

Supplemental Informations

PRR2, a pseudo-response regulator, promotes salicylic acid and camalexin accumulation during plant immunity

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Legends of Supplemental Informations

Supplemental Information 1

Molecular characterization of *prp2* mutant lines. *The prp2* mutants were identified in the *Arabidopsis* T-DNA insertion lines collections from plant resource centers. **(A)** Schematic illustration of the T-DNA insertions into the genomic region of *PRR2* gene. The *prp2-1* (Col) and *prp2-2* (WS) mutants contain a T-DNA insertion in the first intron, 629bp and 425bp downstream of the transcription start site of *PRR2* gene respectively. **(B)** Semi-quantitative analysis of *PRR2* transcripts accumulation was carried out on homozygous lines using primers *F1* and *R1* (see panel A for primers positions). Actin 2 was used as a control. Data indicate that *PRR2* transcripts can be detected in both insertional mutants, but they accumulate at a lower level than that observed in wild-types, indicating a down-regulation of *PRR2* expression in the mutants. **(C)** Quantitative RT-qPCR was performed to measure the relative *PRR2* gene expression level in mutants compared to WT. As shown, *PRR2* expression is respectively decreased by about 3 and 7 fold in *prp2-1* and *prp2-2* compared to Col and WS.

Supplemental Information 2

Molecular characterization of Over-Expressing *PRR2* transgenic lines. **(A)** Relative expression of *PRR2-3HA* transcript in *p35S::CDSPRR2.1* (*OE-PRR2.1*) and *p35S::CDSPRR2.2* (*OE-PRR2.2*) lines compared to WT (Col) by quantitative RT-PCR. **(B)** Western blot of total protein from WT (Col), *OE-PRR2.1* and *OE-PRR2.2*. Total proteins were extracted and separated in SDS-PAGE before blotting and

immunodetection using antibodies raised against the 3HA epitope. Presence and length of the 3HA-tagged PRR2 protein is indicated by the arrow.

Supplemental Information 3

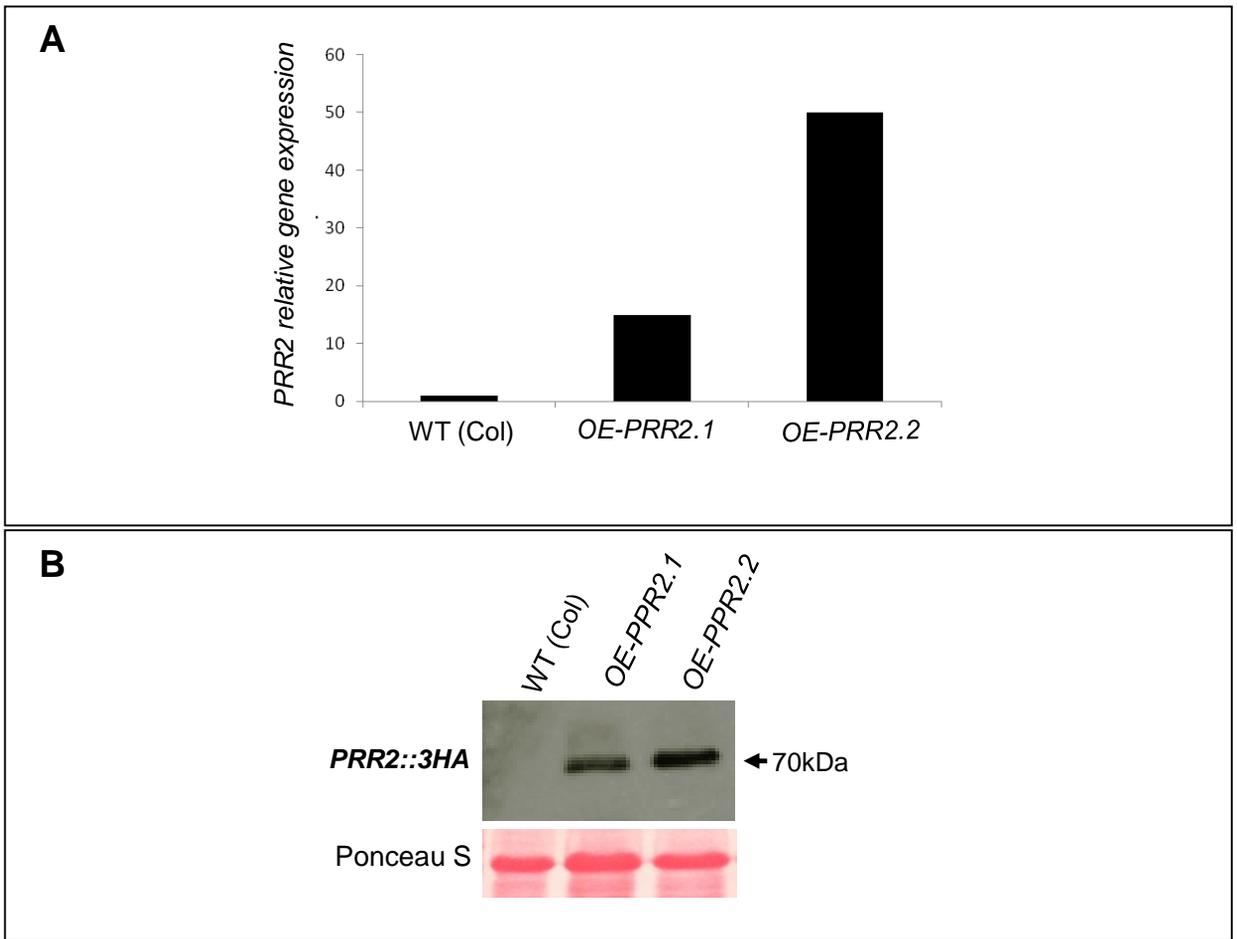
(A) Phenotype of 6-day-old WT (Col), *OE-PRR2.1*, *OE-PRR2.2*, *prp2.1* and WT (WS) and *prp2.2* plants grown in standard conditions. All plants were photographed from the same distance. (B) Phenotype of 4-week-old WT (Col0), *prp2.1*, *OE-PRR2.1*, *OE-PRR2.2* and WT (WS) and *prp2.2* plants grown in standard conditions. All plants were photographed from the same distance.

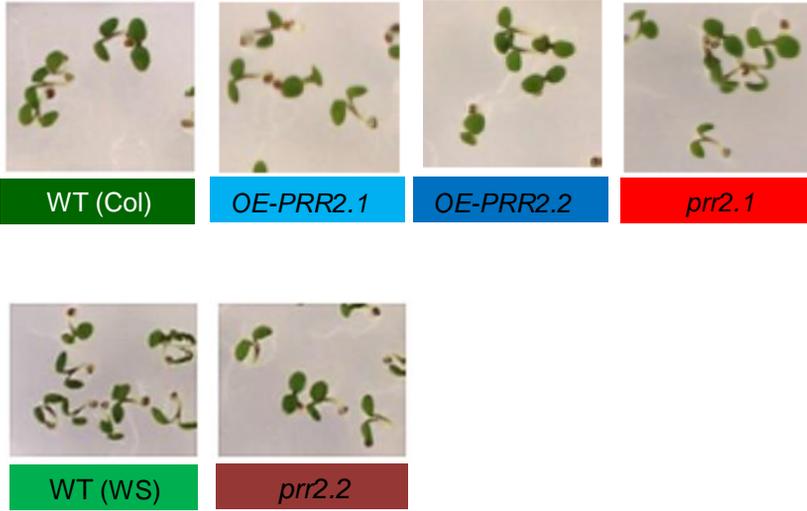
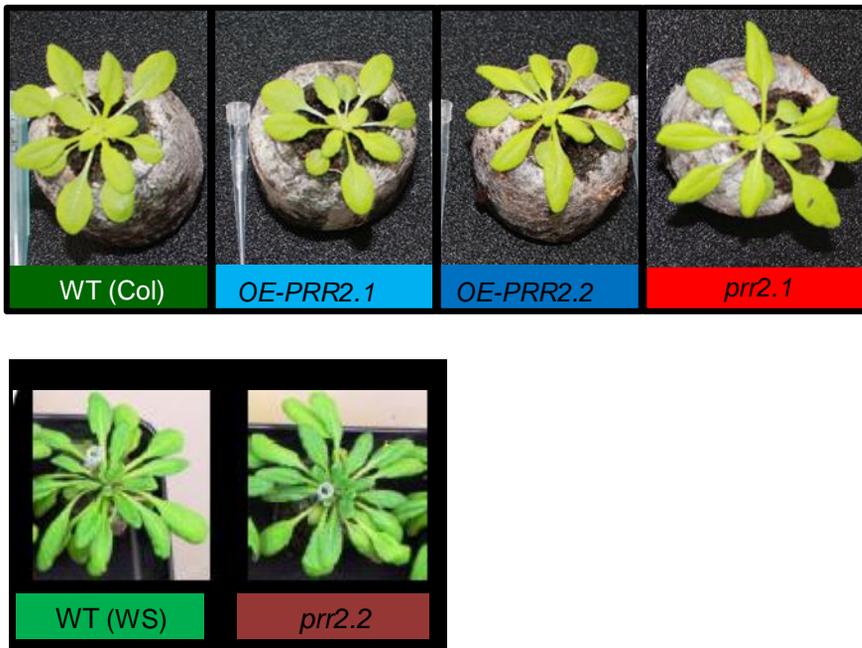
Supplemental Information 4

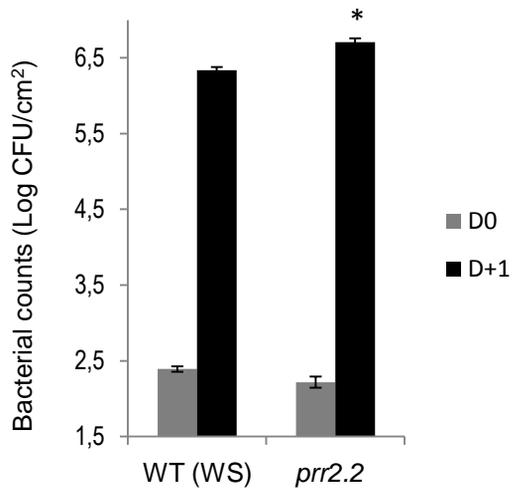
Quantifications of *in planta* bacterial growth in WT (WS) and *prp2.2* mutant line. Bacterial growth quantifications were performed at 0 and 1 dpi with *Pst* DC3000 (A) or *Pseudomonas syringae pv maculicola* (B). Data are representative of 5 replicates of three independent experiments. P values were calculated using the two-tailed Mann-Whitney U-test to indicate significant differences between the genotypes. Asterisks illustrate $p < 0.05$. Error bars represent SEM.

Table S1: List of primers used for quantitative PCR and molecular characterization of the *prp2* genotypes.

Name	Sequence	
<i>ACTIN8</i>	5'-CACCCGAGAGGAAGTACAGTG-3'	Forward
	5'-CATACTCTGCCTTAGAGATCCACA-3'	Reverse
<i>PRR2</i>	5'- GAGCGATTCCACCTTTGTT-3'	Forward
	5'- CAACCCCATGCATTACCG -3'	Reverse
<i>WRKY6</i>	5'- GCAACAGCAACAACAGAACAA -3'	Forward
	5'- TGCCTTGGTACTATCGTCTCC-3'	Reverse
<i>MYB51</i>	5'- GGCCAATTATCTTAGACCTGACA -3'	Forward
	5'- CCACGAGCTATAGCAGACCATT -3'	Reverse
<i>CYP71B15 / PAD3</i>	5'- CACCACTGATCATCTCAAAGGA -3'	Forward
	5'- CGGTCATTCCCCATAGTGTT -3'	Reverse
<i>CBP60G</i>	5'- ATCGCAGCACATCGACTTT -3'	Forward
	5'- GTGGACCGTTGAGCTTGAA -3'	Reverse
<i>PromPRR2</i>	5'- CATACGGTTTTTATAAGTAATGAAGC - 3'	Forward
	5'- CATAGCTGGTTTTGCATCGTTTTTG -3'	Reverse



A**B**

A *Pseudomonas syringae* pv *tomato* (DC3000)**B** *Pseudomonas syringae* pv *maculicola*