Cell Wall Localization of Two DUF642 Proteins, BIIDXI and TEEBE, during Meloidogyne incognita Early Inoculation

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The root-knot nematode Meloidogyne incognita infects a variety of plants, including Arabidopsis thaliana. During migration, root-knot nematodes secrete different proteins to modify cell walls, which include pectolytic enzymes. However, the contribution of host cell wall proteins has not been described during this process. The function of two DUF642 cell wall proteins, BIIDXI (BDX, At4g32460) and TEEBE (TEB, At2g41800), in plant development could be related to the regulation of pectin methyl esterification status in the cell walls of different tissues. Accordingly, the expression of these two genes is up-regulated by auxin. BDX and TEB were highly induced during early M. incognita inoculation. Moreover, cell wall localization of the proteins was also induced. The cell wall localization of BDX and TEB DUF642 proteins during M. incognita early inoculation suggested that these two proteins could be involved in the regulation of the degree of pectin methylation during cell separation.

Keywords: Arabidopsis thaliana, DUF642 proteins, Meloidogyne incognita, plant cell wall

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DUF642 cell wall-related family of proteins that has been exclusively described in spermatophytes (Vázquez-Lobo et al., 2012). Transcriptomic analyses of *M. incognita* early infection (3 dpi) revealed an important induction of *At*1g29980 expression, a DUF642 gene (Barcalá et al., 2010). Ten genes of this family are in the *Arabidopsis thaliana* genome, and the expression of *BIIDXI* (*BDX*, *At*4g32460) and *TEEBE* (*TEB*, *At*2g41800) are highly induced by auxin in the different root zones (Salazar-Iribe et al., 2016; Zúñiga-Sánchez et al., 2014). In this study, we determined the effect of the inoculation of *M. incognita* juveniles on the expression and subcellular localization of *BDX* and *TEB* in roots from 5-d-old seedlings of *A. thaliana*. The same analysis was performed with the false root-knot nematode *Nacobus aberrans*, which also induces a specialized feeding site known as the syncytium; however, despite having a wide host range, this nematode has not been found infecting brassica crops such as broccoli and cabbage in Mexico and also cannot complete its life cycle in the *A. thaliana* ecotype Columbia (unpublished data; E. Zavaleta, personal communication).

The transgenic plants PROBDX::GFP were transformed with a construction that include an 1983 bp fragment upstream *BDX* ATG, PROTEB::GFP lines include an 890 bp fragment upstream of *TEB* ATG; at least three independent lines were analyzed showing the same expression pattern. These lines were used for expression analyses while for determination of subcellular localization PROBDX::BDX-GFP and PROTEB::TEB-GFP lines (Salazar-Iribe et al., 2016; Zúñiga-Sánchez et al., 2014) were analyzed. The seeds from these transgenic plants and wt plants (col) were germinated after imbibition at 5°C for 2-3 days and then grown vertically on petri dishes with 0.5 X Murashige and Skoog (MS) basal medium containing 0.5% sucrose and 0.6% agar. Seed germination was conducted in a CONVIRON growth chamber under long-day conditions at 20-22°C.

Second stage juveniles (J2s) of the root-knot nematodes *M. incognita* and *N. aberrans* were obtained from egg masses extracted from *Capsicum annum* and *Solanum lycopersicum* roots, respectively. Grown as described previously, 7-d-old seedlings were inoculated with 250 J2s of *M. incognita* or *N. aberrans* per plant. DR5::GFP transgenic plants were used for the control of the inoculation. As a negative control, the solution preserving nematodes was used for plant infection; the solution was centrifuged at low speed and the supernatant without nematodes was used for the inoculation. Both transgenic and wt plants were inoculated under the same conditions.

GFP was imaged at 485-545 nm with an Olympus. To-kyo, Japan. Images of plant tissues without internal fluorescence were used as controls. Images were assembled using Photoshop version 5.0 (California, USA.).

Visualization of the auxin responsive reporter DR5::GFP (Fig. 1A) was evident at two days after inoculation with J2s of *M. incognita* (Fig. 1B). Auxin was in all types of cells of the *M. incognita* inoculated roots (Fig. 1B), but no change in auxin pattern was detected in plants inoculated with *N. aberrans* (Fig. 1C). In three independent experiments, an important increase was detected in expression of both *BDX* and *TEB* (Fig. 1E, H) compared with non-inoculated roots (Fig. 1D, G). Changes in cell type expression pattern were also promoted by nematode inoculation. For *BDX*, a strong GFP fluorescence was observed only in the vascular tissue of the control roots (Fig. 1D), whereas the fluorescence was highly detected in other cell types of the infected roots (Fig. 1E). In *A. thaliana* plants inoculated with *N. aberrans*, a slight induction of auxin (Fig. 1C) or expression of *BDX* (Fig. 1F) and *TEB* (Fig. 1I) was observed at 2 days after inoculation. BDX and TEB proteins have been detected in cell wall proteomes from different plant tissues, including cell suspension cultures (Jamet et al., 2006). Cell wall localization of these two proteins in epidermal cells from primary roots treated with auxin has also been described (Salazar-Iribe et al., 2016). In non-treated roots, BDX was located intracellularly in cortex and endodermic cells (Fig. 2A). *Meloidogyne incognita* inoculation promoted BDX localization to cell walls in different cell types (Fig. 2B). For TEB, no fluorescence was detected in control roots but was localized in the cell walls in different cell types of inoculated roots (Fig. 2C, D).

The accumulation and redistribution of auxin during root-knot nematode inoculation are processes directly involved with plant susceptibility to infection (Kyndt et al., 2016). An important increase in auxin was detected in the roots of the seedlings inoculated with *M. incognita*, whereas almost no increase in auxin was observed for *N. aberrans* inoculation (Fig. 1). This absence of increase in auxin is consistent with previous observations that *A. thaliana* is not susceptible to *N. aberrans* infection. Additionally, a differential regulation in the expression of plant cell wall-related genes between susceptible and resistant *S. lycopersicum* to *M. incognita* infection has been described (Shukla et al., 2017). The up-regulation of *BDX* and *TEB* promoters observed in roots during *M. incognita* juvenile migration suggested a potential transcriptional regulation of the expression of these genes upon nematode compatible infection (Fig. 1). By contrast, no changes in *TEB* and *BDX* expression were observed during *N. aberrans* inoculation; this response is similar to that described in resistant *S. ly-
corpersicum in which no significant alteration is observed in the expression of genes involved in the modulation of cell wall architecture (Shukla et al., 2017).

The relevance of cell wall-modifying enzymes as phytonematode effectors that aid in nematode invasion and migration into and within hosts has been extensively documented in different plants (Ibrahim et al., 2011; Shivakumar et al., 2017). The pectolytic enzymes secreted by the nematode are involved in cell separation by promoting the degradation of the middle lamella, which is the pectin layer that fills the space between adjacent cells. However, no role of plant cell wall proteins has been described during this process. The promotion of the expression and cell wall localization of BDX and TEB by *M. incognita* J2 inoculation suggests that the early plant-pathogen interaction also involves the induction of plant proteins that participate in cell wall remodeling. These results also suggest that host plant cell wall proteins are involved in the cellular separa-
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Ition for the intercellular migration of J2s. Plant cells are firmly adhered to one another by the middle lamellae, and the degradation of the low methylated pectin by PGs and PLs promotes cell separation. Pectinesterases or PMEs and the regulators are involved in maintaining a low degree of pectin methylesterification. The role of DUF642 proteins in pectin methyl esterase regulation has been described in A. thaliana (BDX and TEB), Vitis quinquangulataris (VqDUF642) and Amaranthus hypochondriacus (AhDGR2). An increase in PME activity in leaves from overexpressed VqDUF642 A. thaliana plants has been described, with the conclusion that VqDUF642 is involved in plant development and defense against pathogenic infections (Xie and Wang, 2016). The cell wall localization of BDX and TEB DUF642 proteins during M. incognita inoculation suggested that these two proteins could be involved in the regulation of the degree of pectin methylation during cell separation, as was previously described during testa rupture (Zúñiga-Sánchez et al., 2014).

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