

CURRENT PERSPECTIVE ESSAY

The JAZ Proteins Link Jasmonate Perception with Transcriptional Changes

The oxylipin jasmonic acid (JA) and its metabolites, collectively known as jasmonates, are important plant signaling molecules that mediate biotic and abiotic stress responses as well as aspects of growth and development. Although it is well known that JA regulates transcription, the mechanism of this regulation has been largely unknown. Recently, this situation changed dramatically with the discovery of a novel family of transcriptional regulators called jasmonate ZIM-domain (JAZ) proteins. Three groups independently identified members of this family in *Arabidopsis* and showed that they function as repressors of JA-regulated transcription. Furthermore, genetic, molecular, and biochemical studies indicate that the JAZ proteins are degraded by the ubiquitin protein ligase SCF^{COI1} in response to JA. In addition, one group showed that JAZ proteins interact with the positive regulator of JA signaling, MYC2. These data support an important new model of jasmonate signaling and response. In this essay, we summarize this recent work and discuss some remaining questions concerning JA signal transduction.

THE ROLE OF SCF^{COI1} IN JASMONATE SIGNALING

Coronatine is a phytotoxin produced by the plant pathogen *Pseudomonas syringae* that is structurally related to methyl jasmonate (MeJA) and produces very similar effects when applied to plants (Feys et al., 1994). The *Arabidopsis coronatine-insensitive1* (*coi1*) mutant was isolated in a forward genetic screen exploiting coronatine's inhibitory effect on root elongation (Feys et al., 1994). This mutation confers insensitivity to coronatine and MeJA, as well as male sterility (Feys et al., 1994). Subsequent studies demonstrated that *coi1* mutants are completely deficient in every aspect of jasmonate response (Xie et al., 1998), indicating that *COI1* has an essential function in JA signaling. Ultimately, it was demonstrated that *COI1* encodes an F-box protein closely related to the auxin receptor/F-box protein TIR1 (Xie et al., 1998). This finding suggested that COI1 might act as part of an SCF (Skp/Cullin/F-box) E3 ubiquitin ligase to mediate jasmonate signaling (Xie et al., 1998). Indeed, COI1 was shown to associate with components of SCF complexes, including ASK1, ASK2, CUL1, and RBX1 (Xu et al., 2002). Further support for the importance of the ubiquitin proteasome pathway in JA signaling came from observations that mutations in genes required for SCF function, such as *AXR1*, *CUL1*, and *JAI4/SGT1b*, result in reduced jasmonate responses (Devoto et al., 2002; Xu et al., 2002; Gray et al., 2003; Lorenzo et al.,

2004; Lorenzo and Solano, 2005; Ren et al., 2005). Based on these data, it was hypothesized that SCF^{COI1} targets proteins for degradation by the 26S proteasome to promote jasmonate-mediated transcriptional changes.

JASMONATE TRANSCRIPTIONAL RESPONSES

Large-scale transcriptional profiling has identified a number of COI1-dependent jasmonate-regulated genes in *Arabidopsis* (Devoto et al., 2005; Jung et al., 2007). Moreover, a number of transcription factors that control these changes have been identified, including ERF1, WRKY70, ORA47, ORA59, and MYC2 (reviewed in Wasternack, 2007). Other recent work has demonstrated that WRKY18 (Xu et al., 2006) and two previously uncharacterized transcription factors, At1g74930 and At3g53600, which belong to the AP2/EREBP and C2H2 families, respectively (Wang et al., 2007), are positive regulators of JA responses to wounding and are dependent on COI1. The most well-characterized transcription factor in jasmonate signaling to date is MYC2, which was identified in two independent genetic screens from the mutants *methyl-jasmonate resistant1* and *jasmonate-insensitive1* (Berger et al., 1996; Lorenzo et al., 2004). MYC2 positively regulates genes involved in wounding responses but negatively regulates genes involved in pathogen defense (Lorenzo et al., 2004). Interestingly, ERF1 also differentially regulates these two branches of the JA signaling pathway but with the opposite effect of MYC2 (Lorenzo et al., 2003, 2004). Several studies have demonstrated that MYC2 preferentially binds to G-box or G-box-related hexamers in the promoters of target genes (Abe et al., 1997; de Pater et al., 1997; Yadav et al., 2005; Chini et al., 2007; Dombrecht et al., 2007). A number of genes encoding transcription factors involved in JA-mediated transcriptional regulation have these G-box motifs in their 5' regulatory sequences, including *MYC2* and *ERF1* (Dombrecht et al., 2007).

IDENTIFICATION OF THE JAZ PROTEINS

Three groups independently identified a family of proteins in *Arabidopsis* that function as repressors in the JA signaling pathway using either genetic or transcript profiling approaches. These proteins are known as JAZ proteins because their transcripts increase in abundance upon jasmonate treatment and they share a 28-amino acid ZIM domain of unknown function (Chini et al., 2007; Thines et al., 2007). In the study by Thines et al. (2007), eight members of the JAZ family were

CURRENT PERSPECTIVE ESSAY

identified as rapid JA-responsive genes in microarray experiments that involved JA treatment of the jasmonate synthesis mutant *opr3*. Database searches identified five additional genes that encoded related proteins, bringing the total number of family members to 12 (Thines et al., 2007). In a separate study, Chini et al. (2007) characterized the mutant *jasmonate-insensitive3-1* (*jai3-1*) they had identified in a forward genetic screen. Positional cloning of the *jai3-1* mutation identified a base change in an intron acceptor site of the gene now known as *JAZ3*. The *jai3-1* mutation results in the production of a truncated protein that causes a jasmonate-insensitive phenotype and impaired transcriptional responses to jasmonate (Chini et al., 2007). Finally, Yan et al. (2007) selected candidate JA signaling genes from microarray experiments and made transgenic plants overexpressing these candidate genes. One of the transgenic lines, overexpressing a predicted alternatively spliced transcript from a gene they called *Jasmonate-Associated1* (*JAS1*), reduced MeJA sensitivity in root elongation assays. Using the nomenclature of Chini et al. (2007) and Thines et al. (2007), *JAS1* is also known as *JAZ10*.

JAZ PROTEINS ARE SUBSTRATES OF SCF^{COI1}

Working with different members of this protein family, all three groups found that expression of truncated proteins lacking all or part of a conserved domain in the C terminus reduced plant sensitivity to jasmonate, whereas expression of the full-length proteins had little effect (Chini et al., 2007; Thines et al., 2007; Yan et al., 2007). Subsequent studies found that full-length *JAZ1*, *JAZ6*, and *JAI3/JAZ3* proteins are degraded following jasmonate treatment in a proteasome-dependent manner (Chini et al., 2007; Thines et al., 2007). However, these same proteins were stabilized in the *coi1-1* background, implicating SCF^{COI1} in a JAZ degradation pathway (Chini et al., 2007; Thines et al., 2007). Truncated versions of *JAZ1*, *JAZ6*, and *JAZ3*, lacking their respective C-terminal domains, were resistant to jasmonate-induced degradation (Chini et al., 2007; Thines et al., 2007). Moreover, expression of the truncated version of *JAI3/JAZ3*, either in transient expression assays or in the *jai3-1* background, blocked the degradation of full-length *JAZ3*, *JAZ1*, and *JAZ9* (Chini et al., 2007). These results suggest that the dominant jasmonate-insensitive phenotype of plants expressing truncated JAZ proteins is due to stabilization not only of the truncated protein but additional JAZ family members as well. The data are consistent with a model in which the JAZ proteins are repressors of jasmonate signaling. Furthermore, mutants harboring null T-DNA insertion mutations in *JAZ2*, *JAZ5*, *JAZ7*, and *JAZ9* failed to exhibit jasmonate-related phenotypes, suggesting that these genes function redundantly (Thines et al., 2007). Interestingly, RNA interference lines that knocked down expression of *JAS1/JAZ10* exhibited hypersensitivity when treated with MeJA compared with wild-type plants (Yan et al., 2007). Perhaps the *JAS1/JAZ10* RNA interference construct targets additional JAZ genes leading to the jasmonate phenotype.

Genetic data suggest that the JAZ proteins serve as substrates for SCF^{COI1}. To test this idea further, both the Browse and Solano groups examined the ability of *JAZ1* and *JAI3/JAZ3* to interact with *COI1* using in vitro pull-down assays, and both groups demonstrated an interaction (Chini et al., 2007; Thines et al., 2007). This interaction was confirmed for *JAZ1* using yeast two-hybrid (Y2H) assays (Thines et al., 2007). Interestingly, Chini et al. (2007), demonstrated that it is the N-terminal half of *JAI3/JAZ3* that interacts with *COI1*. This is somewhat surprising given that C-terminal domain deletions stabilize these proteins.

An important issue in JA signaling is the chemical nature of the signal. Although exogenous JA elicits a response, earlier data strongly suggest that it is not the active signaling molecule. The JASMONIC ACID RESISTANT1 (*JAR1*) protein is an adenylating enzyme that catalyzes the conjugation of JA to amino acids such as L-isoleucine (JA-Ile) (Staswick and Tiryaki, 2004). Because *jar1* mutants are resistant to the inhibitory effects of MeJA on growth, conjugated forms of JA are predicted to be the active molecules (Staswick et al., 1992). Indeed, the *COI1*-*JAZ1* protein interaction in Y2H assays was promoted by the inclusion of JA-Ile but not of JA alone, MeJA, or another precursor called 12-oxo-phytodienoic acid (Thines et al., 2007). In vitro pull-down assays confirmed the Y2H results and demonstrated that a JA-Leu derivative could promote the *COI1*-*JAZ1* interaction but was at least 50-fold less effective than JA-Ile, whereas JA-Phe and JA-Trp had no activity (Thines et al., 2007).

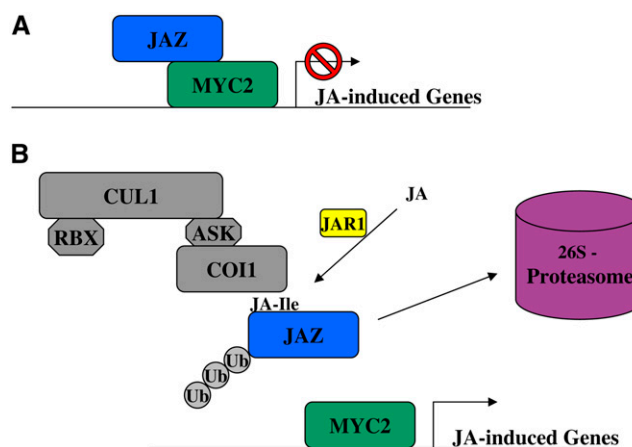


Figure 1. Model for Jasmonate Response in *Arabidopsis*.

(A) MYC2 (green) binds to G-box motifs adjacent to JA-regulated genes. JAZ proteins (blue) repress MYC2 activity through a direct interaction between the C terminus of JAZ and the N terminus of MYC2.

(B) JA is conjugated to L-Ile by *JAR1* (yellow). JA-Ile promotes the interaction of JAZ proteins with SCF^{COI1} (gray), leading to polyubiquitination (Ub) and subsequent degradation by the 26S proteasome (pink). Degradation of the JAZ repressors frees MYC2 to mediate JA-responsive transcriptional changes.

CURRENT PERSPECTIVE ESSAY

JAZ PROTEINS LINK SCF^{COI1} WITH TRANSCRIPTIONAL ACTIVATION

The JAZ proteins do not have an obvious DNA binding domain, suggesting that their effects on transcription could be indirect (Chini et al., 2007). Pull-down and Y2H assays indicate that the C-terminal domain of JAI3/JAZ3 interacts with sequences at the N terminus of MYC2 (Chini et al., 2007). Therefore, it is possible that JAI3/JAZ3 directly blocks MYC2 function. JAI3/JAZ3 degradation via SCF^{COI1} in response to JA would permit MYC2 to activate or repress downstream target genes in JA signaling pathways. Interestingly, several of the JAZ genes are themselves upregulated in response to jasmonate or are constitutively expressed in untreated plants expressing MYC2 from the cauliflower mosaic virus 35S promoter (Chini et al., 2007). These findings indicate that a negative feedback mechanism may limit the response after the initial jasmonate perception.

CONCLUSIONS AND FUTURE STUDIES

The discovery of the JAZ proteins and the elucidation of their role in JA signaling are exciting and important because it links several active areas of research into a coherent signaling pathway (Figure 1). The data presented in these recent articles imply that JAZ proteins bind and repress the transcription factors that modulate transcription of JA-responsive genes. In summary, JAR1 conjugates JA to JA-Ile, which in turn promotes the interaction between SCF^{COI1} and the JAZ repressors. This interaction results in the degradation of the JAZ proteins and subsequent derepression of transcription factors, such as MYC2. It is striking how similar the JA signaling pathway is to the auxin signaling pathway established only a few years ago with the discovery that SCF^{TIR1} serves as an auxin receptor (Dharmasiri et al., 2005a; Kepinski and Leyser, 2005). Indeed, COI1 is the closest known relative to the TIR1/AFB family of F-box proteins (Dharmasiri et al., 2005b), and based on this similarity, it has been suggested that COI1 may be a JA receptor (Parry and Estelle, 2006; Tan et al., 2007). Now, with the identification of COI1 substrates and the recent determination of the structure of TIR1, it should be possible to test this possibility directly. However, even with the establishment of COI1 as a potential site of JA perception, some fascinating questions remain. For example, what constitutes the JA receptor? Does JA-Ile bind COI1, JAZs proteins, or are both proteins required to form the receptor? How many ligands derived from JA promote the interaction of COI1 and JAZ proteins and is there any specificity with respect to different JA conjugates and members of the JAZ family? Coronatine is a phytotoxin from the plant pathogen *Pseudomonas* with structural similarity to JA-Ile. Does it directly promote COI1–JAZ interactions, and how many other pathogens produce molecules that can directly activate JA responses through this signaling mechanism?

To date, Aux/IAA proteins are the only known substrates of SCF^{TIR1/AFB} complexes, but it is possible that there are others.

Are the JAZ proteins the only substrates SCF^{COI1}? Histone deacetylases can negatively regulate transcription, have been shown to interact with COI1 in planta (Devoto et al., 2002), and have been implicated in JA signaling (Zhou et al., 2005). Could they also be substrates for COI1? On the other side of the signaling equation, how many transcription factors can the JAZ repressors bind and is there any specificity for repressor-transcription factor binding? As these questions and others are addressed in the future, we can expect new insights into the mechanism of JA signaling and its importance in various aspects of plant growth and survival.

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REFERENCES

- Abe, H., Yamaguchi-Shinozaki, K., Urao, T., Iwasaki, T., Hosokawa, D., and Shinozaki, K.** (1997). Role of arabidopsis MYC and MYB homologs in drought- and abscisic acid-regulated gene expression. *Plant Cell* **9**: 1859–1868.
- Berger, S., Bell, E., and Mullet, J.E.** (1996). Two methyl jasmonate-insensitive mutants show altered expression of AtVsp in response to methyl jasmonate and wounding. *Plant Physiol.* **111**: 525–531.
- Chini, A., Fonseca, S., Fernandez, G., Adie, B., Chico, J.M., Lorenzo, O., Garcia-Casado, G., Lopez-Vidriero, I., Lozano, F.M., Ponce, M.R., Micol, J.L., and Solano, R.** (2007). The JAZ family of repressors is the missing link in jasmonate signalling. *Nature* **448**: 666–671.
- de Pater, S., Pham, K., Memelink, J., and Kijne, J.** (1997). RAP-1 is an Arabidopsis MYC-like R protein homologue that binds to G-box sequence motifs. *Plant Mol. Biol.* **34**: 169–174.
- Devoto, A., Ellis, C., Magusin, A., Chang, H.S., Chilcott, C., Zhu, T., and Turner, J.G.** (2005). Expression profiling reveals COI1 to be a key regulator of genes involved in wound- and methyl jasmonate-induced secondary metabolism, defence, and hormone interactions. *Plant Mol. Biol.* **58**: 497–513.
- Devoto, A., Nieto-Rostro, M., Xie, D., Ellis, C., Harmston, R., Patrick, E., Davis, J., Sherratt, L., Coleman, M., and Turner, J.G.** (2002). COI1 links jasmonate signalling and fertility to the SCF ubiquitin-ligase complex in Arabidopsis. *Plant J.* **32**: 457–466.

CURRENT PERSPECTIVE ESSAY

- Dharmasiri, N., Dharmasiri, S., and Estelle, M.** (2005a). The F-box protein TIR1 is an auxin receptor. *Nature* **435**: 441–445.
- Dharmasiri, N., Dharmasiri, S., Weijers, D., Lechner, E., Yamada, M., Hobbie, L., Ehrismann, J.S., Jurgens, G., and Estelle, M.** (2005b). Plant development is regulated by a family of auxin receptor F box proteins. *Dev. Cell* **9**: 109–119.
- Dombrecht, B., Xue, G.P., Sprague, S.J., Kirkegaard, J.A., Ross, J.J., Reid, J.B., Fitt, G.P., Sewelam, N., Schenk, P.M., Manners, J.M., and Kazan, K.** (2007). MYC2 differentially modulates diverse jasmonate-dependent functions in *Arabidopsis*. *Plant Cell* **19**: 2225–2245.
- Feys, B., Benedetti, C.E., Penfold, C.N., and Turner, J.G.** (1994). *Arabidopsis* mutants selected for resistance to the phytotoxin coronatine are male sterile, insensitive to methyl jasmonate, and resistant to a bacterial pathogen. *Plant Cell* **6**: 751–759.
- Gray, W.M., Muskett, P.R., Chuang, H.W., and Parker, J.E.** (2003). *Arabidopsis* SGT1b is required for SCF(TIR1)-mediated auxin response. *Plant Cell* **15**: 1310–1319.
- Jung, C., Lyou, S.H., Yeu, S., Kim, M.A., Rhee, S., Kim, M., Lee, J.S., Choi, Y.D., and Cheong, J.J.** (2007). Microarray-based screening of jasmonate-responsive genes in *Arabidopsis thaliana*. *Plant Cell Rep.* **26**: 1053–1063.
- Kepinski, S., and Leyser, O.** (2005). The *Arabidopsis* F-box protein TIR1 is an auxin receptor. *Nature* **435**: 446–451.
- Lorenzo, O., Chico, J.M., Sanchez-Serrano, J.J., and Solano, R.** (2004). JASMONATE-INSENSITIVE1 encodes a MYC transcription factor essential to discriminate between different jasmonate-regulated defense responses in *Arabidopsis*. *Plant Cell* **16**: 1938–1950.
- Lorenzo, O., Piqueras, R., Sanchez-Serrano, J.J., and Solano, R.** (2003). ETHYLENE RESPONSE FACTOR1 integrates signals from ethylene and jasmonate pathways in plant defense. *Plant Cell* **15**: 165–178.
- Lorenzo, O., and Solano, R.** (2005). Molecular players regulating the jasmonate signalling network. *Curr. Opin. Plant Biol.* **8**: 532–540.
- Parry, G., and Estelle, M.** (2006). Auxin receptors: A new role for F-box proteins. *Curr. Opin. Cell Biol.* **18**: 152–156.
- Ren, C., Pan, J., Peng, W., Genschik, P., Hobbie, L., Hellmann, H., Estelle, M., Gao, B., Peng, J., Sun, C., and Xie, D.** (2005). Point mutations in *Arabidopsis* Cullin1 reveal its essential role in jasmonate response. *Plant J.* **42**: 514–524.
- Staswick, P.E., Su, W., and Howell, S.H.** (1992). Methyl jasmonate inhibition of root growth and induction of a leaf protein are decreased in an *Arabidopsis thaliana* mutant. *Proc. Natl. Acad. Sci. USA* **89**: 6837–6840.
- Staswick, P.E., and Tiryaki, I.** (2004). The oxylipin signal jasmonic acid is activated by an enzyme that conjugates it to isoleucine in *Arabidopsis*. *Plant Cell* **16**: 2117–2127.
- Tan, X., Calderon-Villalobos, L.I., Sharon, M., Zheng, C., Robinson, C.V., Estelle, M., and Zheng, N.** (2007). Mechanism of auxin perception by the TIR1 ubiquitin ligase. *Nature* **446**: 640–645.
- Thines, B., Katsir, L., Melotto, M., Niu, Y., Mandaokar, A., Liu, G., Nomura, K., He, S.Y., Howe, G.A., and Browse, J.** (2007). JAZ repressor proteins are targets of the SCF(COI1) complex during jasmonate signalling. *Nature* **448**: 661–665.
- Wang, Z., Cao, G., Wang, X., Miao, J., Liu, X., Chen, Z., Qu, L.J., and Gu, H.** (2007). Identification and characterization of COI1-dependent transcription factor genes involved in JA-mediated response to wounding in *Arabidopsis* plants. *Plant Cell Rep.* **27**: 125–135.
- Wasternack, C.** (2007). Jasmonates: An update on biosynthesis, signal transduction and action in plant stress response, growth and development. *Ann. Bot. (Lond.)* **100**: 681–697.
- Xie, D.X., Feys, B.F., James, S., Nieto-Rostro, M., and Turner, J.G.** (1998). COI1: An *Arabidopsis* gene required for jasmonate-regulated defense and fertility. *Science* **280**: 1091–1094.
- Xu, L., Liu, F., Lechner, E., Genschik, P., Crosby, W.L., Ma, H., Peng, W., Huang, D., and Xie, D.** (2002). The SCF(COI1) ubiquitin-ligase complexes are required for jasmonate response in *Arabidopsis*. *Plant Cell* **14**: 1919–1935.
- Xu, X., Chen, C., Fan, B., and Chen, Z.** (2006). Physical and functional interactions between pathogen-induced *Arabidopsis* WRKY18, WRKY40, and WRKY60 transcription factors. *Plant Cell* **18**: 1310–1326.
- Yadav, V., Mallappa, C., Gangappa, S.N., Bhatia, S., and Chattopadhyay, S.** (2005). A basic helix-loop-helix transcription factor in *Arabidopsis*, MYC2, acts as a repressor of blue light-mediated photomorphogenic growth. *Plant Cell* **17**: 1953–1966.
- Yan, Y., Stolz, S., Chetelat, A., Reymond, P., Pagni, M., Dubugnon, L., and Farmer, E.E.** (2007). A downstream mediator in the growth repression limb of the jasmonate pathway. *Plant Cell* **19**: 2470–2483.
- Zhou, C., Zhang, L., Duan, J., Miki, B., and Wu, K.** (2005). HISTONE DEACETYLASE19 is involved in jasmonic acid and ethylene signaling of pathogen response in *Arabidopsis*. *Plant Cell* **17**: 1196–1204.

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