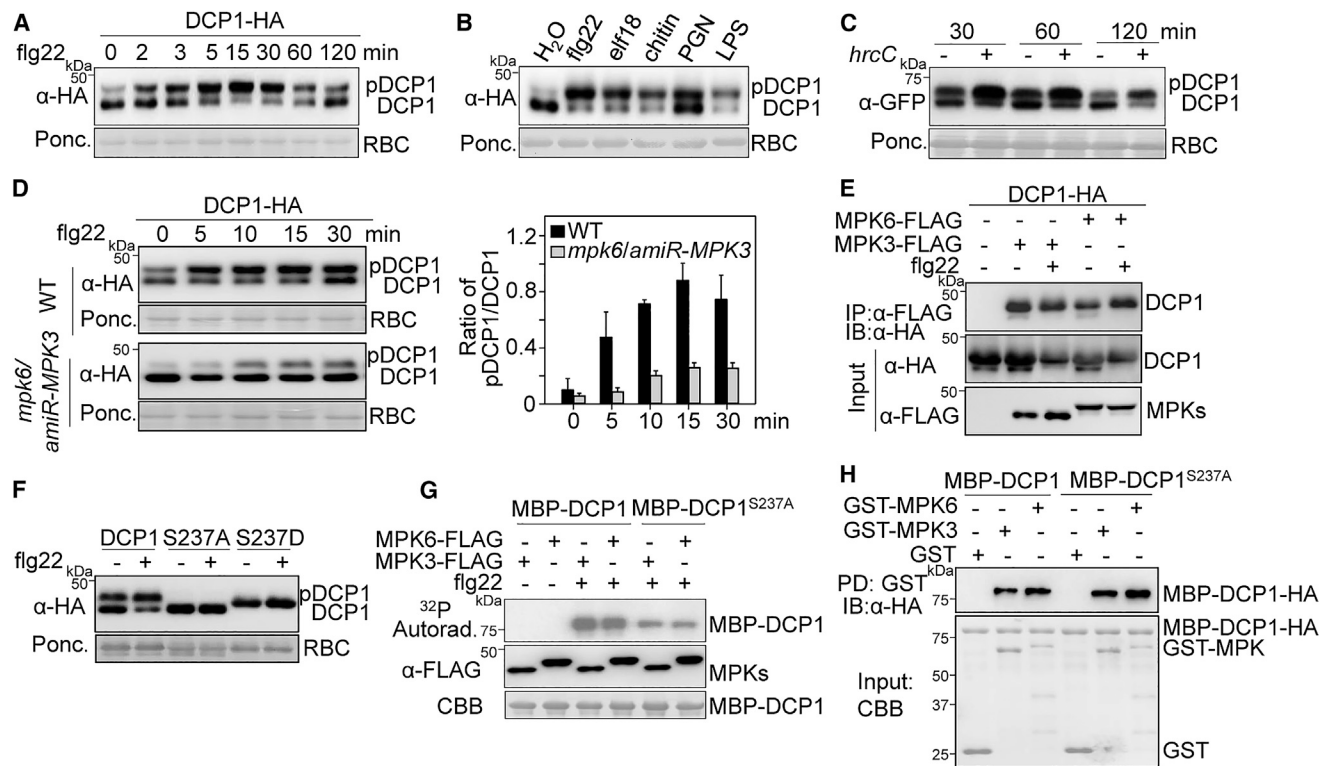


Figure 3. MAMP-Activated MPK3/MPK6 Phosphorylate DCP1 on Ser²³⁷

(A) Mobility shift of DCP1-HA upon flg22 treatment. Protoplasts expressing DCP1-HA were treated with 100 nM flg22 for indicated times. Protein expression was analyzed with an α -HA immunoblot. The intensity of shifted and unshifted polypeptide bands was quantified by ImageJ software, and the percentage of shifted to total proteins (pDCP1/DCP1) is shown on the bottom as a bar graph.

(B) DCP1-HA mobility shift induced by multiple MAMPs. Protoplasts expressing DCP1-HA were treated with 100 nM flg22, 100 nM elf18, 50 μ g/mL chitin, 50 μ g/mL PGN, or 50 μ g/mL LPS for 15 min. The quantification is shown on the bottom.

(C) DCP1 mobility shift induced by *Pst hrcC* in 35S::DCP1-GFP transgenic plants. Leaves from 4-week-old plants were hand-inoculated with H₂O (–) or *hrcC* (+) at OD₆₀₀ = 0.1 for α -GFP immunoblot. The quantification is shown on the bottom.

(D) The flg22-induced DCP1 phosphorylation depends on MPK3 and MPK6. Protoplasts from WT Col-0 and the *mpk6/amiR-MPK3* plants were transfected with DCP1-HA and treated with 100 nM flg22 for the indicated times. The quantification is shown on the right. Data in (A)–(D) are shown as mean $\pm$ SD (n = 3).

(E) DCP1 associates with MPK3 and MPK6 in *N. benthamiana*. *Agrobacterium* carrying 35S::DCP1-HA was co-infiltrated with *Agrobacterium* carrying 35S::MPK3-FLAG, 35S::MPK6-FLAG, or an empty vector (Ctrl.) into leaves of *N. benthamiana*. Leaf samples were collected 2 dpi for co-immunoprecipitation (coIP) assay with α -FLAG (IP: α -FLAG), and the proteins were analyzed by IB with α -HA (top panel). The input control is shown on the bottom two panels.

(F) Ser²³⁷ is important for flg22-induced DCP1 phosphorylation *in vivo*. Protoplasts expressing HA-tagged DCP1, DCP1^{S237A} (phospho-inactive), or DCP1^{S237D} (phospho-mimetic) were treated without or with 100 nM flg22 for 15 min.

(G) Ser²³⁷ is important for MPK3/MPK6-mediated DCP1 phosphorylation. Protoplasts expressing MPK3-FLAG or MPK6-FLAG were treated with 100 nM flg22 for 15 min. MAPK proteins were immunoprecipitated with α -FLAG antibody and subjected to an *in vitro* kinase assay by using MBP-DCP1 or MBP-DCP1^{S237A} as a substrate in the presence of [γ -³²P]-ATP. Proteins were analyzed by SDS-PAGE autoradiography (upper panel), and MAPK expression is shown by immunoblot (middle panel). MBP-DCP1 proteins are shown by Coomassie brilliant blue (CBB) staining.

(H) Both MBP-DCP1 and MBP-DCP1^{S237A} interact with GST-MPK3 and GST-MPK6 *in vitro*. GST, GST-MPK3, or GST-MPK6 immobilized on glutathione Sepharose beads was incubated with MBP-DCP1 or MBP-DCP1^{S237A} proteins. Beads were washed and pelleted for immunoblot analysis with α -HA antibody. PD, pull-down. The above experiments were repeated three times with similar results.

See also Figure S2.

DCP2 upon flg22-activated MAPK-mediated phosphorylation. Remarkably, we consistently observed that flg22 treatment enhanced DCP1 interaction with XRN4 (Figures 4D and S3D), a 5'–3' exonuclease that degrades the decapped mRNAs (Souret et al., 2004). In addition, XRN4 associated with DCP1^{S237D} stronger than that with DCP1^{S237A} in *Arabidopsis* and *N. benthamiana* (Figures 4E and S3D). Similarly, *in vitro* pull-down assays showed that DCP1^{S237D} had a stronger affinity

with XRN4 than DCP1^{S237A} (Figure 4F). Collectively, the results suggest that PRR activation upon MAMP perception triggers rapid phosphorylation of the decapping co-activator DCP1 by MAPKs, which subsequently leads to DCP1 dissociation from the decapping enzyme DCP2 and association with the exonuclease XRN4. MAMP-induced DCP1 dissociation from DCP2 and association with XRN4 might facilitate the engagement of XRN4 in mRNA degradation once mRNAs are

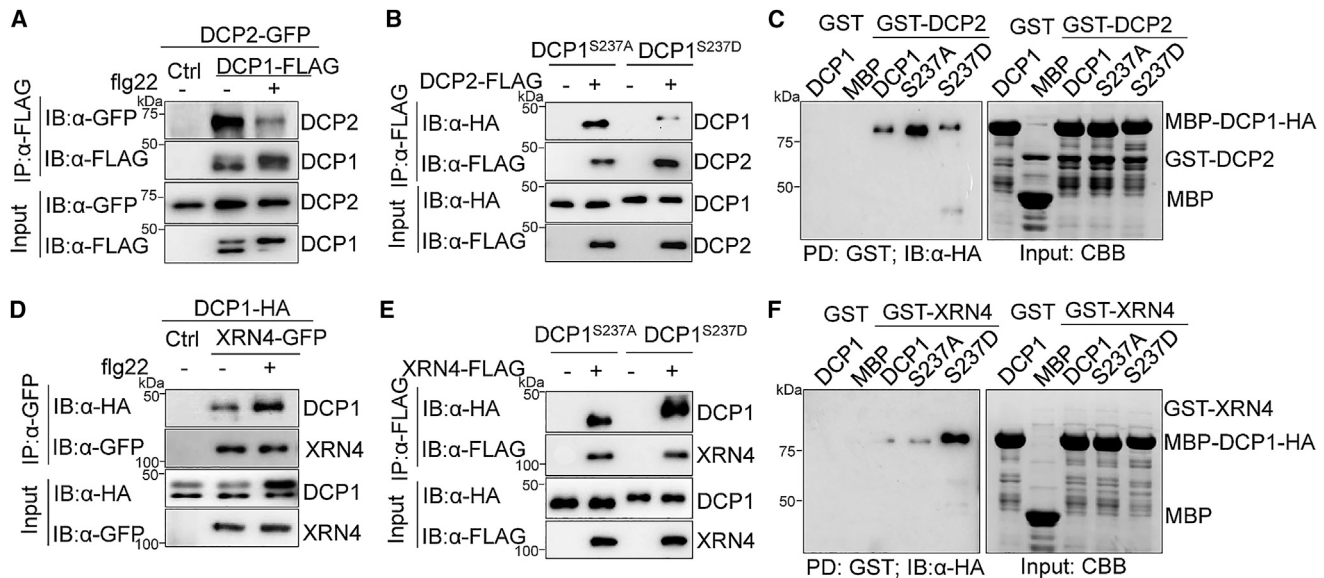


Figure 4. MAMP-Induced DCP1 Association Dynamics with DCP2 and XRN4

(A) Treatment with flg22 induces DCP1-DCP2 dissociation. Protoplasts co-expressing DCP1-FLAG and DCP2-GFP or a vector control were treated with 100 nM flg22 for 15 min. Co-IP was performed with α-FLAG (IP: α-FLAG), and the proteins were analyzed by IB with α-GFP or α-FLAG (top two panels). The input control is shown on the bottom two panels.

(B) DCP1^{S237A} has a higher affinity with DCP2 than DCP1^{S237D} in *Arabidopsis* protoplast co-IP assay.

(C) DCP1^{S237A} shows higher affinity with DCP2 than DCP1^{S237D} *in vitro*. GST or GST-DCP2 immobilized on glutathione Sepharose beads was incubated with MBP, MBP-DCP1, MBP-DCP1^{S237A}, or MBP-DCP1^{S237D} proteins. Beads were washed and pelleted for IB with α-HA (left). Input proteins are shown by CBB staining (right).

(D) Treatment with flg22 enhances DCP1-XRN4 association in *Arabidopsis* protoplast co-IP assay.

(E) DCP1^{S237D} has a higher affinity with XRN4 than DCP1^{S237A} in *Arabidopsis* protoplast co-IP assay.

(F) DCP1^{S237D} shows higher affinity with XRN4 than DCP1^{S237A} *in vitro*. The experiment was carried out as described in (C). The above experiments were repeated three times with similar results.

See also Figure S3.

decapped by DCP2. It is likely that mRNAs are decapped before the DCP1-DCP2 dissociation upon flg22 treatment.

The flg22-Induced DCP1 Phosphorylation Is Required for Its Function in Plant Immunity and P-Body Disassembly

To determine the biological significance of MAMP-induced DCP1 phosphorylation in plant immunity, we complemented the *dcp1-1* mutant with GFP-tagged WT DCP1 or DCP1^{S237A} under the control of its native promoter. Both *pDCP1::DCP1-GFP* and *pDCP1::DCP1^{S237A}-GFP* complementation lines rescued the seedling lethality of the *dcp1-1* mutant and were indistinguishable from WT Col-0 plants at mature stage (Figures S4A, S4B, and S4C), suggesting that phosphorylation at Ser²³⁷ is not required for DCP1 function in plant growth. When challenged with bacterial pathogen *Pst* or *Psm*, the population of bacterial growth in *pDCP1::DCP1-GFP* transgenic lines (L12 and L20) was similar with that in WT plants (Figure 5A). Notably, the *pDCP1::DCP1^{S237A}-GFP* complementation lines (L22 and L31) supported more bacterial growth of *Pst* or *Psm* at 3 dpi than WT and *pDCP1::DCP1-GFP* transgenic lines (Figure 5A). The disease symptom also appeared more severe in the *pDCP1::DCP1^{S237A}-GFP* lines than that in WT and *pDCP1::DCP1-GFP* lines (Figure 5B). In addition, the *pDCP1::DCP1^{S237A}-GFP* transgenic lines

displayed compromised induction of immune-related genes *FRK1* and *PP2C* upon flg22 treatment (Figure 5C). The *pDCP1::DCP1^{S237A}-GFP* transgenic lines deposited less callose upon flg22 elicitation than WT and *pDCP1::DCP1-GFP* transgenic lines (Figure 5D). The number of callose deposits in the *pDCP1::DCP1^{S237A}-GFP* lines was about half of what was observed in WT and *pDCP1::DCP1-GFP* transgenic lines. Furthermore, the flg22 treatment primed plant resistance against *Pst* infection in both WT and *pDCP1::DCP1-GFP* transgenic lines. However, this flg22-induced resistance was compromised in *pDCP1::DCP1^{S237A}-GFP* lines (Figure 5E). These data suggest that phosphorylation of DCP1 at Ser²³⁷ by flg22-activated MPK3/MPK6 is essential for its function in plant immunity.

To determine whether flg22-induced P-body disassembly is dependent on DCP1 phosphorylation at Ser²³⁷, we compared the dynamics of DCP1-GFP- and DCP1^{S237A}-GFP-labeled P-bodies before and after flg22 treatment. Consistently, flg22 treatment reduced both size and number of DCP1-GFP-labeled P-bodies (Figure 5F). However, the reduction was not observed for DCP1^{S237A}-GFP-labeled P-bodies after flg22 treatment (Figure 5F). This result indicates that MAPK-mediated DCP1 phosphorylation is required for flg22-induced P-body disassembly. To gain insights into the possible consequence of MAMP-induced DCP1 association with XRN4, we evaluated the

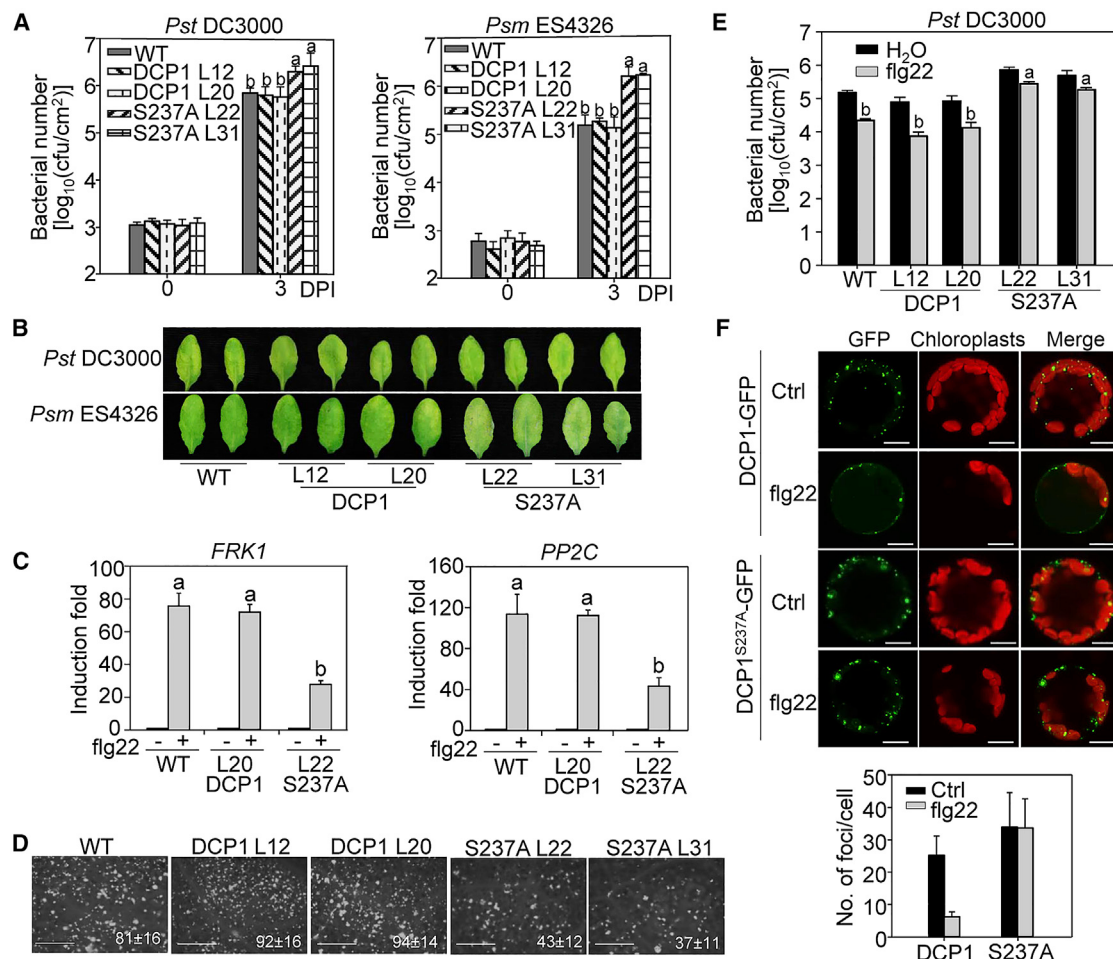


Figure 5. The flg22-Induced DCP1 Phosphorylation Is Required for Its Function in Plant Immunity

(A) DCP1^{S237} is critical for DCP1 function in bacterial disease resistance. Leaves of 4-week-old WT Col-0, *pDCP1::DCP1-GFP/dcp1-1* and *pDCP1::DCP1^{S237A}-GFP/dcp1-1* plants were hand-inoculated with *Pst* (left) or *Psm* (right) at OD₆₀₀ = 5×10^{-4} for bacterial growth assay at 0 and 3 dpi.

(B) Disease symptom of infected leaves. Pictures of inoculated leaves from (A) were taken at 3 dpi.

(C) Reduced immune marker gene expression in *pDCP1::DCP1^{S237A}-GFP/dcp1-1* plants. Leaves of 4-week-old plants were infiltrated with 100 nM flg22 or H₂O for 60 min for qRT-PCR analysis. Data in (A) and (C) are shown as mean $\pm$ SD from three independent repeats. The different letters indicate significant differences according to one-way ANOVA test ($p < 0.01$).

(D) Reduced callose deposition in *pDCP1::DCP1^{S237A}-GFP/dcp1-1* plants. Callose deposits were stained with aniline blue at 24 h after 100 nM flg22 or H₂O treatment in 4-week-old plants. The number of callose deposits per mm² is shown as mean $\pm$ SD from three biological repeats. Bar, 0.5mm. The above experiments were repeated three times with similar results.

(E) Compromised flg22-mediated resistance to *Pst* infection in *pDCP1::DCP1^{S237A}-GFP/dcp1-1* plants. Leaves from 4-week-old plants were hand-inoculated with H₂O or 100 nM flg22 and 24 h later hand-inoculated with *Pst* at OD₆₀₀ = 5×10^{-4} . Bacterial counting was performed at 3 dpi.

(F) flg22-induced P-body disassembly is blocked in *p35S::DCP1^{S237A}-GFP/Col-0* plants. Protoplasts isolated from *p35S::DCP1-GFP/Col-0* and *p35S::DCP1^{S237A}-GFP/Col-0* plants were treated without or with 100 nM flg22 for 30 min. Bar, 10 μ m. Quantification is shown on the bottom as mean $\pm$ SD ($n = 8$). See also Figure S4.

mRNA decay of *ERF1*, *AT2G25250*, *AT1G10120*, and *AT1G30210*, which have been shown to be the targets of XRN4 (Nguyen et al., 2015; Rymarquis et al., 2011), in *pDCP1::DCP1-GFP* and *pDCP1::DCP1^{S237A}-GFP* complementation lines. Interestingly, the mRNA stability of these genes tended to be higher in *pDCP1::DCP1^{S237A}-GFP* than that in *pDCP1::DCP1-GFP* complementation lines (Figure S4D). This result suggests that the interaction between phosphorylated DCP1 and XRN4 may promote XRN4 to degrade its substrates.

DCP1 Phosphorylation-Dependent mRNA Decay Contributes to flg22-Triggered Gene Downregulation

MAMP-induced P-body disassembly may be due to a reduced mRNA pool stored in P-bodies after PTI elicitation. These P-body-associated and translationally suppressed mRNAs are, therefore, either fated for degradation or re-entry to the translation cycle (Decker and Parker, 2012). This P-body-mediated posttranscriptional regulation may allow hosts to launch a rapid and effective immune response by degrading the transcripts of negative regulators and/or timely translation of positive

regulators. We first examined whether flg22 treatment could induce a global translational change. The degree of active translation was estimated by calculating the percentage of cellular RNAs in the polysome fraction (PL%) resolved by sucrose gradient. Results showed no significant difference of PL% between samples treated without or with flg22 for 30, 60, or 180 min (Figures 6A and 6B). Thus, flg22 treatment does not seem to trigger a detectable remodeling of global protein translation. However, we cannot exclude the possibility that specific immune-related transcripts originally stored in P-bodies may be reengaged with polysomes for translation upon MAMP treatment. A recent study shows that elf18 induces translational reprogramming of a suite of *Arabidopsis* genes (Xu et al., 2017).

We then determined whether flg22 perception might regulate the mRNA decay. According to the previous reports (Gutierrez et al., 2002; Narsai et al., 2007), we identified seven genes (*AT3G45970/EXPL1*, *AT2G46690/SAUR32*, *AT2G24570/WRKY17*, *AT3G11410/PP2CA*, *AT3G23030/IAA2*, *AT3G26810/AFB2*, and *AT1G78080/RAP2.4*) that possess highly unstable mRNAs by monitoring the transcript level in the presence of transcriptional inhibitor cordycepin (Figure S5). Significantly, the mRNA decay rates of four genes (*EXPL1*, *SAUR32*, *PP2CA*, and *IAA2*) were accelerated upon flg22 treatment within the 90-min period examined (Figure 6C), whereas the mRNA decay of the other three genes (*WRKY17*, *AFB2*, and *RAP2.4*) was not changed upon flg22 treatment. To determine whether the observed flg22-induced mRNA decay is linked with MAPK-mediated DCP1 phosphorylation, we compared the mRNA decay rates of these genes in transgenic plants expressing DCP1 phospho-inactive mutant (*DCP1^{S237A}*) or phospho-mimetic mutant (*DCP1^{S237D}*). Consistently, the flg22-induced mRNA decay rates of *EXPL1*, *SAUR32*, *PP2CA*, and *IAA2* were higher in *pDCP1::DCP1^{S237D}-GFP/dcp1-1* than those in *pDCP1::DCP1^{S237A}-GFP/dcp1-1* lines (Figure 6D). These results indicate that MAMP treatment induces DCP1 phosphorylation-dependent decay of specific mRNAs.

The mRNA decay constitutes an important step in the regulation of gene expression by maintaining proper mRNA levels both in the steady-state and under perturbations. We determined whether DCP1 and mRNA decay were involved in flg22-triggered downregulation of immune genes. Among flg22-downregulated genes (Li et al., 2015), 6 of 15 examined genes, including *AT1G53830/PME2*, *AT1G73830/BEE3*, *AT2G01850/XTH27*, *AT2G31070/TCP10*, *AT2G45450/ZPR1*, and *AT3G30180/BR6OX2*, showed de-repression in the *dcp1-1* mutant compared to WT plants (Figures 6E and S6A), indicating that flg22-triggered downregulation of these genes depends on DCP1. We further determined whether flg22 treatment promoted mRNA decay of these genes. Among them, *BEE3*, *BR6OX2*, *XTH27*, and *ZPR1* showed accelerated mRNA decay upon flg22 treatment (Figures 6F and S6B). Notably, flg22-induced mRNA decay of these genes was considerably blocked in the *pDCP1::DCP1^{S237A}-GFP/dcp1-1* plants compared to WT plants (Figure 6F). Together, the results indicate that DCP1 phosphorylation-dependent mRNA decay in P-bodies attributes to flg22-triggered gene downregulation (Figure 6G).

DISCUSSION

Activation of PRRs induces a large-scale and dynamic transcriptome reprogramming, culminating in immunity to invading pathogens. The multi-layered transcriptional and posttranscriptional regulation of immune responsive genes is central for launching an effective defense response in hosts. We provide evidence that the components of P-bodies, the cytoplasmic granules that serve as locales for translationally repressed mRNA turnover and storage, play an important role in plant PRR-mediated immunity. P-body assembly is modulated in a dynamic manner during plant immune activation, and P-body components DCP1 and DCP2 are essential in plant immunity. PRR-activated MPK3 and MPK6 directly phosphorylate DCP1, a co-activator for decapping enzyme DCP2, and stimulate DCP1 association with XRN4, an exonuclease executing 5'-to-3' degradation of decapped mRNA (Figure 6G). Complementation analysis indicates that MAMP-induced DCP1 phosphorylation by the MAPK cascade is essential for its function in plant immunity. PRR-activated DCP1 phosphorylation also regulates flg22-induced mRNA decay. Thus, our studies provide a mechanism of immune gene downregulation upon PRR activation and reveal that DCP1 phosphorylation-dependent mRNA decay is an essential host immune response.

Pathogen-induced transcriptome reprogramming leads to dynamic up- and downregulation of a large number of immune-related genes (Li et al., 2016). Downregulation of negative regulators is as crucial as upregulation of positive regulators for hosts to launch an effective defense response. We hypothesize that DCP1 phosphorylation-dependent mRNA decay may facilitate the degradation of the transcripts of negative regulators. The homologs of several DCP1-dependent and flg22 downregulated genes have been shown to negatively regulate plant immunity. *AT1G73830/BEE3* encodes a basic helix-loop-helix (bHLH) transcription factor, which is closely related to *BEE2* and *HBI1*. Both *BEE2* and *HBI1* negatively regulate plant PTI and mediate trade-off between PTI and hormone brassinosteroid (BR)-mediated growth (Fan et al., 2014; Malinovsky et al., 2014). BR perception and signaling inhibit PTI responses (Albrecht et al., 2012; Belkhaoui et al., 2012). *AT3G30180/BR6OX2*, a key BR biosynthetic gene, is downregulated by multiple MAMPs (Jiménez-Góngora et al., 2015). Apparently, suppression of BR biosynthesis by turnover of BR biosynthetic genes is associated with PRR activation. The transient degradation of mRNA upon flg22 treatment may include both positive and negative regulators in the immune signaling, which, in turn, regulates some other immune responses, including gene expression, callose deposition, and immunity.

Nonsense-mediated mRNA decay (NMD) is part of mRNA quality control that prevents translation of aberrant proteins by degrading their mRNAs. It has been shown that NMD is involved in another branch of plant immunity triggered by pathogen effectors by intracellular nucleotide-binding domain leucine-rich repeat (NLR) proteins. NMD controls the constitutive turnover of the transcripts of several Toll/interleukin-1 receptor (TIR) domain-containing NLRs, the overexpression of which often leads to plant autoimmunity (Gloggnitzer et al., 2014). Thus, impairment of NMD leads to a constitutive activation of plant

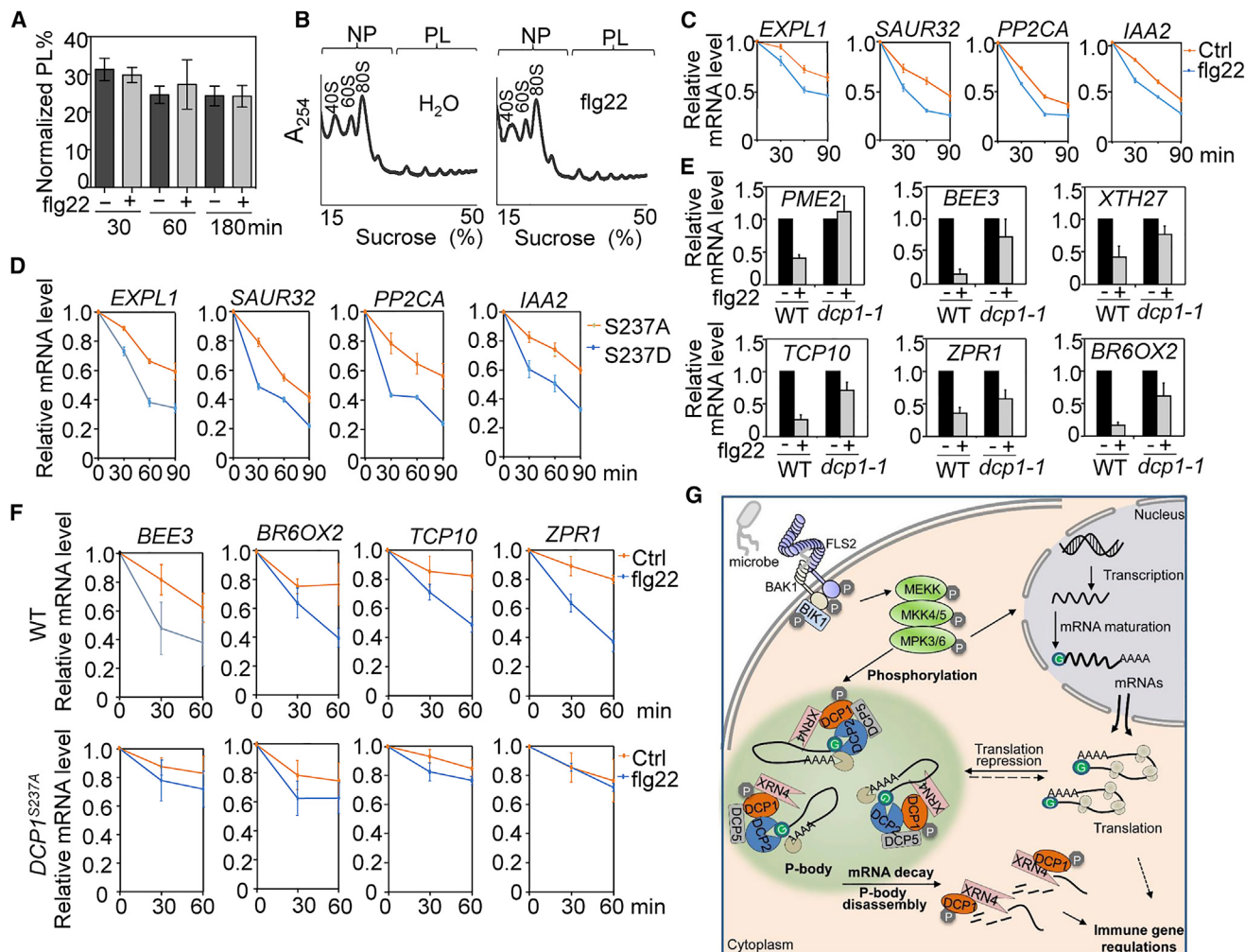


Figure 6. DCP1 Phosphorylation-Dependent mRNA Decay Contributes to flg22-Triggered Gene Downregulation

(A) No detectable global mRNA translational changes upon flg22 treatment. The bar graph shows the RNA loading percentage in polysome fraction (PL%) in WT Col-0 seedlings treated without or with 100 nM flg22 for different times. The data are shown as mean percentage $\pm$ SE from three biological replicates.

(B) A representative polysome profile of WT seedlings treated without or with 100 nM flg22 for 60 min. Ribosome subunits (40S and 60S), mono-ribosome (80S), non-polysomal (NP), and polysomal (PL) fractions are marked. The data are shown with a representative absorbance profile of fractionated ribosomes from three independent biological replicates. A254 indicates absorbance at 254 nm.

(C) Enhanced mRNA decay upon flg22 treatment. Ten-day-old WT seedlings were pre-treated with transcriptional inhibitor cordycepin at 150 μ M for 30 min and then treated without or with 100 nM flg22 for 0, 30, 60, and 90 min. The relative mRNA level at different time points is shown as the ratio to the mRNA level at 0 min (right before flg22 treatment), which is set as 1. *ACTIN2* was used as an internal control.

(D) *pDCP1::DCP1^{S237D}-GFP/dcp1-1* transgenic plants have a faster mRNA decay rate than *pDCP1::DCP1^{S237A}-GFP/dcp1-1*. Ten-day-old seedlings were treated with cordycepin for indicated times for qRT-PCR analysis.

(E) DCP1 is required for flg22-induced gene downregulation. Five-day-old WT and *dcp1-1* seedlings were treated with 100 nM flg22 for 60 min for qRT-PCR analysis. Gene expression level was normalized with *ACTIN2*.

(F) DCP1 phosphorylation-dependent mRNA decay is essential for downregulation of flg22-repressed genes. Ten-day-old seedlings from WT and *pDCP1::DCP1^{S237A}-GFP/dcp1-1* plants were pretreated with cordycepin for 30 min and then treated without or with 100 nM flg22 for indicated times. Data from (C)–(F) are shown as means $\pm$ SD from three independent biological replicates.

(G) A model of P-body dynamics in plant immunity. Plant immunity is initiated by the recognition of MAMP by cell-surface-resident PRR complex, which activates the convergent MAPK cascade. The activated MPK3/MPK6 directly phosphorylate DCP1, the co-activator of decapping enzyme DCP2 in P-bodies, and leads to DCP1 disassociation from DCP2, and association with XRN4, an exonuclease degrading decapped mRNAs. Degradation of mRNAs stored in P-bodies results in P-body disassembly and downregulation of some immune-related genes. Certain translationally repressed mRNAs in P-bodies may also move to polysomes for translation. This P-body-mediated posttranscriptional regulation may provide hosts a means to launch a rapid and effective immune response by degrading the transcripts of negative regulators and/or timely translation of positive regulators.

See also Figures S5 and S6.

immune responses. NMD also serves as a defense system to viral replication by eliminating (+) strand RNAs (Garcia et al., 2014). We show here that the 5'-to-3' exonucleolytic decay, a major mRNA decay pathway, plays an important role in plant cell surface-resident PRR-mediated immunity. Unlike NMD, P-body components positively modulate PTI responses. In addition, the P-body-localized *Arabidopsis* tandem zinc finger 9 (TZF9) is required for PTI responses (Maldonado-Bonilla et al., 2014). Interestingly, TZF9 is also a substrate of MAMP-activated MPK3 and MPK6 (Maldonado-Bonilla et al., 2014). In contrast to the positive role of MPK3 and MPK6, MPK4 negatively regulates plant defense (Petersen et al., 2000). MPK4 phosphorylates protein associated with topoisomerase II 1 (PAT1), a decapping enhancer in RNA decay (Roux et al., 2015). Similar to *mpk4*, the *pat1* mutant displays constitutive defense and autoimmunity in an NLR protein SUMM2-dependent manner (Roux et al., 2015). Flg22-treatment induced PAT1-GFP-labeled P-bodies, likely due to the increased abundance of PAT1 proteins after MAMP treatment (Figures S1C and S1D) (Roux et al., 2015). Apparently, differential phosphorylation of mRNA decay components by MAPKs may specifically affect the fates of targeted immune-related genes upon MAMP treatments.

P-body components also play important roles in plant abiotic stress responses (Perea-Resa et al., 2016; Xu and Chua, 2012). Osmotic-stress-activated subclass I SnRK2s phosphorylate varicose (VCS), an mRNA decapping activator, to regulate mRNA decay under osmotic stress conditions (Soma et al., 2017). Upon dehydration stress, activated MPK6 phosphorylates DCP1 at Ser²³⁷, leading to an enhanced association with DCP5, and likely DCP2, to promote mRNA decapping (Xu and Chua, 2012). Here, we observed that MAMP-activated MPK3 and MPK6 phosphorylate DCP1, leading to its disassociation from DCP2 and association with XRN4. It is intriguing that flg22- and dehydration-stress-activated MPK3/MPK6 phosphorylate DCP1 at the same site but could lead to different interaction dynamics with DCP2 and DCP5. As both elicitors could induce global changes of numerous proteins, it is possible that flg22 or dehydration stress could induce specific proteins or their modifications to differentially affect P-body composition, assembly, and function. In the mRNA decay process, DCP-mediated decapping is followed by exoribonuclease-mediated 5'-to-3' mRNA degradation. In *Drosophila*, DCP1 directly interacts with exoribonuclease 1 and functions as a molecular link between DCP2 activation and the subsequent mRNA turnover (Braun et al., 2012). Our data suggest that upon MAMP perception, DCP1 is rapidly phosphorylated by MAPKs and disengaged with DCP2, then interacts with XRN4, likely to promote the degradation of a subset of mRNAs, and, hence, leads to the disassembly of P-bodies and plant immune activation (Figure 6G). DCP1 and XRN4 association could promote exonuclease activity of XRN4. Alternatively, DCP1 and XRN4 association could increase the accessibility of the decapped mRNAs to XRN4. The interaction between DCP1 and DCP5 (an RNA binding protein) (Xu and Chua, 2009) (Figure S3B) could hold XRN4 in close vicinity to the decapped mRNAs and make the decapped mRNA substrates more accessible to XRN4.

The mechanism underlying P-body assembly and disassembly is still poorly understood. It has been suggested that

the dynamics of assembly results from either protein-protein interaction dynamics or the availability of mRNAs as a platform for protein binding (Standart and Weil, 2018). Accordingly, we observed flg22-induced DCP1 phosphorylation at Ser237 contributes to the degradation of a subset of mRNAs, which may explain the disassembly of P-bodies upon MAMP treatment. Consistently, flg22-induced P-body disassembly was not observed with DCP1^{S237A}-GFP. Furthermore, a subset of P-body-stored translationally repressed mRNAs could re-engaged with polysomes for translation in response to MAMP elicitation. Taken together, our data uncover that modulation of P-body assembly dynamics and DCP1 phosphorylation-dependent mRNA decay activated by a canonical MAPK cascade downstream of multiple PRRs provide a means for hosts to launch a prompt defense response by posttranscriptional reprogramming of primary immune genes.

STAR★METHODS

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SUPPLEMENTAL INFORMATION

Supplemental Information can be found online at <https://doi.org/10.1016/j.celrep.2019.07.054>.

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AUTHOR CONTRIBUTIONS

X.Y., B.L., L.S., and P.H. conceived the project, designed experiments, and analyzed data. X.Y., B.L., G.-J.J., and S.J. performed experiments and analyzed data. D.J., J.-C.J., and S.-H.W. analyzed data, provided critical feedback, and helped shape the research. X.Y., B.L., L.S., and P.H. wrote the manuscript with input from all co-authors.

DECLARATION OF INTERESTS

The authors declare no competing interests.

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STAR★METHODS

KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Antibodies		
Anti-HA-Peroxidase	Roche	Cat # 12013819001; RRID:AB_439705
Anti-FLAG-Peroxidase	Sigma-Aldrich	Cat # A8592; RRID:AB_259529
Anti-GFP	Roche	Cat # 11814460001; RRID:AB_390913
Anti-Mouse IgG HRP-linked antibody	Cell Signaling	Cat # 7076; RRID:AB_330924
Anti-FLAG M2 Affinity gel	Sigma-Aldrich	Cat # 2220; RRID:AB_10063035
Protein G Agarose	Roche	Cat # 5015952001
Bacterial and Virus Strains		
<i>E.coli</i> MC1061	Li et al., 2015	N/A
<i>Agrobacterium tumefaciens</i> GV3101	Li et al., 2015	N/A
<i>E. coli</i> BL21	Li et al., 2015	N/A
<i>Pseudomonas syringae</i> pv. <i>tomato</i> DC3000 (<i>Pst</i>)	He et al., 2006	N/A
<i>P. syringae</i> pv. <i>maculicola</i> ES4326 (<i>Psm</i>)	Li et al., 2014a	N/A
<i>Pseudomonas syringae</i> pv. <i>tomato</i> DC3000 (<i>Pst</i>) type III secretion mutant <i>hrcC</i>	He et al., 2006	N/A
Chemicals, Peptides, and Recombinant Proteins		
PD184161	A.G. Scientific	Cat # P-1713
Calf intestinal alkaline phosphatase (CIP)	New England BioLabs	Cat # M0290S
Lambda protein phosphatase (λ PP)	New England BioLabs	Cat # P0753S
Cordycepin	Sigma-Aldrich	Cat # C9137
RiboZol RNA Extraction Reagent	AMRESCO	Cat # N580
RNase-free DNase I	New England BioLabs	Cat # M0303L
flg22	Genscript	N/A
elf18	Genscript	N/A
chitin	Sigma-Aldrich	Cat # C9752
peptidoglycan	Sigma-Aldrich	Cat # 77140
lipopolysaccharide	Sigma-Aldrich	Cat # L2012
IPTG	Sigma-Aldrich	Cat# I6758
Pierce glutathione agarose	Thermo Scientific	Cat# 16101
amylose resin	New England BioLabs	Cat# E8021L
Critical Commercial Assays		
M-MuLV Reverse Transcriptase	New England BioLabs	Cat # M0253L
iTaq SYBR green Supermix	Bio-Rad	Cat # 1725124
Experimental Models: Organisms/Strains		
<i>Arabidopsis thaliana</i> Col-0 wild-type	Li et al., 2015	N/A
<i>Arabidopsis thaliana</i> Ler ecotype	Li et al., 2015	N/A
<i>mpk3</i>	ABRC	SALK_151594
<i>mpk6</i>	ABRC	Salk_073907
<i>mpk4</i>	ABRC	CS5205, Ler background
<i>fls2</i>	ABRC	Salk_141277
<i>dcp1-1</i>	ABRC	GABI-844B03
<i>p35S::DCP1-GFP/Col-0</i>	This paper	N/A
<i>p35S::DCP1^{S237A}-GFP/Col-0</i>	This paper	N/A
<i>pDCP1::DCP1-GFP/dcp1-1</i>	This paper	N/A

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
<i>pDCP1::DCP1^{S237A}-GFP/dcp1-1</i>	This paper	N/A
<i>pDCP1::DCP1^{S237D}-GFP/dcp1-1</i>	This paper	N/A
<i>mpk6/amiR-MPK3</i>	This paper	N/A
Oligonucleotides		
Primers for cloning, point mutation and VIGS, see Table S1	This paper	N/A
Primers for genotyping and RT-PCR, see Table S1	This paper	N/A
Primers for qRT-PCR, see Table S1	This paper	N/A
Recombinant DNA		
<i>pYL156 (pTRV-RNA2)</i>	Li et al., 2014a	N/A
<i>pTRV-RNA1</i>	Li et al., 2014a	N/A
<i>pYL156-GFP</i>	Li et al., 2014a	N/A
<i>pHBT</i>	He et al., 2006	N/A
<i>pGST</i>	Lu et al., 2010	N/A
<i>pMAL-c2</i>	Lu et al., 2010	N/A
<i>pCB302</i>	Li et al., 2014a	N/A
<i>pHBT-amiRNA-ICE1</i>	Li et al., 2014b	N/A
<i>pTA7002-AvrPto</i>	He et al., 2006	N/A
<i>pYL156-DCP1</i>	This paper	N/A
<i>pYL156-DCP2</i>	This paper	N/A
<i>pHBT-DCP1-GFP</i>	This paper	N/A
<i>pHBT-DCP1-HA</i>	This paper	N/A
<i>pHBT-DCP1-FLAG</i>	This paper	N/A
<i>pHBT-DCP2-GFP</i>	This paper	N/A
<i>pHBT-DCP2-FLAG</i>	This paper	N/A
<i>pHBT-XRN4-GFP</i>	This paper	N/A
<i>pHBT-XRN4-HA</i>	This paper	N/A
<i>pHBT-XRN4-FLAG</i>	This paper	N/A
<i>pHBT-DCP5-GFP</i>	This paper	N/A
<i>pHBT-DCP5-HA</i>	This paper	N/A
<i>pHBT-PAT1-GFP</i>	This paper	N/A
<i>pHBT-DCP1^{S237A}-HA</i>	This paper	N/A
<i>pHBT-DCP1^{S237D}-HA</i>	This paper	N/A
<i>pHBT-MEKK1-FLAG</i>	Li et al., 2015	N/A
<i>pHBT-MPK3-FLAG</i>	Li et al., 2014a	N/A
<i>pHBT-MPK6-FLAG</i>	Li et al., 2014a	N/A
<i>pCB302-35S::DCP1-GFP</i>	This paper	N/A
<i>pCB302-35S::DCP1^{S237A}-GFP</i>	This paper	N/A
<i>pCB302-35S::DCP1^{S237D}-GFP</i>	This paper	N/A
<i>pCB302-35S::DCP2-FLAG</i>	This paper	N/A
<i>pCB302-35S::XRN4-HA</i>	This paper	N/A
<i>pCB302-35S::MPK3-FLAG</i>	This paper	N/A
<i>pCB302-35S::MPK6-FLAG</i>	This paper	N/A
<i>pCB302-pDCP1::DCP1-GFP</i>	This paper	N/A
<i>pCB302-pDCP1::DCP1^{S237A}-GFP</i>	This paper	N/A
<i>pCB302-pDCP1::DCP1^{S237D}-GFP</i>	This paper	N/A
<i>pMAL-DCP1</i>	This paper	N/A
<i>pMAL-DCP1^{S237A}</i>	This paper	N/A
<i>pMAL-DCP1^{S237D}</i>	This paper	N/A

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
pGST-DCP2	This paper	N/A
pGST-XRN4	This paper	N/A
pGST-MPK3	This paper	N/A
pGST-MPK6	This paper	N/A
pTA7002-amiR-MPK3	This paper	N/A
Software and Algorithms		
Olympus Fluoview Viewer	Olympus	Version 3.0
ImageJ	NIH	https://imagej.nih.gov/ij/

LEAD CONTACT AND MATERIALS AVAILABILITY

Further information and requests for resources should be directed to and will be fulfilled by the Lead Contact, Ping He (pinghe@tamu.edu). We will distribute the plasmids and transgenic plants freely to the scientific community upon request.

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Arabidopsis thaliana and growth conditions

All *Arabidopsis* plants used in this study were in the Columbia-0 (Col-0) background, except *mpk4* mutant which was in the *Ler* background. The various mutants and transgenic lines used in this study were described in the [Key Resources Table](#). *Arabidopsis* lines were grown in soil (Metro Mix 366) in a growth room at 23°C, 45% humidity and 85 $\mu\text{E m}^{-2}\text{s}^{-1}$ light with a 12-hr light/12-hr dark photoperiod. Four-week-old plants were used for protoplast isolation and defense-related assays. Seedlings were germinated on 1/2 Murashige and Skoog (MS) plates with 1% sucrose and 0.8% agar, and grown under the same growth condition as above for 10 days. The seedlings were transferred to a 6-well tissue culture plate with 2 mL H₂O for overnight, and then used for indicated assays.

Nicotiana benthamiana and growth conditions

Nicotiana benthamiana was grown in greenhouses in soil under a 12-hr light/12-hr dark photoperiod at 23°C.

Bacterial strains

The various bacteria strains used in this study were described in the [Key Resources Table](#). *Pseudomonas syringae* pv. tomato (*Pst*) DC3000 and *Pst* DC3000 *hrcC* were grown on the King's B medium plates with 50 $\mu\text{g/ml}$ rifampicin. *P. syringae* pv. *maculicola* ES4326 (*Psm*) was grown on the King's B medium plates with 50 $\mu\text{g/ml}$ Streptomycin. All the *Pseudomonas* strains were grown on plates at 28°C for 2 days for further inoculum preparation.

METHOD DETAILS

Constructs and transgenic plant generation

To generate VIGS *DCP1* and *DCP2* constructs, fragments of *DCP1* (~450bp) and *DCP2* (~430bp) were PCR-amplified from *Arabidopsis* Col-0 cDNA, digested with EcoRI and KpnI, and ligated into the *pYL156* (*pTRV-RNA2*) vector. The *DCP1*, *DCP2*, *DCP5*, *XRN4* and *PAT1* genes were amplified from Col-0 cDNA and introduced into the plant gene expression vector *pHBT* with an HA, FLAG or GFP epitope-tag at the C terminus. The point mutations of *DCP1*^{S237A} and *DCP1*^{S237D} were generated by site-directed mutagenesis. To generate constructs for *E. coli* fusion protein isolation, *DCP1*, *DCP1*^{S237A}, *DCP1*^{S237D}, *DCP2*, and *XRN4* were released from *pHBT* vector and subcloned into a modified *pGST* or *pMAL-c2* vector. To generate the 35S promoter-driven genes in the binary vector, *DCP1*, *DCP1*^{S237A}, *DCP1*^{S237D}, *DCP2*, and *XRN4* were sub-cloned into the *pCB302* binary vector with a 35S promoter and an HA, FLAG or GFP epitope-tag at the C terminus. To construct the native promoter-driven genes in the binary vector, the *DCP1*-GFP fragment and its mutants were released from *pHBT*-35S::*DCP1*-GFP, and ligated into the *pCB302* vector. The native promoter of *DCP1* (~660 bp upstream of the start codon) was amplified from Col-0 genomic DNA and ligated into the abovementioned *pCB302*-35S::*DCP1*-GFP construct. The DNA fragments amplified from PCR were fully sequenced to validate the sequences after cloning into the perspective vectors. The *MEKK1*, *MKP*, *MPK3*, *MPK4* and *MPK6* clones in the *pHBT* vector were reported previously (Li et al., 2014a, Li et al., 2015). To generate transgenic plants, the standard protocol for *Agrobacterium tumefaciens*-mediated floral dip was used. The transgenic plants were selected by glufosinate-ammonium (50 $\mu\text{g/ml}$). Multiple transgenic lines were analyzed by immunoblotting for protein expression and two lines with a single T-DNA insertion and similar protein expression levels were chosen for further phenotypic and molecular characterization.

Agrobacterium-mediated floral dipping

Agrobacterium tumefaciens GV3101 containing the binary vector was cultured at 28°C in LB liquid medium with 50 µg/ml Kanamycin and 25 µg/ml Gentamicin. Bacteria were harvested by centrifugation at 3000 rpm for 15 min and the pellet was suspended with buffer containing 50 mM MES (pH 5.5–5.7), 5% sucrose and 200 µl/L silwetL-77 at the density of $OD_{600} = 0.8$. *Arabidopsis* flower buds were dipped thoroughly to the bacteria solution and then the dipped plants were covered with a dome for 24 hr to maintain high humidity. After that, plants were placed in the greenhouses under 12-hr light/12-hr dark light period and seeds were harvested for transgenic plants selection.

Elicitor and chemical inhibitor treatments

MAMPs were used in a final concentration of 100 nM for flg22, 100 nM for elf18, 50 µg/mL for chitin, 50 µg/mL for PGN and 50 µg/mL for LPS. 5 µM of the MEK inhibitor PD184161 was added to protoplasts 1 hr before flg22 treatment. Calf intestinal alkaline phosphatase (CIP) and Lambda protein phosphatase (λPP) treatments were performed following the instruction. The working concentration of transcriptional inhibitor cordycepin was 150 µg/mL for RNA decay assay.

Generation of mpk6/amiR-MPK3 plants and treatment

Artificial microRNA (amiRNA) was constructed as previously described (Li et al., 2014b). In brief, primers for cloning amiRNAs were generated according to the website (<http://wmd3.weigelworld.org/cgi-bin/webapp.cgi>). The cognate fragments were cloned into the modified *pHBT-amiRNA-ICE1* vector (Li et al., 2014b), and then subcloned into *pTA7002-AvrPto* vector (He et al., 2006) with XhoI and StuI digestion to generate *pTA7002-amiR-MPK3* construct. The *pTA7002-amiR-MPK3* construct was transformed into the *mpk6* mutant (SALK_062471) and the transgenic plants were selected with hygromycin resistance. To silence MPK3 expression, the transgenic plants were sprayed with 30 µM dexamethasone supplemented with 0.01% Silwet L-77 at 6 days and 3 days before protoplast isolation.

Callose deposition

Leaves of four-week-old plants were hand-infiltrated with 500 nM flg22 or ddH₂O with a needleless syringe. The treated leaves were harvest 24 h later and then fixed with FAA solution (10% formaldehyde, 5% acetic acid and 50% ethanol), cleared in 95% ethanol, rinsed twice with 75% ethanol and ddH₂O, and stained for 30 min in 0.01% aniline blue solution (150 mM KH₂PO₄, pH 9.5). The leaf samples were mounted in 50% glycerol and callose deposits were visualized under a UV filter with a fluorescence microscope. The number of callose deposits was counted using ImageJ software and expressed as number/mm² with the mean ± SD from at least three different leaves.

Bacterial infection assay

Pseudomonas syringae pv. *tomato* DC3000 (*Pst*) and *P. syringae* pv. *maculicola* ES4326 (*Psm*) strains were cultured for overnight at 28°C in the King's B medium with appropriate antibiotics. Bacteria were harvested by centrifugation at 3500 rpm for 5 min and washed twice with ddH₂O. The pellet was resuspended with 10 mM MgCl₂ and adjusted to the density of $OD_{600} = 5 \times 10^{-4}$. Leaves of four-week-old plants were hand-infiltrated with bacterial suspension using a needleless syringe. For flg22 protection assay, leaves were pre-infiltrated with 100 nM flg22 or water as a control for 24 h before bacterial pathogen infiltration. To measure bacterial growth, six leaf discs separated as three repeats were ground and serial dilutions were plated on Petri dish plates with medium (1% tryptone, 1% sucrose, 0.1% glutamic acid and 1.5% agar) containing corresponding antibiotics. The plates were cultured in a 28°C incubator for two days and bacterial colony forming units (cfu) were counted.

Virus-induced gene silencing assay

Plasmids containing binary TRV vectors *pTRV-RNA1*, *pTRV-RNA2* (*pYL156*), *pYL156-GFP*, *pYL156-DCP1*, or *pYL156-DCP2* were introduced into *Agrobacterium tumefaciens* strain GV3101 by electroporation. *Agrobacterium* culture was grown at 28°C in LB medium containing 50 µg/ml Kanamycin and 25 µg/ml Gentamicin. Bacteria were pelleted by centrifugation, and re-suspended in the induction medium (10 mM MgCl₂, 10 mM MES and 200 µM acetosyringone). Cell suspensions were adjusted to a final concentration $OD_{600} = 1.5$ and then incubated at room temperature for at 3 hr. *Agrobacteria* containing *pTRV-RNA1* and derivatives of *pTRV-RNA2* were mixed at a 1:1 ratio, infiltrated into the first pair of true leaves of two-week-old soil-grown plants with a needleless syringe. Two weeks after infiltration, *pYL156-DCP1*, *pYL156-DCP2* and *pYL156-GFP* (control) inoculated plants were used for PTI response assays and pathogen infection.

RNA isolation, qRT-PCR analysis, and mRNA half-life determination

To determine flg22-induced immune gene expression, leaves of four-week-old plants were hand-infiltrated with 100 nM flg22 or ddH₂O with a needleless syringe and samples were harvested at 1 hour post infiltration (hpi). To measure the unstable mRNA decay rate, ten-day-old seedlings from 1/2 MS plates were transferred to ddH₂O for overnight. Before the transcriptional inhibitor cordycepin (150 µg/ml) treatment, seedlings were pre-soaked in the incubation buffer (1 mM PIPES, pH = 6.25, 1 mM sodium

citrate, 1 mM KCl, 15 mM sucrose, and 0.08% Silwet L-77) for 30 min. Total RNAs from plant samples were extracted by RiboZol RNA Extraction Reagent and quantified with NanoDrop (Thermo Scientific). RNA was treated with RNase-free DNase I to remove DNA contamination. The first strand cDNA was synthesized by reverse transcription with M-MuLV reverse transcriptase and oligo (dT) primer. qRT-PCR analysis was performed using iTaq SYBR green Supermix with an ABI GeneAmp® PCR System 9700 following a standard protocol. The expression of each gene was normalized to the expression of internal control *ACTIN2*. The percentages of mRNAs at each time point calculated relative to the zero time point value were plotted against time and a regression curve was obtained.

Confocal microscopy and image analysis

For transient protoplast expression, protoplasts were transfected with GFP-tagged *DCP1*, *DCP5*, *XRN4* or *PAT1* vectors and were incubated for 12 hr in a 12-well tissue culture plate followed by flg22 treatment. For *pCB302-35S::DCP1-GFP/Col-0* and *pCB302-35S::DCP1^{S237A}-GFP/Col-0* transgenic plants, five-day-old seedlings or protoplasts isolated from four-week-old plants were used for flg22 treatment, and leaves of four-week-old plants were used for *Pst hrcC* infiltration. The fluorescence images were taken by the FLUOVIEW FV1200 (Olympus) confocal system. The excitation wavelength is 488 nm for GFP, and 635 nm for Chlorophyll, which serves as the internal control. Emission is detected at 505–525 nm for GFP, and 660–680 nm for Chlorophyll. Images were converted to 8-bit grayscale with ImageJ software. To quantify the number of P-bodies, the threshold was adjusted to allow the cytoplasmic foci above a certain intensity to be detected. The same threshold was applied to all samples in one experiment. The number of P-bodies was counted using the “analyse particles” function and particle size was limited to those greater than 0.2 μm^2 to get rid of noises. Quantification data are shown as mean $\pm$ SD from at least 10 cells or images.

In vivo co-immunoprecipitation (Co-IP) assay

Arabidopsis protoplasts were transfected with a pair of constructs tested (the empty vector as the negative control) and incubated for 12 hr. Samples were collected by centrifugation and lysed with Co-IP buffer (20 mM Tris-HCl, pH7.5, 100 mM NaCl, 1 mM EDTA, 10% Glycerol, 0.5% Triton X-100 and protease inhibitor cocktail from Roche) by vortexing. For Co-IP in *Nicotiana benthamiana*, leaves of three-week-old soil-grown plants were hand-infiltrated with different pairs of *Agrobacterium tumefaciens* carrying indicated vectors. Overnight cultured bacteria were harvested by centrifugation and re-suspended in buffer (10 mM MES, pH5.7, 10 mM MgCl_2 , 200 μM acetosyringone) at $\text{OD}_{600} = 1.5$. Leaf samples were harvested two days post inoculation and subjected to homogenization with Co-IP buffer. Protein extract was pre-incubated with protein-G-agarose beads for 1 hr at 4°C with gentle shaking on a rocker. Immunoprecipitation was carried out with an α -GFP or α -FLAG antibody for 2 hr and then with protein-G-agarose beads for another 2 hr at 4°C. The beads were collected and washed three times with washing buffer (20 mM Tris-HCl, pH7.5, 100 mM NaCl, 1 mM EDTA, 0.1% Triton X-100). The immunoprecipitated proteins and input proteins were analyzed by immunoblotting with indicated antibodies.

In vitro pull-down and immunocomplex kinase assays

Fusion proteins were expressed in *E. coli* BL21 strain using LB medium supplemented with 0.25 mM Isopropyl β -D-1-thiogalactopyranoside (IPTG). Glutathione-S-transferase (GST), GST-MPK3, GST-MPK4 and GST-MPK6 were purified with Pierce glutathione agarose, and maltose binding protein (MBP), MBP-DCP1, MBP-DCP1^{S237A} and MBP-DCP1^{S237D} proteins were purified using amylose resin according to standard protocols. MBP fusion proteins (tagged with HA) were pre-incubated with prewashed glutathione agarose in 300 μL incubation buffer (20 mM Tris-HCl, pH7.5, 100 mM NaCl, 0.1 mM EDTA and 0.5% Triton X-100) for 0.5 hr at 4°C. After centrifugation, the supernatant was collected and incubated with prewashed GST, GST-MPK3 or GST-MPK6 beads for another 1 hr. The beads were collected and washed three times with washing buffer (20 mM Tris-HCl, pH7.5, 300 mM NaCl, 0.1 mM EDTA and 0.1% Triton X-100). Proteins were detected with an α -HA antibody by immunoblotting. For immunocomplex kinase assay, FLAG-tagged MPK3, MPK4, and MPK6 were expressed in Col-0 protoplasts for 12 hr and then treated with 100 nM flg22 for 15 min. The activated MPK proteins were immunoprecipitated using α -FLAG agarose following Co-IP procedure. The beads were spun down and re-suspended in 50 mM Tris-HCl, pH7.5. The MPK-bounded beads were incubated with MBP-DCP1 or MBP-DCP1^{S237A} fusion proteins in the kinase reaction buffer (20 mM Tris-HCl, pH7.5, 20 mM MgCl_2 , 5 mM EDTA, 1 mM DTT and 100 μM ATP) in the presence of 5 μCi [^{32}P]- γ -ATP for 2 hr at room temperature. The reactions were stopped by adding SDS sample buffer, and protein phosphorylation was visualized by autoradiography in 10% SDS-PAGE.

Analysis of translation efficiency

Ribosome profiles were performed as described previously (Liu et al., 2012). In brief, polysomal RNA was extracted with polysome extraction buffer. The resuspended mixture was incubated on ice for 5 min, and then spun at 15000 g for 5 min at 4°C. Supernatant was loaded on a continuous sucrose gradient (15%–50%) and spun at 210000 g for 3.5 hr at 4°C. The distribution of the nucleic acids was examined by a UV₂₅₄ absorbance profile (model #UV-6, ISCO). Total RNAs from non-polysome fractions (NP) or polysome fractions (PL) were used for the calculation of PL% (PL/NP+PL) after normalization by spike-in RNA DAP (Affymetrix, GeneChip Poly-A RNA Control Kit).

QUANTIFICATION AND STATISTICAL ANALYSIS

In general, data are represented by mean $\pm$ SD. The numbers of biological repeats (n) are provided in each figure legend. One-way analysis of variance (ANOVA) was performed to check for statistically significant differences. For qRT-PCR analysis, each biological repeat is based on an RNA sample extracted from four seedlings.

DATA AND CODE AVAILABILITY

This manuscript did not generate new datasets or code.