

ARTICLE ADDENDUM

## Getting to PTI of bacterial RNAs: Triggering plant innate immunity by extracellular RNAs from bacteria

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### ABSTRACT

Defense against diverse biotic and abiotic stresses requires the plant to distinguish between self and non-self signaling molecules. Pathogen/microbe-associated molecular patterns (PAMPs/MAMPs) are pivotal for triggering innate immunity in plants. Unlike in animals and humans, the precise roles of nucleic acids in plant innate immunity are unclear. We therefore investigated the effects of infiltration of total *Pseudomonas syringae* pv. tomato DC3000 (*Pto* DC3000) RNAs into *Arabidopsis* plants. The pathogen population was 10-fold lower in bacterial RNAs pre-treated *Arabidopsis* plants than in the control. Bacterial RNAs purity was confirmed by physical (sonication) and chemical (RNase A and proteinase K digestion) methods. The perception of bacterial RNAs, especially rRNAs, positively regulated mitogen-activated protein kinase (MAPK) and induced a reactive oxygen species burst, callose deposition, salicylic acid (SA) and jasmonic acid (JA) signaling, and defense-related genes. Therefore, bacterial RNAs function as a new MAMP that activates plant innate immunity, providing a new paradigm for plant-microbe interactions.

**Abbreviations:** ET, ethylene; ETI, effector-triggered immunity; JA, jasmonic acid; MAPK, mitogen-activated protein kinase; PRR, pattern recognition receptor; PTI, PAMP-triggered immunity; SA, salicylic acid

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Unlike in vertebrates, a circulating immune system has not been developed in plants. Plants are continuously subjected to invasion by pathogens and insects. Therefore, to overcome these attacks, plants must produce highly effective and specialized immune responses.<sup>1-4</sup> The onset of innate immunity in plants is first activated by the recognition of non-self components.<sup>5</sup> Foreign non-self signals or molecular patterns are perceived by pattern recognition receptors (PRRs) in plants.<sup>6</sup> Two major innate immune systems, pathogen/microbe-associated molecular patterns (PAMPs/MAMPs)-triggered immunity (PTI) and effector-triggered immunity (ETI), are widely known in plants.<sup>7</sup>

Flagellin, peptidoglycan, lipopolysaccharide (LPS), translation elongation factor-Tu (EF-Tu), cold-shock protein, and fungal chitin are key molecular patterns that activate plant innate immunity.<sup>8-10</sup> In animals, nucleic acids play a role in eliciting innate immunity. However, little is known about the effects of nucleic acids on innate immunity in plants. To shed light on the functions of nucleic acids in this process in plants, we infiltrated total bacterial RNAs from *Pseudomonas syringae* pv. tomato DC3000 (*Pto* DC3000) into *Arabidopsis thaliana* leaves and subsequently inoculated the same plants with the same bacterium.<sup>11</sup> Notably, pre-infiltration of bacterial RNAs into plants resulted in reduced pathogen population density and activated downstream signaling pathways, as triggered by the typical PTI, in addition to increasing superoxide anion

production and callose deposition and positively regulating mitogen-activated protein kinase (MAPK) and defense-related gene expression.<sup>11</sup>

Intriguingly, plant innate immunity was elicited by total bacterial RNAs but not by *Arabidopsis* RNA, suggesting that the host plant can recognize bacterial RNAs as a “non-self” signaling molecule.<sup>11</sup> However, this suggestion can be countered by the fact that bacterial RNAs are structurally similar to plant ATP.<sup>12</sup> Perhaps when bacterial RNAs are placed into the apoplast of *Arabidopsis*, they are hydrolyzed and their degradation elicits the plant ATP-type defense response. However, this explanation may be inaccurate because sheared bacterial RNAs did not modulate plant innate immunity.<sup>11</sup> This result suggests that certain structural features of bacterial RNAs are the major determinants of plant innate immunity.

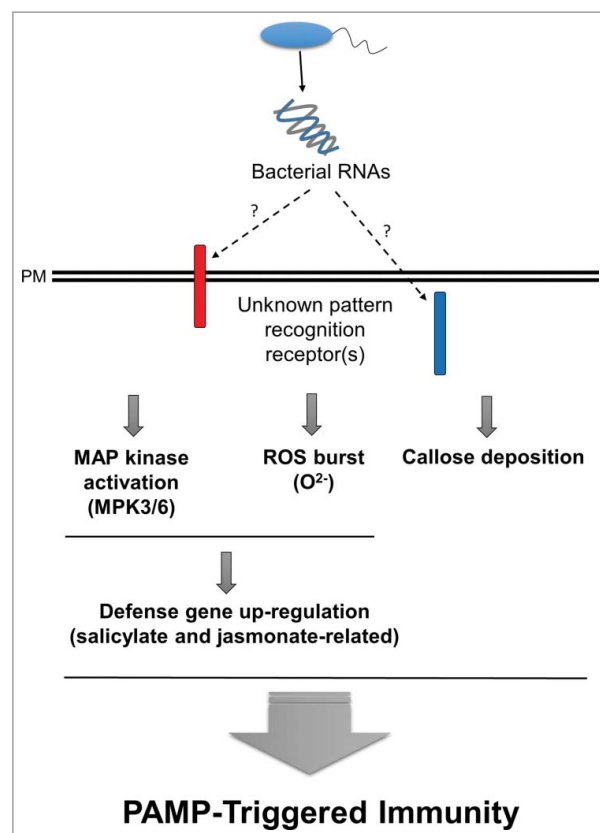
The major bottleneck of this study was that any contamination of the bacterial RNAs had to be eliminated. To demonstrate the purity of the bacterial RNAs, we used both physical and chemical approaches. When bacterial RNAs are sheared by sonication, they become fragmented and degraded, and any contaminating peptide(s)/proteins remain in the sheared bacterial RNAs sample. Despite the possible existence of peptide(s)/proteins in sheared bacterial RNAs, no plant innate immunity was elicited by this RNA sample, suggesting that intact bacterial RNAs are needed to activate plant innate immunity. To support the function of bacterial RNAs, purified bacterial RNAs were

treated with RNase A before infiltration into *Arabidopsis* leaves. The pathogen population was 10-fold higher in leaves treated with RNase A + bacterial RNAs than in those treated with bacterial RNAs alone.<sup>11</sup> In addition, 5-fold more pathogen colonies were observed in leaves treated with bacterial RNAs + proteinase K than in those treated with bacterial RNAs alone.<sup>11</sup> These results clearly demonstrate that bacterial RNAs are indeed implicated in the modulation of innate immunity in *Arabidopsis*.

Although the pathogen level was higher in leaves treated with RNase A + bacterial RNAs versus RNAs alone, this level failed to reach that of the control plants. Moreover, we observed a difference in pathogen population density between plants treated with bacterial RNAs alone and those treated with bacterial RNAs + proteinase K.<sup>11</sup> It is possible that the activity of RNase A and proteinase K was not completely diminished during the pre-incubation process and that the residual activity affected the pathogen population density. The levels of the pathogen population in plants treated with RNase A alone vs. the water control were significantly different, whereas the same level was detected in plants treated with proteinase K alone versus the water control.<sup>11</sup> This finding, along with the observations that ribonucleases modulate plant defense responses<sup>12</sup> and that protease inhibitors widely participate in improving plant defense responses to pathogens and insects,<sup>13</sup> suggests that protease may negatively affect plant innate immunity. Based on our results, RNase A and proteinase K may be involved in innate immunity in *Arabidopsis*.

To investigate whether plant defense hormones function in plant innate immunity triggered by bacterial RNAs, we used representative *Arabidopsis* mutant lines for salicylic acid (SA), jasmonic acid (JA), and ethylene (ET) signaling, finding that SA and JA signaling pathways are required for bacterial RNAs-mediated *Arabidopsis* innate immunity.<sup>11</sup> A result of our study concerning plant defense signaling pathways differs from that of a previous study. In contrast to *flg22*- and *elf18*-induced resistance, we found that bacterial RNAs-induced resistance to *Pto* DC3000 was completely abolished in *npr1*, *NahG*, and *jar1-1* mutant plants. The *Pto* DC3000 population was only significantly reduced when the quadruple mutant *dde2 ein2 pad4 sid2* was used, while no changes were observed in single mutants.<sup>14</sup> This difference may be explained as follows: 1) Different PTI-enhancing components were used in the experiments (i.e., *flg22* or *elf18* vs. bacterial RNAs); 2) The pathogen incubation time after pre-treatment with bacterial RNAs was 4 times longer and the pathogen (*Pto* DC3000) concentration was approximately 100 times higher in our study than in the previous study; and 3) In our study, the *Pto* DC3000 population was assessed in systemic leaves rather than local leaves. Collectively, these parameters led to significant differences between our results and those of the previous study, which is not surprising.

In conclusion, our study revealed that pre-infiltrating *Arabidopsis* with bacterial RNAs positively modulated innate immunity in response to *Pto* DC3000. Despite the strong ability of bacterial RNAs to function as an elicitor of plant innate immunity, several unanswered questions remain: How are bacterial RNAs perceived by specific plant



**Figure 1.** A schematic representation of plant innate immunity elicited by bacterial RNAs as a pathogen-associated molecular pattern (PAMP) in plant. Bacterial RNAs may be perceived by either intercellular or intracellular plant pattern recognition receptor(s) (red or blue bar, respectively). Inside the plant cell, bacterial RNAs affect the downstream pathways. Activation of the MAP kinase signaling cascade, reactive oxygen species (ROS) including superoxide anion burst, callose deposition, and defense-related gene expression levels results in elicitation of PAMP-triggered immunity. PM = plasma membrane.

receptor(s)? How can bacterial RNAs survive in the plant apoplast under acidic condition? Our intriguing findings are listed in Fig. 1. Based on the results, the well-known PRRs of chitin, flagella, and EF-Tu do not appear to be receptors of bacterial RNAs. Inside the plant cell, bacterial RNAs positively regulate the MAP kinase signaling cascade, as well as superoxide anion levels, callose deposition, SA and JA signaling pathways, and the associated downstream defense-related gene expression levels. The discovery of the role of bacterial RNAs as an elicitor of plant innate immunity provides new insights into this important process. To better understand the underlying mechanisms by which bacterial RNAs play a major role in the regulation of plant innate immunity, further investigations are needed.

### Disclosure of potential conflicts of interest

No potential conflicts of interest are disclosed.

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