Remorins: Essential Regulators in Plant-Microbe Interaction and Cell Death Induction

Programmed cell death (PCD) is a genetically controlled process triggered by developmental or environmental cues. Plant-microbe interactions often lead to PCD in plant host cells, triggered by the hypersensitive response. Plants may recognize microbes through receptor-like kinases in the plasma membrane or detect pathogen effectors through resistance (R) proteins in the cytoplasm. Both pathways activate immune responses and trigger PCD (Fig. 1).

In this issue of *Plant Physiology*, Cai et al. (2020) describe a new role for remorin (REM) in regulating PCD. REMs are plant-specific proteins localized on the inner side of the plasma membrane and plasmodesmata. REMs accumulate in detergent-insoluble fractions of the plasma membrane called lipid rafts (Raffaele et al., 2009). REMs have a highly conserved C terminus with a coiled-coil domain and a divergent N terminus, the sequence of which permits them to be classified into six groups (Raffaele et al., 2007). The roles of REMs in plant-microbe interaction have been demonstrated in various species (Fig. 1). Potato (*Solanum tuberosum*) and tobacco (*Nicotiana benthamiana*) group I REMs physically interact with virus movement proteins and inhibit virus movement (Raffaele et al., 2009; Fu et al., 2018). Arabidopsis (*Arabidopsis thaliana*) group IV REMs increase susceptibility to viruses, as the *atrem4.1 atrem4.2* double mutant is more resistant to geminivirus infection (Son et al., 2014). In legumes, orthologs in *Medicago truncatula* and *Lotus japonicus* of a group II REM are required for nodulation and interact with symbiotic receptor-like kinases (Lefebvre et al., 2010; Tóth et al., 2012).

Cai et al. (2020) report that transgenic tomato (*Solanum lycopersicum*) overexpressing a group I REM, SIREM1, was more susceptible to the necrotrophic fungus *Botrytis cinerea*. Additionally, *Agrobacterium tumefaciens*-mediated transient expression of SIREM1 in tobacco leaves induced PCD, as assayed by Trypan Blue staining and cell electrolyte leakage. Overexpression of other group I REMs, SIREM1.4, SIREM1.5, SIREM1.6, and SIREM1.7, but not SIREM1.2/SIREM1.3, in tobacco leaves resulted in similar symptoms, suggesting at least partially conserved roles for group I SIREMs in PCD induction. Consistent with previous reports that REMs form homooligomers and heterooligomers (Lefebvre et al., 2010; Tóth et al., 2012; Son et al., 2014; Fu et al., 2018), SIREM1 physically interacted with itself and SIREM1.4, SIREM1.5, SIREM1.6, and SIREM1.7. However, no interaction between SIREM1 and SIREM1.2/SIREM1.3 was detected.

Cai et al. (2020) further performed quantitative proteomics analysis using tobacco leaves with or without SIREM1 overexpression. In the proteins up-regulated by SIREM1 overexpression, the authors identified a set of proteins involved in PCD regulation, including the known plasma membrane NADPH oxidase NtRBOHB. Quantitative real-time PCR showed that higher abundance of NtRBOHB is attributable to a higher transcript level. Consistent with the molecular data, reactive oxygen species including hydrogen peroxide and superoxide were more abundant in leaves overexpressing SIREM1. In addition, the authors identified three novel cell death regulators through the proteomics analysis, including CYSTEINE-RICH AND TRANSMEMBRANE DOMAIN-CONTAINING PROTEIN A-LIKE, BLUE COPPER PROTEIN-LIKE, and NUCLEAR CAP-BINDING PROTEIN SUBUNIT 2. Transient expression of these proteins in tobacco leaves enhanced cell death, confirming their positive roles in PCD induction (Fig. 1).

To reveal the functional domains that are required for REM function, Cai et al. (2020) generated a series of mutated forms of SIREM1 and transiently expressed

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Figure 1. REMs participate in plant-microbe interaction and PCD. REMs polymerize and physically interact with receptor-like kinases (RLK) and pathogen effectors; the latter activate *R* proteins and trigger immunity. The C terminus of REM (REM-C) is required for plasma membrane (PM) localization and protein interaction, while phosphorylation occurs at the N terminus (REM-N). REMs induce reaction oxygen species (ROS) and up-regulate protein abundance of CYSTEINE-RICH AND TRANSMEMBRANE DOMAIN-CONTAINING PROTEIN A-LIKE (CRTD), BLUE COPPER PROTEIN-LIKE (BCPL), NUCLEAR CAP-BINDING PROTEIN SUBUNIT 2 (NCP2), and RBOHB in tobacco, which all contribute to programmed cell death. Whether REM induction of cell death is triggered by RLKs is not known.
them in tobacco leaves. They concluded that amino acids 173 to 187 at the C terminus are essential for the plasma membrane localization of SIREM1 and induction of cell death. Additionally, the 35 amino acids at the C terminus of SIREM1 were necessary for interaction with SIREM1.4 (Cai et al., 2020). These results are consistent with previous findings in other species, including StREM1.3 in potato (Perraki et al., 2012), LjSYMREM1 in L. japonicus (Tóth et al., 2012), and NbREM1 and NbREM4 in tobacco (Fu et al., 2018; Albers et al., 2019). Together, these results demonstrate that the conserved C terminus of REMs is responsible for plasma membrane localization and protein interactions.

Lastly, Cai et al. (2020) investigated the effect of phosphorylation on SIREM function. REMs were originally identified as phosphorylated proteins, and they are phosphorylated by various protein kinases in vitro (Tóth et al., 2012; Son et al., 2014; Albers et al., 2019). Cai et al. (2020) identified five phosphorylated sites of SIREM1 expressed in tobacco leaves using mass spectrometry. Interestingly, all phosphorylated sites were localized in the N terminus of the protein, which is consistent with earlier findings in LjSYMREM1 (Tóth et al., 2012) and NbREM4 (Albers et al., 2019). Both the phosphorylation-inactive and -active forms of SIREM1 exhibited decreased efficiency in PCD induction. The active form of SIREM1 also failed to interact with the native SIREM1 in a yeast two-hybrid assay, suggesting that these sites are necessary for SIREM1 function. It would be interesting to further dissect how phosphorylation of SIREM1 regulates the protein activity in plants.

Earlier studies suggest that REMs participate in upstream signaling during plant-microbe interaction, as they physically interact with receptor-like kinases and pathogen effectors. Cai et al. (2020) have provided evidence that REMs up-regulate PCD regulators at transcript and protein levels and induce PCD, a downstream defense mechanism. Further studies may reveal whether REMs directly participate in PCD induction by interacting with PCD signaling proteins or through the plant defense response.

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