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Title

Different threats, same response

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1 Plant immunity /631/449/2169 2 /631/449/2169/2673 Effectors in plant pathology 3 Pattern recognition receptors in plants /631/449/2169/2107 4 /631/61/212/2019 Transcriptomics 5 6 **Strapline:** Plant immunity 7 8 Title: Different threats, same response 9 10 Authors: Benjamin J. Cole and Susannah G. Tringe 11 12 Standfirst (37w, 250c): Plants can experience a range of challenges, from 13 osmotic stress to pathogen attack, requiring different types of responses. Despite 14 this variety, two recent studies of plant transcriptomes reveal a surprising 15 commonality in the genes induced by stress. 16 17 Main text (873 words): Plants cannot run away from their enemies. Nor can they 18 relocate when conditions change for the worse. Therefore plants have evolved an 19 elaborate system of defences to identify, respond, and adapt to stresses ranging 20 from limited nutrient availability to pathogen attack, and to distinguish them from 21 harmless stimuli like commensal microbes. In this issue of Nature Plants, two groups 22 from Switzerland explore these stress responses in greater molecular detail^{1,2}, 23 systematically characterizing changes in gene expression triggered by a wide variety of challenges in the model plant Arabidopsis thaliana. As a result they 24 25 unveiled features shared across diverse stress responses, and identified genes not 26 previously implicated in defence. 27 28 In the first study from Bjornson et al¹, a diverse set of molecules (Pathogen-, 29 Microbe-, or Damage-Associated Molecular Patterns, or P/M/DAMPs) that elicit a 30 strong, broad-range immune response (PTI for pattern-triggered immunity) were 31 applied to Arabidopsis plants. These molecules are derived from bacteria, fungi, 32 insects, or even plant tissue, and their perception by plant membrane receptors 33 (Pattern Recognition Receptors, or PRRs) signals an imminent danger: a foreign 34 organism is present or plant cellular damage has occured³. After application, the 35 authors measured global transcriptomic responses in short time scales (within 3 36 hours). Remarkably, they observed that the two largest clusters of differential expression were the genes modified by the most active elicitor only (flg22), and 37 genes modified by any of the 7 tested elicitors. It points to a large overlap and a 38 39 core of coordinated gene expression in response to pathogen perception. 40 41 Even more intriguing, when these gene expression phenotypes were compared to those induced by other environmental stresses (e.g. light, heat, salt), a large 42 43 overlap was observed, suggesting that a General Stress Response (GSR) is central

44 to P/M/DAMP-induced immunity, especially in the very early stages of exposure

45 (figure 1). Notably, impairment of the GSR also interfered with PTI, indicating a tight 46 coupling between these responses. Lastly, the study focused on genes that are strongly impacted by P/M/DAMP elicitors, but not affected by abiotic stress, termed 47 48 "Core Immunity Response", or CIR genes. Among these, they identified a set of 49 calcium channels (Glutamate Receptors, or GLRs) that, when mutated, dampen the 50 plant immune response to a bacterial pathogen. 51 52 Moving beyond elicitors, independent work by Maier and colleagues² brought a 53 diverse panel of live bacterial strains to bear on plants. The goal was to explore 54 commonalities and differences among members of the plant microbiome, including 55 pathogenic and commensal species, in how they impact plant gene expression. This 56 study focused on longer time scales - days instead of minutes or hours - and also 57 identified a surprisingly small set of just 24 genes, termed General Non-Self 58 Response, or GNSR (figure 1). These genes are induced by all bacterial strains, 59 regardless of the nature of each strain's relationship with the plant. The study also 60

- identified a core set of metabolites (especially those derived from the amino acid,
 tryptophan) that change in abundance when the plant is colonized by bacteria, and
- 62 found that some of those metabolites are regulated by GNSR genes.
- 63

64 This pair of studies highlights the utility of a systematic approach to characterizing

plant stress responses, particularly when using global transcriptome profiling
 technology. While previous work had examined transcriptional responses to

67 bacterial and fungal species⁴, none had done systematic, controlled comparisons

68 across different stresses and elicitors, clarifying which differences are due to true

69 differential stress response as opposed to experimental features like plant growth

70 setup, timing or mechanism of stress exposure. The data not only reveal

71 commonalities among diverse biotic stimuli, but will enable the identification of

72 differential responses to specific elicitors and bacteria, opening the door to

73 understanding, amongst other things, how plants differentiate between friend and

74 foe.

75

76 While the new gene expression resources produced by these studies shed new light

77 on the nature of plant stress responses, they also raise questions about how

78 immune responses are elicited within the spatial context of the leaf or root. Do

79 these core sets of genes behave the same way in all cell types of the plant? Are

80 there distinct features of cells directly impacted by microorganisms that are not

81 shared with those that are distal to the site of invasion? Do genes that respond

82 strongly to a biotic stimulus have a more broad expression profile across cell types,

83 or does the increased expression result from specialized action of a subset of cells?

84 These studies investigated gene expression changes across all tissues in the

85 phyllosphere (Maier, et al) or in whole seedlings (Bjornson et al.). However, tissue

86 type or spatial features could play a significant role in determining the magnitude

87 and quality of the immune response⁵. Indeed, Bjornson et al. note differences in

88 89 90	expression patterns of glutamate receptor genes of interest with regard to their bias toward stomatal guard cells.
91 92 93 94 95 96 97 98 99 100 101	New single-cell approaches, increasingly applied to plants, offer the ability to identify distinct responses of individual cell types and thereby address these important questions. Droplet-based single cell RNA-seq ⁶ and spatial transcriptomics ⁷ methods could reveal the temporal and spatial evolution of transcriptional responses ^{8,9} , either reinforcing the "core" nature of the identified gene sets or breaking them into differentially regulated modules. Such a finer-scale dissection of potentially heterogeneous responses to biotic stimuli could identify cells or tissues that house these "core" responses, further enabling their study and manipulation.
102 103 104 105 106 107 108	Author Information: Lawrence Berkeley National Laboratory LBL · Joint Genome Institute, Walnut Creek, California, USA Email: <u>bjcole@lbl.gov</u> , <u>sgtringe@lbl.gov</u>
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Figure 1: Global transcriptome analysis reveals partially overlapping core gene sets induced in response to diverse stimuli.

Microbes and purified immunity-triggering elicitors trigger an acute reprogramming 135 136 in the plant transcriptome, as do many abiotic stresses. Successful host/microbe 137 interactions result from microbes suppressing some plant immune responses, 138 however a common core set of genes (GNSR) are still influenced by microbial presence. A General Stress Response (GSR) is commonly activated by biotic and 139 abiotic stimuli alike, while a Core Immune Response (CIR) is triggered by 140 141 microorganisms or other immune elicitors. These gene sets are not mutually 142 exclusive, supporting the idea that core stress pathways are rapidly activated 143 regardless of the nature of the threat.