

MEKK1, MKK1/MKK2 and MPK4 function together in a mitogen-activated protein kinase cascade to regulate innate immunity in plants

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Mitogen-activated protein kinase (MAPK) cascades play important roles in regulating plant innate immune responses. In a genetic screen to search for mutants with constitutive defense responses, we identified multiple alleles of *mpk4* and *mekk1* that exhibit cell death and constitutive defense responses. Bimolecular fluorescence complementation (BiFC) analysis showed that both MPK4 and MEKK1 interact with MKK1 and MKK2, two closely related MAPK kinases. *mekk1* and *mekk2* single mutant plants do not have obvious mutant phenotypes. To test whether MKK1 and MKK2 function redundantly, *mekk1 mekk2* double mutants were generated. The *mekk1 mekk2* double mutant plants die at seedling stage and the seedling-lethality phenotype is temperature-dependent. Similar to the *mpk4* and *mekk1* mutants, the *mekk1 mekk2* double mutant seedlings accumulate high levels of H₂O₂, display spontaneous cell death, constitutively express *Pathogenesis Related (PR)* genes and exhibit pathogen resistance. In addition, activation of MPK4 by *flg22* is impaired in the *mekk1 mekk2* double mutants, suggesting that MKK1 and MKK2 function together with MPK4 and MEKK1 in a MAP kinase cascade to negatively regulate innate immune responses in plants.

Keywords: MAPK, innate immunity, MPK4, MEKK1, MKK1, MKK2

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Introduction

Eukaryotes use different receptors to recognize microbial pathogens [1]. There are two types of immune receptors in plants. One type recognizes microbe-associated molecular patterns (MAMPs) and initiates downstream immune responses. For example, *Arabidopsis* FLS2 is required for the recognition of bacterial flagellin and EFR is required for the recognition of bacterial elongation factor EF-Tu [2, 3]. Both FLS2 and EFR belong to the leucine-rich-repeat (LRR)-containing receptor-like kinase family. The other type of immune receptors is the plant resistance (R) proteins. They are thought to recog-

nize effector proteins that are delivered by pathogens to suppress MAMP-triggered immune responses. Compared with MAMP-triggered immune responses, R protein-mediated resistance responses are generally much stronger and often lead to hypersensitive response (HR), a localized programmed cell death event that restricts pathogen growth.

In *Arabidopsis*, several mutations in R genes have been identified that lead to constitutively activated downstream immune responses. Constitutive activation of these immune receptors has detrimental effects on plant growth and development. *ssi4*, *slh1* and *snc1* all contain mutations in TIR-NB-LRR-type R genes [4-6]. Both *ssi4* and *slh1* mutants show spontaneous cell death. Although *snc1* mutant plants do not show cell death phenotypes, they accumulate high levels of salicylic acid (SA) and have reduced stature. Recently, a semi-dominant mutant *uni-1D* was found to contain a gain-of-function mutation in CC-NB-LRR-type R genes [7]. The *uni-1D* mutation causes constitutive activation of defense responses and

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morphological defects through accumulation of abnormal levels of cytokinin. Constitutive activation of MAMP receptor FLS2 by growing plants on medium containing flg22 (a peptide derived from *Escherichia coli* flagellin) also results in inhibition of plant growth [2]. To prevent the detrimental effects caused by activation of defense responses, plant immune responses must be under tight negative regulations in the absence of pathogen infection.

Activation of mitogen-activated protein kinase (MAPK) cascades is a common mechanism for regulating innate immune responses in both animals and plants [8]. *Ara-*

bidopsis MPK4 is an important negative regulator of systemic acquired resistance (SAR). *mpk4* mutant plants in Landsberg ecotype (Ler) accumulate high levels of SA and constitutively express *Pathogenesis Related (PR)* genes and exhibit pathogen resistance, but have no necrotic lesions [9]. *Arabidopsis* MEKK1, a MAPK kinase kinase, also functions as a negative regulator of plant immune responses. Knockout mutants of *MEKK1* constitutively express *PR* genes and exhibit a seedling-lethality phenotype because of spontaneous cell death [10-12].

To search for additional negative regulators of plant innate immune response, we looked for mutants that con-

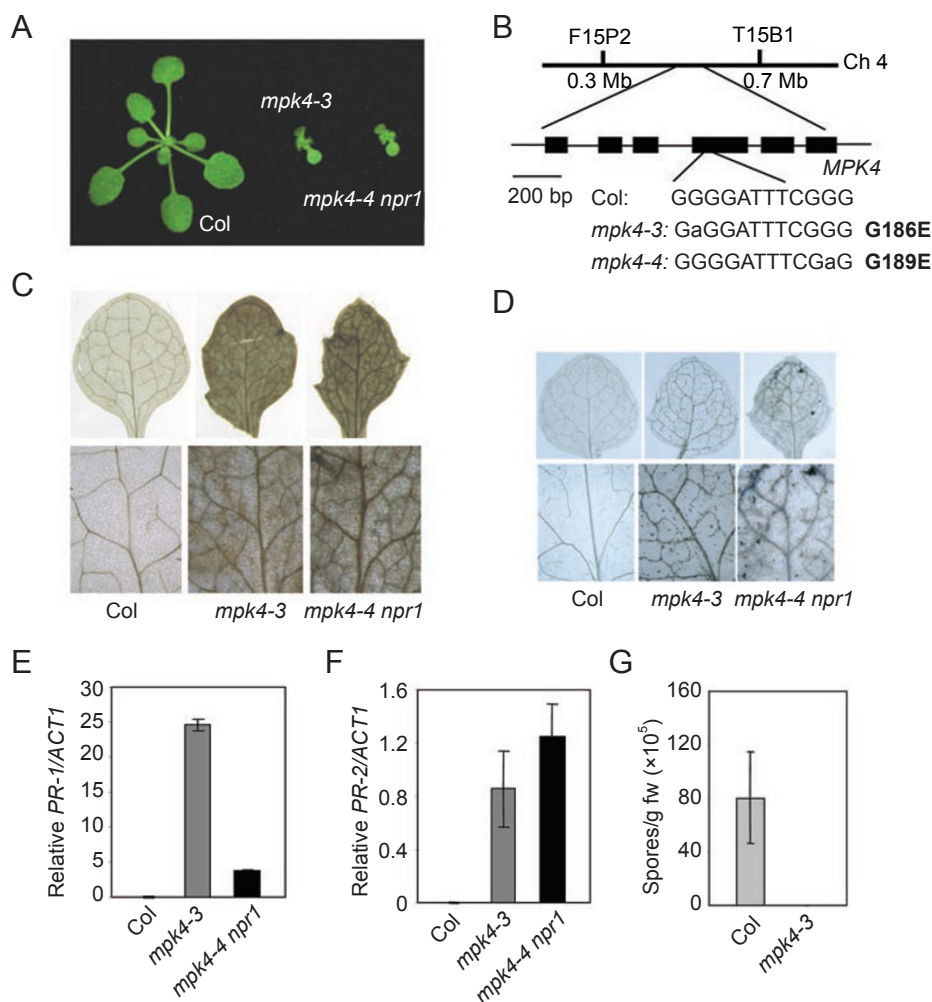


Figure 1 Analysis of *mpk4-3* and *mpk4-4 npr1-1* mutants. **(A)** Phenotypes of wild-type, *mpk4-3* and *mpk4-4 npr1-1*. Plants were grown on soil and photographed 3 weeks after planting. **(B)** Map position and mutations in *mpk4-3* and *mpk4-4*. **(C)** DAB staining of wild-type, *mpk4-3* and *mpk4-4 npr1-1* mutant seedlings. **(D)** Trypan blue staining of wild-type, *mpk4-3* and *mpk4-4 npr1-1* mutant seedlings. **(E)** *PR-1* and **(F)** *PR-2* expression in the wild-type, *mpk4-3* and *mpk4-4 npr1-1* mutant seedlings. Error bars represent standard deviation from three measurements. Values were normalized to the expression of *ACTIN1*. **(G)** Growth of *H. parasitica* Noco2 on wild type and *mpk4-3*. Three-week-old seedlings were sprayed with *H. parasitica* Noco2 spores (10^5 spores/ml). Infection was scored 7 days after inoculation by counting the number of conidia spores per gram of leaf samples. Error bars represent standard deviation from three measurements.

stitutively express *PR* genes and exhibit pathogen resistance. We screened an EMS-mutagenized mutant population and identified multiple mutant alleles of *mpk4* and *mekk1*. Unlike the original *mpk4* mutant identified in Ler background, the *mpk4* mutant alleles we obtained in the Columbia background were more like the *mekk1* knockout mutants and died at seedling stage because of spontaneous cell death. We found that both MPK4 and MEKK1 interact with two MAPK kinases, MKK1 and MKK2, *in vivo*, and *mekk1 mkk2* double mutant plants exhibit very similar phenotypes to *mekk1* and *mpk4* mutants. Our results suggest that a MAP kinase cascade consisting of MEKK1, MKK1/MKK2 and MPK4 negatively regulates plant innate immunity.

Results

Identification and characterization of alleles of *mpk4*

To identify mutants that are constitutively resistant to pathogens, progeny from individual EMS-mutagenized M1 mutant lines in Columbia background were analyzed for resistance to *Hyaloperonospora parasitica* Noco2 (*H. p.* Noco2). One of the mutants resistant to *H. p.* Noco2 was found to die at the seedling stage (Figure 1A). We mapped the mutation to a 400-kb region on chromosome IV, where *MPK4* is localized. Sequence analysis revealed a point mutation in the mutant; thus we named it *mpk4-3* (Figure 1B). 3, 3'-Diaminobenzidine (DAB) staining showed that the *mpk4-3* plants also accumulated high levels of H₂O₂ (Figure 1C). Trypan blue staining showed that there was extensive cell death in the *mpk4-3* mutant (Figure 1D). Both *PR-1* and *PR-2* are constitutively expressed in the *mpk4-3* mutant (Figure 1E and 1F). In addition, the *mpk4-3* mutant plants are constitutively resistant to *H. p.* Noco2 (Figure 1G).

We also identified another allele of *mpk4* (*mpk4-4*) in a suppressor screen of *npr1-1* (*nonexpresser of PR genes 1, 1*) to look for mutants that constitutively express the *BGL2-GUS* reporter gene in the *npr1-1* mutant background. The *npr1-1* mutation blocks SA-induced *PR* gene expression and the *BGL2-GUS* reporter gene expression. Like *mpk4-3*, the *mpk4-4* mutant accumulates high levels of H₂O₂, shows extensive cell death and exhibits seedling-lethality phenotypes (Figure 1A, 1C and 1D). Whereas *PR-2* is constitutively expressed in the *mpk4-4 npr1-1* mutant plants, *PR-1* expression is blocked by the *npr1-1* mutation (Figure 1E and 1F). Through similar mapping and targeted gene-sequencing approaches, two other mutants that constitutively express the *BGL2-GUS* reporter gene in the *npr1-1* mutant background and exhibit seedling-lethality phenotypes were later found to be alleles of *mekk1* (Figure 2). Interesting-

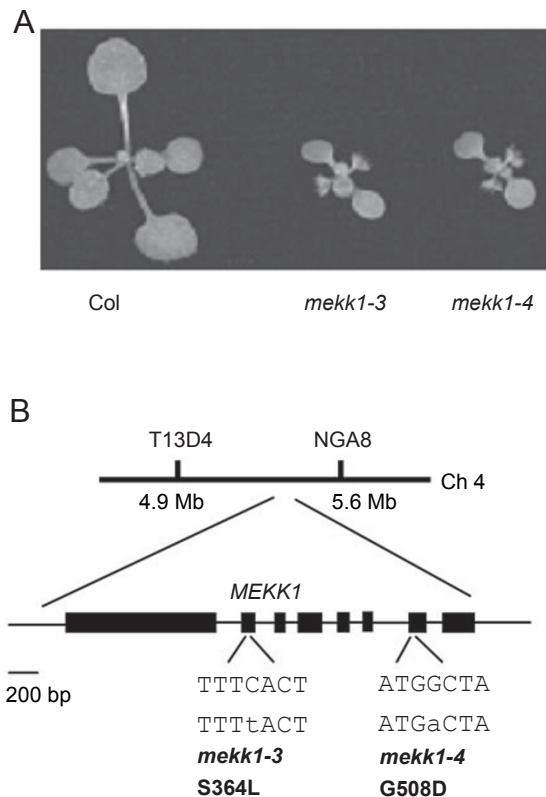


Figure 2 Analysis of *mekk1-3* and *mekk1-4* mutants. **(A)** Phenotypes of wild type, *mekk1-3* and *mekk1-4*. Plants were grown on soil and photographed 3 weeks after planting. **(B)** Map position and mutations in *mekk1-3* and *mekk1-4*.

ly, the seedling-lethality phenotypes of *mpk4* and *mekk1* alleles are temperature dependent (observed at 22 °C). At 28 °C, these mutant plants can complete their life cycles and set seeds.

MKK1 and MKK2 interact with MPK4 and MEKK1 in vivo

MKK1 and MKK2 are two closely related MAP kinase kinases that interact with MPK4 and MEKK1 in yeast two-hybrid assays [13-15]. To test whether MKK1 and MKK2 associate with MPK4 and MEKK1 *in vivo*, we analyzed their interactions using a bimolecular fluorescence complementation (BiFC) approach [16]. MKK1 and MKK2 were fused to the N-terminal fragment of YFP (MKK1-YFP^N and MKK2-YFP^N), whereas MEKK1 and MPK4 were fused to the C-terminal fragment of YFP (MEKK1-YFP^C and MPK4-YFP^C). If two proteins associate with each other, a fluorescent YFP complex would be formed. As shown in Figure 3, YFP fluorescence was observed on the plasma membrane of *Arabidopsis* mesophyll protoplasts co-transformed with the MKK1-YFP^N

and MEKK1-YFP^C constructs. Similar results were observed when MKK2-YFP^N and MEKK1-YFP^C constructs were co-transformed into the *Arabidopsis* mesophyll protoplasts, suggesting that MKK1 and MKK2 interact with MEKK1 on the plasma membrane. As negative controls, MKK1-YFP^N or MKK2-YFP^N was co-transformed with the YFP^C vector and no YFP fluorescence was observed. Co-transformation of MEKK1-YFP^C and the YFP^N vector into protoplasts did not result in YFP fluorescence either.

In contrast, YFP fluorescence was observed on the plasma membrane and in the nucleus of *Arabidopsis* mesophyll protoplasts co-transformed with the MPK4-YFP^C and MKK1-YFP^N or MKK2-YFP^N constructs (Figure 3), suggesting that MPK4 interacts with MKK1 and MKK2 on the plasma membrane and in the nucleus. As negative controls, MPK4-YFP^C was transformed into protoplasts

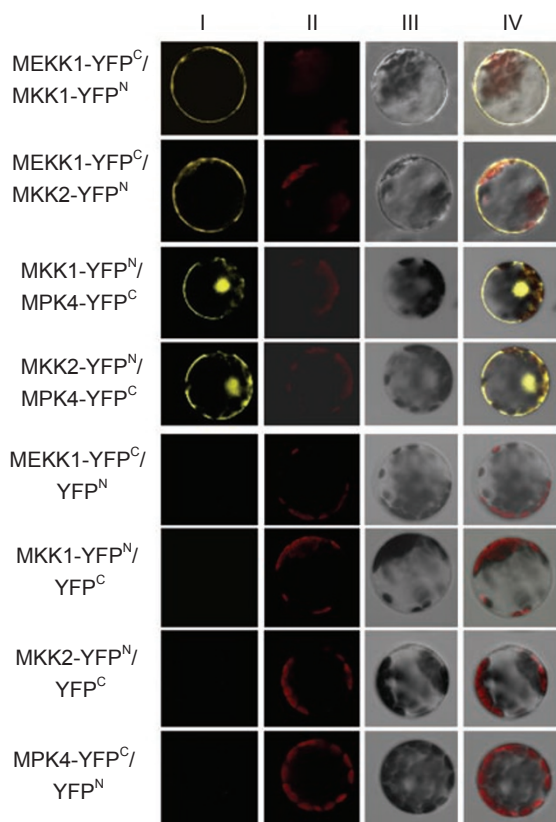


Figure 3 Analysis of MKK1 and MKK2 interactions with MEKK1 and MPK4 by BiFC. Epifluorescence (I), chloroplast autofluorescence (II), bright field (III) and merged (IV) images of *Arabidopsis* mesophyll protoplasts co-transformed with constructs expressing different fusion proteins (top four panels). Vectors expressing the N- or C-terminal domains of YFP alone with each corresponding fusion construct were used as negative controls (lower four panels).

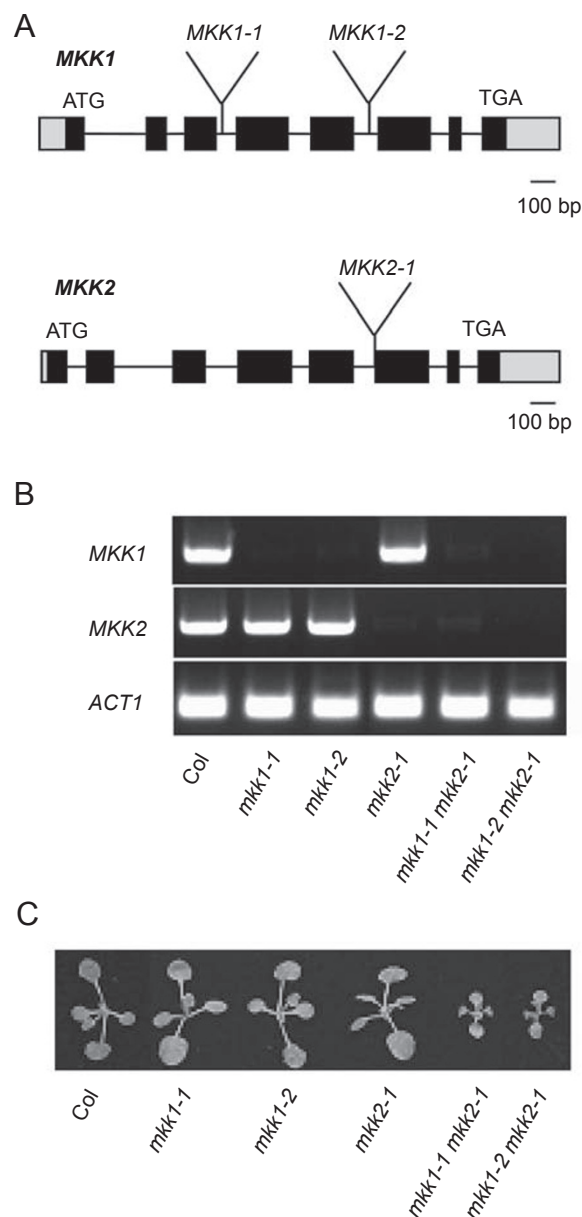


Figure 4 Analysis of the *mkk1* and *mkk2* insertion mutants. **(A)** Positions of T-DNA insertions within the *MKK1* and *MKK2* genes in *mkk1* and *mkk2* mutants. **(B)** RT-PCR analysis of *MKK1* and *MKK2* expression in the knockout mutants. **(C)** Phenotypes of wild-type, *mkk1*, *mkk2* and *mkk1 mkk2* double mutants. Plants were grown on soil and photographed 3 weeks after planting.

together with the YFP^N vector and no YFP fluorescence was observed.

mkk1 mkk2 double mutant plants exhibit lethality at seedling stage

To test whether MKK1 and MKK2 have similar functions as MEKK1 and MPK4 in regulating cell death and

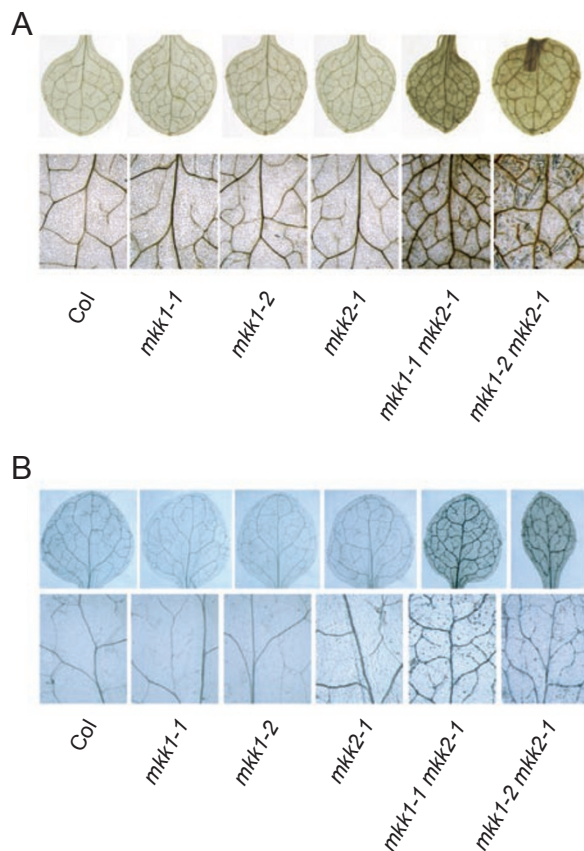


Figure 5 DAB staining (A) and trypan blue staining (B) of wild-type, *mkk1*, *mkk2* and *mkk1 mkk2* double mutant seedlings.

defense responses, T-DNA insertion knockout mutants of *MKK1* and *MKK2* were obtained from ABRC. The locations of the T-DNA insertion are shown in Figure 4A. RT-PCR analysis of *MKK1* and *MKK2* in the knockout mutant indicates that expression of the target genes is abolished by the T-DNA insertions (Figure 4B). As *MKK1* and *MKK2* have high sequence similarity and single knockout mutants do not exhibit any obvious mutant phenotypes, *mkk1 mkk2* double mutants were obtained by crossing the single mutants. As shown in Figure 4C, *mkk1-1 mkk2-1* and *mkk1-2 mkk2-1* double mutant plants are dwarf and small. Similar to the *mpk4* and *mek11* mutants, *mkk1 mkk2* double mutant plants exhibit a seedling-lethality phenotype.

As only one *mkk2* allele was identified, we tested whether the morphological phenotypes observed in the *mkk1-1 mkk2-1* double mutant co-segregate with the *mkk2-1* mutation. The progeny of F2 lines that are heterozygous for the *mkk2-1* mutation and homozygous for the *mkk1-1* mutation were planted. Among 160 plants, 36 displayed *mkk1 mkk2* double mutant morphology. PCR analysis showed that all 36 plants were homozygous for the *mkk2-1* mutation, indicating that the morphological phenotypes observed in the *mkk1-1 mkk2-1* double mutant co-segregate with the *mkk2-1* mutation.

Analysis of cell death and H_2O_2 accumulation in *mkk1 mkk2* plants

To test whether cell death occurred in the *mkk1 mkk2*

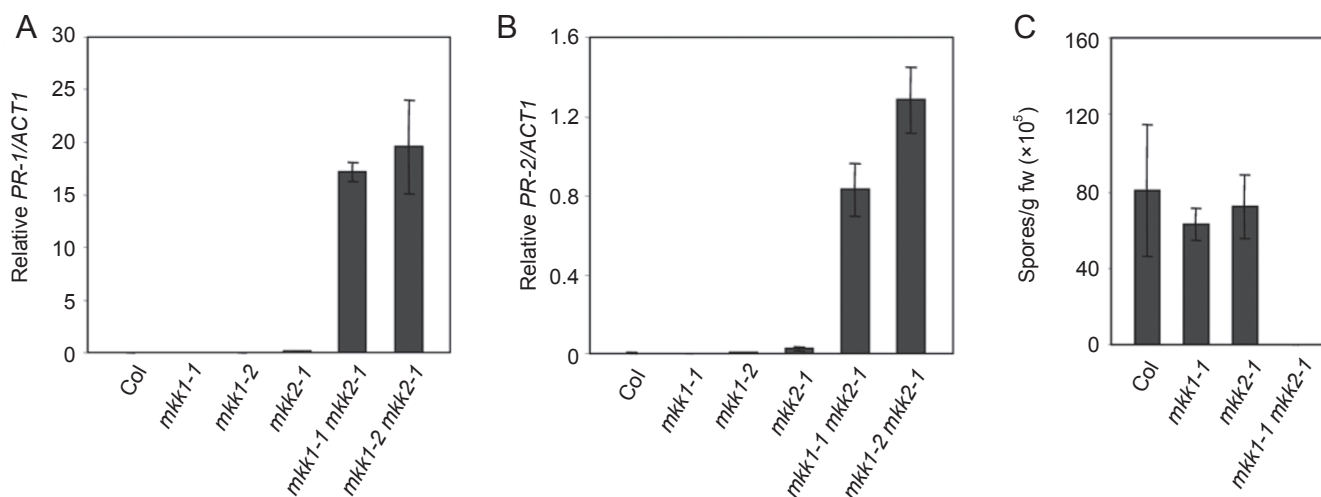


Figure 6 Characterization of the *mkk1 mkk2* double mutant plants. *PR-1* (A) and *PR-2* (B) expression in wild-type, *mkk1*, *mkk2* and *mkk1 mkk2* double mutants. Error bars represent standard deviations from three measurements. Values were normalized to the expression of *ACTIN1*. (C) Growth of *H. parasitica* Noco2 on wild-type, *mkk1*, *mkk2* and *mkk1 mkk2* double mutants. Three-week-old seedlings were sprayed with *H. parasitica* Noco2 spores (10^5 spores/ml). Infection was scored 7 days after inoculation by counting the number of conidia spores per gram of leaf samples. Error bars represent standard deviations from three measurements.

double mutants, trypan blue staining was performed on wild type and mutant seedlings. As shown in Figure 5A, there was extensive cell death in the leaves of the *mkk1 mkk2* double mutant plants, particularly along the veins. In contrast, cell death was not observed in the *mkk1* and *mkk2* single mutants. To check the H₂O₂ levels in the mutant plants, we performed DAB staining on the mutant seedlings. As shown in Figure 5B, strong staining was observed in the *mkk1 mkk2* double mutants, but not in the wild-type and the single mutants, indicating that the *mkk1 mkk2* mutant seedlings accumulated high levels of H₂O₂.

Constitutive defense responses in *mkk1 mkk2* plants

To test whether defense responses were constitutively activated, the expression of *PR* genes in the *mkk1*, *mkk2* and *mkk1 mkk2* mutant plants was determined by real-time RT-PCR. As shown in Figure 6A and 6B, high levels of *PR-1* and *PR-2* were constitutively expressed in the *mkk1 mkk2* double mutant plants. In contrast, *PR-1* and *PR-2* were not constitutively expressed in the wild-type and the single mutant plants, suggesting that loss of function of both *MKK1* and *MKK2* leads to constitutive activation of defense responses.

To test whether the *mkk1* and *mkk2* single mutants and the *mkk1 mkk2* double mutants have enhanced pathogen resistance, mutant seedlings were challenged with the virulent oomycete pathogen *H. p. Noco2*. As shown in Figure 6C, the *mkk1 mkk2* double mutant seedlings showed strong enhanced resistance against *H. p. Noco2* compared with the wild-type and the single mutants, confirming that defense responses were constitutively activated in the *mkk1 mkk2* mutant plants.

MKK1 and *MKK2* are required for *MPK4* activation

MEKK1 has previously been shown to be required for the activation of *MPK4* in response to the treatment of the flagellin peptide *flg22*. To test whether *MKK1* and *MKK2* are also required for the activation of *MPK4* in response to *flg22*, we analyzed the kinase activity of immunoprecipitated *MPK4* from wild-type and mutant plants. Myelin-basic protein was used as substrate for the assay. As shown in Figure 7A, *MPK4* kinase activity was at very low levels when the plants were not treated with *flg22*. Spraying with *flg22* resulted in quick activation of *MPK4* in the wild-type plants. Whereas the activation of *MPK4* was not affected in the single mutants of *mkk1* and *mkk2*, activation of *MPK4* was severely compromised in the *mkk1-1 mkk2-1* double mutant, suggesting that *MKK1* and *MKK2* are required for the activation of *MPK4* by *flg22*. Previously Meszaros *et al.* [17] showed that activation of *MPK4* was blocked in *mkk1* single

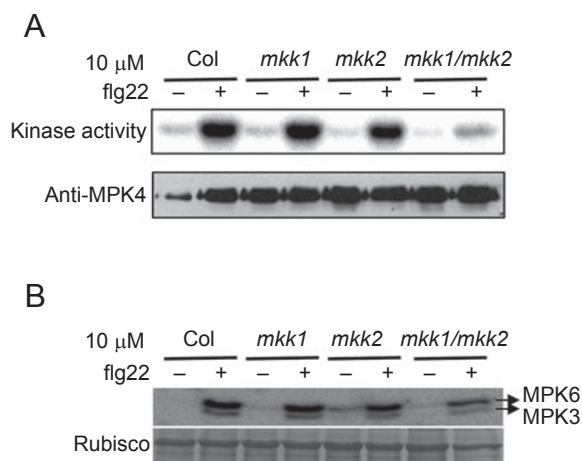


Figure 7 MAPK kinase activities in wild-type, *mkk1*, *mkk2* and *mkk1 mkk2* double mutants. **(A)** Analysis of *MPK4* activity upon *flg22* treatment by immunocomplex kinase assay. Top panel: *MPK4* kinase activity. Bottom panel: *MPK4* protein levels detected by immunoblot analysis. **(B)** Analysis of *MPK3* and *MPK6* activity upon *flg22* treatment by in-gel kinase assay. In both **(A)** and **(B)**, 2-week-old seedlings were grown on 1/2 MS medium at 22 °C. Seedlings were treated with 10 μM *flg22* for 15 min and subsequently collected in liquid nitrogen. Immunocomplex kinase assays were performed as described in the experimental procedures. In-gel kinase assays were performed as described previously [28].

knockout mutant plants. A recent report by Qiu *et al.* [18] found that *MPK4* activity was not affected in *mkk1* single mutant plants, which is consistent with our findings. As Meszaros *et al.* used 1 μM *flg22* while Qiu *et al.* used 10 μM *flg22* for induction, the difference in *MPK4* activation is probably because of the different concentrations of *flg22* used.

In addition to *MPK4*, *MPK3* and *MPK6* were activated upon *flg22* treatment [19]. To test whether *MKK1* and *MKK2* are required for the activation of *MPK3* and *MPK6*, we used an in-gel kinase assay with myelin basic protein as the imbedded kinase substrate to measure the activity of these two kinases in the wild-type and mutant plants. As shown in Figure 7B, activation of *MPK3* and *MPK6* is not affected in either the *mkk1* and *mkk2* single mutants or the *mkk1-1 mkk2-1* double mutant, suggesting that *MKK1* and *MKK2* are not required for the activation of *MPK3* and *MPK6* by *flg22*. In a recent report, Qiu *et al.* found that activation of *MPK3* and *MPK6* by *flg22* was compromised in the *mkk1* and *mkk2* single mutant and double mutant plants. We used in-gel kinase assays, whereas Qiu *et al.* performed kinase assays on immunoprecipitated *MPK3* and *MPK6*. The difference in *MPK3* and *MPK6* activities observed in these two studies might

be because of the different assays used.

Discussion

Arabidopsis MEKK1 is a negative regulator of cell death and defense responses. Knockout mutants of *MEKK1* displayed a seedling-lethality phenotype [10–12]. MPK4 has been suggested to be the MAP kinase downstream of MEKK1 because activation of MPK4 by flg22 or H₂O₂ is impaired in the *mekk1* mutants [10–12]. However, the previously reported *mpk4-1* mutant plants do not exhibit cell death and seedling-lethality phenotypes that were observed in the *mekk1* mutants [9]. Whereas *mekk1* mutants are in the Columbia background, *mpk4-1* is in the Landsberg background. The phenotypic difference between *mpk4-1* and *mekk1* mutants could be because of the genetic background difference. Here we report the identification of two new alleles of *mpk4* (*mpk4-3* and *mpk4-4*) in the Columbia background. Like the *mekk1* mutants, *mpk4-3* and *mpk4-4* exhibit spontaneous cell death and seedling-lethality phenotypes, suggesting that MPK4 and MEKK1 may function in the same MAPK kinase cascade.

MAP kinase cascades consist of three sequentially activated kinases, a MAP kinase kinase kinase (MAPKKK), a MAP kinase kinase (MAPKK) and a MAP kinase [20]. Our data suggest that MKK1 and MKK2 are the MAPKKs that function together with MEKK1 and MPK4. MKK1 and MKK2 interact with both MEKK1 and MPK4 *in vivo*, and *mkk1 mkk2* double knockout mutants display similar phenotypes as the *mekk1* and *mpk4* mutants. Furthermore, activation of MPK4 by flg22 is severely compromised in *mkk1 mkk2* double mutant plants, suggesting that MEKK1, MKK1/MKK2 and MPK4 form a MAP kinase cascade. Similar results were also reported by Qiu *et al.* [18]. As MPK4 is required for normal plant growth in the absence of external signals, one of the functions of the MEKK1, MKK1/MKK2 and MPK4 kinase cascade is most likely to perceive an endogenous plant signal for suppressing spontaneous cell death and defense activation.

Interestingly, interaction between MEKK1 and MKK1 or MEKK1 and MKK2 occurs mainly on the plasma membrane, whereas interaction between MPK4 and MKK1 or MPK4 and MKK2 occurs both on the plasma membrane and in the nucleus, suggesting that perception of the upstream signal occurs initially at the plasma membrane by MEKK1 and is subsequently transduced to the nucleus by the downstream kinases MKK1/MKK2 and MPK4.

One question is how disruption of the MEKK1, MKK1/MKK2 and MPK4 cascade leads to cell death and

activation of defense responses. As *mekk1*, *mkk1 mkk2* and *mpk4* mutants all accumulate high levels of H₂O₂, and H₂O₂ is a known inducer of cell death in plants [21], over-accumulation of H₂O₂ is probably the cause of cell death in these mutants. H₂O₂ is usually produced when plants are under stress conditions. It is important to keep H₂O₂ levels under control to avoid its detrimental effects on plants. The MEKK1, MKK1/MKK2 and MPK4 cascade could be a major mechanism for down-regulating H₂O₂ levels in plants. Disruptions of this MAPK kinase cascade lead to over-accumulation of H₂O₂ and, subsequently, cell death. Downstream of the MAPK kinase cascade, MPK4 has been shown to interact with MKS and phosphorylate two MKS-associated WRKY transcription factors, WRKY25 and WRKY33, *in vitro* [22]. It remains to be determined whether WRKY25 and WRKY33 also function in the regulation of H₂O₂ levels.

The cell death and constitutive activation of defense responses in the *mekk1*, *mpk4* and *mkk1 mkk2* double mutants are very similar to those of the mutant phenotypes caused by auto-activation of R proteins. Many R protein-mediated defense responses are also suppressed at high temperatures, suggesting a potential link between the MEKK1, MKK1/MKK2 and MPK4 kinase cascade and R protein-dependent defense pathways. However, it is unclear whether the high H₂O₂ levels in the *mekk1*, *mpk4* and *mkk1 mkk2* mutants are the cause or result of activation of R protein-mediated signaling pathways.

Arabidopsis MPK4 is activated under a variety of biotic and abiotic stress conditions [15, 23], probably through the MEKK1, MKK1/MKK2 and MPK4 kinase cascade. How different stresses activate the same downstream MAPK kinase cascade is unclear. One possibility is that a common signal is generated under different stress conditions. The signal is perceived by an unknown receptor upstream of MEKK1, which subsequently activates the MEKK1, MKK1/MKK2 and MPK4 kinase cascade. As H₂O₂ is produced under various stress conditions and it can activate MPK4 by itself, it is plausible to hypothesize that H₂O₂ serves as a general stress signal upstream of the MEKK1, MKK1/MKK2 and MPK4 kinase cascade in a negative feedback loop to down-regulate the H₂O₂ level. The mechanisms by which H₂O₂ activates its downstream cascade are yet to be determined by future studies.

Material and Methods

Mutant screen and mapping

To identify mutants that constitutively express *PR-2* (*BGL2*), *npr1-1* plants that contain the *pBGL2-GUS* reporter gene were mutagenized with EMS. The progeny of individual M1 lines were grown on MS medium and GUS staining was performed on half

of the seedlings from each line when they were 2 weeks old. Lines displaying constitutive GUS staining were recorded and the remaining half of the seedlings from these lines were transplanted to soil to set seeds. To identify mutants that are resistant to *H. p. Noco2*, the progeny of individual M1 lines were planted in soil. Two-week-old seedlings were sprayed with *H. p. Noco2* at a concentration of 50 000 spores per ml and scored 7 days after inoculation. Mutants that displayed seedling-lethality phenotypes were propagated with plants that are heterozygous for the mutation.

To map the *mpk4* and *mekk1* mutations in the Col background, the mutants were crossed with wild type Ler plants. The F2 progeny homozygous for the *mpk4* and *mekk1* mutations were identified by their morphology. The locations of the mutations were determined using In-del markers on different chromosomes.

BiFC analysis

The *MEKK1*, *MKK1*, *MKK2* and *MPK4* cDNA was amplified by PCR from Col total cDNA. The PCR products of *MEKK1* and *MPK4* were cloned into pUC-SPYCE that contained the 35S promoter and the C-terminal region of YFP to obtain pMEKK1-YCE and pMPK4-YCE. The PCR products of *MKK1* and *MKK2* were cloned into pUC-SPYNE that contained the 35S promoter and N-terminal region of YFP to obtain pMKK1-YNE and pMKK2-YNE, respectively. All constructs were confirmed by sequencing. Transient expression using *Arabidopsis* mesophyll protoplasts was performed according to a procedure described previously [24].

Mutant characterization

Plants were grown at 22 °C in 16 h light/8 h darkness on soil or 1/2 MS medium. *mekk1-1* was kindly provided by Dr Bögre [17]. *mkk1-2* (SALK_027645) and *mkk2-1* (SAIL_511_H01) were obtained from the ABRC. *mkk1-1* and *mkk1-2* were crossed with *mkk2-1* to generate the *mkk1 mkk2* double mutants. Expression of *PR* genes was analyzed as described previously [25]. Trypan blue and DAB staining were performed on 16-day-old seedlings grown on 1/2 MS medium according to procedures described previously [26, 27].

For the *H. p. Noco2* infection experiments, plants were kept at 18 °C in 12 h light/12 h dark cycles with 95% humidity after being sprayed with the spores of *H. p. Noco2* at a concentration of 50 000 spores per ml water.

Kinase assays

For MPK3, MPK4 and MPK6 kinase assay, 2-week-old seedlings grown on 1/2 MS medium were treated with 10 μM flg22 for 15 min and subsequently collected in liquid nitrogen. The plant tissues were ground and homogenized in extraction buffer containing 50 mM HEPES-KOH, pH 7.4, 50 mM NaCl, 10 mM EDTA, 5 mM NaF, 1 mM Na₃VO₄, 0.1% Triton-X 100, 1 mM DTT, 1 mM PMSF and 1× protease inhibitor cocktail (Sigma). After centrifugation at 14 000 rpm for 15 min at 4 °C, the protein concentration was determined using a Bio-Rad Bradford protein assay kit. An equal amount of total protein for each genotype was used in subsequent assay. For immuno-complex kinase assays, protein extracts were incubated with polyclonal antibodies of MPK4 (Sigma) for 1 h at 4 °C, and 20 μl protein A-agarose (GE) was subsequently added and incubated for another 3 h at 4 °C. The kinase on the beads was collected by centrifugation and washed thrice with extraction buffer and once with kinase buffer (50 mM HEPES-KOH, pH 7.4, 1

mM DTT, 10 mM MgCl₂, 10 mM MnCl₂ and 10 μM ATP). Kinase assays on the immunoprecipitated MPK4 were performed at room temperature for 30 min in 15 μl of kinase buffer containing myelin basic protein (MBP) (Upstate) as the artificial substrate and 2.5 μCi of [γ -³²P]ATP. Reactions were ended by the addition of SDS-PAGE loading buffer. Phosphorylation of MBP was analyzed by autoradiography after SDS-PAGE. The in-gel kinase activity assay was performed according to a procedure described previously [28] using 7 μg of total protein.

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