Bacterial effectors target BAK1-associated receptor complexes
One stone two birds

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Abbreviations: BAK1, BRI1-associated kinase; PAMP, pathogen-associated molecular pattern; PRR, pattern recognition receptor; TTSS, type III secretion system; ETI, effector-triggered immunity

The long-standing association between hosts and microbes has generated some of most intricate relationships. The studies on molecular mechanisms of host-microbe interaction have been revealing many fascinating stories. Here we zoom in on a specific topic on the interplay between bacterial effectors and plant innate immune signaling. In particular, we will summarize our recent discovery that bacterial effector proteins, AvrPto and AvrPtoB, target plant immune signaling receptor complexes to interfere with host immune responses and development.

PAMP-Triggered Immunity; The Front Line of Host Defense Responses

Disease is the exception rather than the rule for most of creatures despite that plants and animals are constantly surrounded by a diverse array of potential pathogens. Through evolution, hosts have developed the capacity to timely detect and mount effective defense responses to dampen potential infections. As an active and first line of defense, innate immunity plays pivotal roles in fending against the onslaught of outside invaders that penetrate or evade the physical barriers (such as skin and waxy epidermal cells). The invading microbes are in part detected through perception of some ‘general’ microbe-derived molecules, which were coined as pathogen-associated molecular patterns (PAMPs) or microbe-associated molecular patterns (MAMPs), by host cell-surface pattern-recognition receptors (PRRs), leading to PAMP-triggered immunity (PTI).1,2 Most identified plant PRRs are receptor-like kinases (RLKs).

Two well-characterized Arabidopsis PAMP receptors are FLAGELLIN-SENSING 2 (FLS2) that perceives a conserved 22-amino acid peptide (flg22) from bacterial flagellin, and EF-TU RECEPTOR (EFR) that recognizes bacterial elongation factor EF-Tu.3,4 Affinity cross-linking assays demonstrated direct binding of flagellin to FLS2 and EF-Tu to EFR, suggesting that FLS2 and EFR are bona fide PAMP receptors.5,5 CERK1, a RLK with three LysM motifs, is involved in fungal chitin perception and signaling in Arabidopsis.6,7 However, there is no direct evidence to support that CERK1 binds to chitin. Instead, a plasma membrane glycoprotein (CEBiP) has been shown to directly bind to chitin in rice.8 CEBiP also contains two extracellular LysM motifs. Recently, it was found that the biogenesis of certain PRRs in plant innate immunity requires the endoplasmic reticulum-quality control (ER-QC) machinery. EFR, but not FLS2, cannot accumulate in the null mutant of Calreticulin 3 gene, which encodes an ER chaperon involving the proper folding of newly synthesized glycoproteins.9 It is intriguing how the ER-QC components are specifically required for the proper accumulation of a subset of PRRs as FLS2 and EFR belong to the same family of RLKs and induce largely overlapping responses.

The intracellular signaling events downstream of PRR perception include the activation of evolutionarily conserved mitogen-activated protein kinase (MAPK) signaling cascades, transcriptional changes, and the production of antimicrobial compounds.10,11 As an early signaling event upon PAMP binding to corresponding PRR, receptor dimerization plays an essential role in triggering intracellular signaling. In plants, FLS2 rapidly dimerizes with BAK1, another RLK, upon flagellin treatment. BAK1 was originally identified as a partner of BRI1 mediating BR signaling.12-15 Interestingly, ER-QC machinery has also been implicated in the biogenesis of BRI1.16 However, BAK1 is not involved in BR binding to BRI1 and flagellin binding to FLS2.14,17 Intriguingly, BAK1 is required for multiple PAMP responses, including flagellin, EF-Tu, lipopolysachride (LPS), peptidoglycan (PGN), bacterial cold-shock protein and oomycete elicitor INF1 in Arabidopsis and Nicotiana benthamiana.14,15 Thus, BAK1 appears to function in distinct receptor signaling complexes to integrate multiple PAMP perception into downstream signaling events. In addition, BAK1 is also required to constrain the spread of cell death induced by virulent bacteria Pseudomonas syringae pv. tomato DC3000.18

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Through its interaction capacity with Pto in a yeast two-hybrid screen.\(^2\) The apparent anomaly that pathogens possess an immune elicitor was satisfactorily explained by a careful observation that AvrPto moderately promoted pathogen growth on the plants lacking Pto or Prf.\(^3\) An outstanding question was how these bacterial proteins contribute to the pathogenesis at the molecular level. The observation that type III effectors collectively suppress PTI led us to engage a versatile cell-based system to screen for individual effectors contributing to the suppression activities on susceptible hosts. AvrPto and AvrPtoB were identified from the screen as suppressors of multiple PAMP-mediated immune responses upstream of MAPK cascade at the plasma membrane. It was proposed that AvrPto and AvrPtoB may target PAMP receptors or other convergent early signaling components to suppress immunity triggered by multiple PAMPs.\(^3\)

An interesting observation that transgenic plants overexpressing AvrPto resemble BR-deficient mutant led to the discovery that AvrPto and AvrPtoB directly target the signaling partner BAK1 involved in multiple PAMP-triggered immune responses and plant hormone BR-mediated signaling.\(^3\) AvrPto interacts with BAK1 in a yeast split-ubiquitin assay designed for detecting membrane protein interaction, and AvrPto and AvrPtoB associate with BAK1 in vivo by co-immunoprecipitation assay. Importantly, extensive mutagenesis and deletion analyses suggest the biological importance of BAK1-AvrPto/AvrPtoB interaction.\(^3\) This targeting leads to the dissociation of flagellin-induced FLS2-BAK1 association, and brassinosteroid-mediated BRI1-BAK1 association, thereby blocking the initiation of flagellin and BR signaling (Fig. 1). It is important to note that AvrPto and AvrPtoB do not interact with BRI1. Thus, suppression of BR signaling by AvrPto does not involve the interaction with BR receptor BRI1. Given the fact that BAK1 is involved in signaling triggered by BR, multiple PAMPs and cell death, by targeting BAK1, AvrPto and AvrPtoB efficiently suppress multiple signaling in plant immunity and development, one stone killing two birds (Fig. 2). Apparently, AvrPto also targets BAK1-independent PAMP signaling since BAK1 is essential in some, but not all PAMP-signaling responses, whereas AvrPto is equally effective in suppressing the immune responses triggered by all PAMPs tested (Fig. 2).

**Conclusions and Challenges**

Great strides are being made in understanding the biological and enzymatic functions of effectors in the past few years with the development of new technologies and more sensitive and quantitative assays. The newly identified effector targets in hosts have been revealed to be often novel and essential

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**Figure 1.** Bacterial effectors suppress PAMP immunity by targeting receptor signaling complexes. (A) In the absence of infection, PAMP receptor FLS2 remains at the resting stage and there is no apparent receptor complex formation. (B) Once challenged by microbes, FLS2 detects the invading microbes by binding to the ligand-bacterial flagellin, which allows FLS2 conformational change and heterodimerization with BAK1 to form a tight receptor complex. The functional PAMP receptor complex initiates downstream signaling, including activation of MAPK cascade, transcription factors and immune genes, thereby leading to plant resistance to microbes. (C) To be pathogenic, some successful pathogens deployed virulence weapons to tear down host immune system. For instance, type III effectors AvrPto and AvrPtoB, secreted from *Pst* DC3000 and delivered into host cells, target BAK1-associated PAMP receptor complexes and disrupt the complex formation, thereby impeding the immune signaling and leading to disease.

**Bacterial Effectors Dampen Host Immunity and More: Breach of Peace**

To be pathogenic, many Gram-negative bacteria have evolved the ability to inject a repertoire of virulence effector proteins into host cells through type III secretion system (TTSS) to modulate diverse host cellular activities and physiology.\(^9\),\(^10\) Although the structural components of TTSS are relatively conserved, the sequences and functions of individual effectors secreted from TTSS can be highly divergent in different species of plant and animal pathogenic bacteria. Some species, such as *Yersinia* of animal pathogens, secrete just a few effectors, whereas *Pseudomonas* species of plant pathogens deliver about thirty effectors.\(^2\),\(^2\) Now, it is certain that effectors collectively determine bacterial pathogenicity by interfering with host defense responses, thus leading to effort-triggered susceptibility (ETS) on susceptible plants. However the contribution of individual effectors to virulence is trivial to observe in plants likely due to the potential functional redundancy and the limitation of diagnostic tools.\(^2\),\(^3\) In some cases, these effectors could be recognized directly or indirectly by plant disease resistance proteins (R proteins) to induce potent effector-triggered immunity (ETI) and serve as so-called avirulence (Avr) factors that turn virulent strains into avirulent ones.\(^2\),\(^9\)\(^-\)\(^1\)

AvrPto was historically classified as an avirulence protein based on its ability of eliciting substantial immune responses in tomato plants carrying the immune sensors Pto and Prf.\(^3\)-\(^6\)AvrPtoB, a sequence distinct effector composing of at least two functional domains (the N terminus responsible for interaction with Pto, and the C terminal E3 ligase domain) was initially identified...
components in innate immunity. For instance, tandem mass spectrometry analysis revealed that OsP from Shigella encodes a novel enzyme, phosphothreonine lyase, in the suppression of MAPK activity.\(^{39}\) HopAI, an OsP family member in plant pathogen \textit{P. syringae}, also dephosphorylates MAPKs probably through the same biochemical activity.\(^{40}\) \textit{P. syringae} effector HopU1, a mono-ADP-ribosyltransferase (ADP-RT), modifies several Arabidopsis RNA-binding proteins, which represent a novel class of ADP-RT substrates. Importantly, the Arabidopsis mutant lacking one of these RNA binding proteins is immunocompromised to pathogen infection, suggesting an essential role of this protein in innate immunity.\(^{41}\) HopM1, another \textit{P. syringae} effector, targets and degrades Arabidopsis MIN7 protein, a member of the ARF family of guanine nucleotide exchange factors involved in vesicle trafficking, which was also found to be an important player in plant innate immunity.\(^{42}\) Currently, profound knowledge on mechanistic bases of effector hijacking host cellular signaling is mostly derived from overexpression of individual effector genes in host cells. In nature, minute amount of effectors is likely delivered into hosts by pathogens. Thus, the interpretation of data needs to be cautious with the particular system deployed. Another complication is whether the secretion of effectors into hosts possesses the host specificity and secretion hierarchy among the repertoire of effectors on different hosts. Multiple dynamic host targets of effectors have also been proven to commonly exist.\(^{43}\) With the rapid development of innovative technologies and in-depth understanding of host cellular signaling pathways, it is brightly believed that biologists will continue to unravel the dynamic and intimate interaction between microbes and hosts.

**Figure 2.** Multiple functions of AvrPto and AvrPtoB as virulence and avirulence factors. BAK1 is a signaling partner of BR receptor BRI1, multiple PAMP receptors PRRs, and unknown receptor for cell death. By targeting BAK1, AvrPto and AvrPtoB efficiently suppress multiple signaling in plant immunity, development and cell death control, one stone killing two birds. Apparently, AvrPto also targets BAK1-independent PAMP signaling. In the presence of Pto and Prf, AvrPto triggers potent effector-mediated immunity in tomato.

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