INTRODUCTION

*Xylella fastidiosa* (Xf) is the causal agent of Pierce’s Disease (PD) of grapevine, a serious and often lethal disease (Hopkins and Purcell, 2002, Chatterjee et al., 2008, Purcell and Hopkins, 1996). This xylem-limited bacterial pathogen colonizes the xylem and in doing so must be able to move efficiently from one xylem vessel element to adjacent vessels (Roper et al. 2007). Xylem conduits are separated by pit membranes (PMs) that are composed of primary cell wall and serve to prevent movement of air embolisms and pathogens within the xylem (Buchanan, 2000). More specifically, PMs are composed of cellulose microfibrils embedded in a meshwork of pectin and hemicellulose (Buchanan, 2000). The pore sizes within that meshwork range from 5 to 20 nM, which will not allow passive passage of Xf cells whose size is 250-500 x 1,000-4,000 nM (Perez-Donoso et al., 2010, Mollenhauer & Hopkins, 1974). Based on functional genomics and *in planta* experimental evidence, Xf utilizes cell wall-degrading enzymes (CWDEs) to actively digest the polymers within the PMs, thereby facilitating its movement throughout the xylem network (Simpson et al. 2000, Roper et al., 2007, Perez-Donoso et al., 2010). It is known that polygalacturonase (PG) is a major pathogenicity factor for Xf (Roper et al., 2007) and that it acts in concert with at least one EGase to breach the PM barrier (Perez-Donoso et al. 2010). EGases are implicated in virulence and colonization of the xylem in other bacterial phytopathogens, such as *Pantoea stewartii* subsp. *stewartii*, *Ralstonia solanacearum* and *Xanthomonas campestris* pv. *campestris* (Gough, 1988, Roberts et al., 1988, Saile et al., 1997, Mohammadi et al., 2012). In our previous study (project # 14-0144-SA), we tested the role of the Xf EGases *in planta* by constructing deletion mutants in two of the EGases (Δ*engXCA1* and Δ*engXCA2*) and mechanically inoculating the modified Xf lines into *Vitis vinifera* cv. Cabernet sauvignon and cv. Chardonnay grapevines. Interestingly, both Δ*engXCA1* and Δ*engXCA2* achieved the same titers (*data not shown*) in the Cabernet sauvignon vines as wild type Xf, yet they were significantly less virulent and elicited fewer PD symptoms (Fig 1A,B).

PD symptom development is tightly correlated with the ability of Xf to degrade specific polysaccharides, namely fucosylated xyloglucans (part of the hemicellulosic component) and weakly esterified homogalacturonans (part of the pectin portion), that make up the intervessel PMs (Sun et al., 2011). In general, pectin is one of the first targets of cell wall digestion for invading pathogens and the resulting oligogalacturonides (OGs), which are smaller pieces of the pectin polymer, that are released are likely used as a carbon source for the invading pathogen. In addition, specific OGs with a degree of polymerization in the size range of 10-15 residues can also serve as signals that trigger host defense responses (Benedetti et al., 2015). These responses include accumulation of reactive oxygen species (ROS), expression of pathogenesis-related proteins, deposition of callose, activation of mitogen-activated protein kinases (MAPKs), among other defense related processes (Boller & Felix, 2009, Benedetti et al., 2015).
Figure 1: Pierce's disease development over 15 weeks in Cabernet Sauvignon grapevines after inoculation with wild-type Temecula 1 (blue), and the ΔengXCA1 (red) or ΔengXCA2 (orange) mutant strains. 1X PBS (green) served as the negative control. All vines were rated on a disease scale of 0-5, where 0 = healthy, 1-4 = increasing degrees of scorching, and 5 = vine death. Both the ΔengXCA1 and ΔengXCA2 mutant strains maintained lower average disease scores per week relative to the wild-type strain. Data are the means of three independent assays with ten replicates each. Bars represent the standard error of the mean.

Tyloses are outgrowths of parenchyma cells that emerge through vessel-parenchyma pits into vessel lumen, and are common in a wide range of species (Bonsen and Kučera 1990; Esau 1977; Tyree and Zimmermann 2002). Tyloses impede fluid penetration (Parameswaran et al. 1985) and induce a permanent state of reduced hydraulic conductivity, and are triggered by abiotic and biotic stresses, such as pathogen infection (Aleemullah and Walsh 1996; Collins et al. 2009; Dimond 1955; Parke et al. 2007). Tylose formation is the predominant vascular occlusion associated with Xf infection (Fig 2A,B), and excessive tylose development has been linked to the extreme susceptibility of Vitis vinifera wine grapes to PD (Fritschi et al. 2008; Sun et al. 2013).

Figure 2: Xylem vessels of V. vinifera grapevines inoculated with Xf. A) Longitudinal section B) cross-section. Grapevine petiole sections were stained with toluidine blue O (0.05%). White arrows and bracket indicate vessels that are completely occluded with tyloses, and yellow arrow indicates a partially occluded vessel. Images taken by J. Rapicavoli (Roper Lab).

Importantly, rates of tylose development in V. arizonica, a resistant species, are much lower than those in V. vinifera, which may reflect differing innate immune responses to the presence of Xf in the xylem. To our knowledge, no one has looked at the molecular mechanisms underlying the differences in response to Xf among
different *V. vinifera* cultivars. Thus, we sought out to better understand this difference in cultivar response to *Xf* in the context of host cell wall degradation and the elicitation of specific defense responses that lead to tylose formation in grapevines. Interestingly, a preliminary analysis of tylose formation in Cabernet Sauvignon vines inoculated with the Δ*engXCA1* mutant using a high resolution microCT technique (a kind of CAT scan) by the McElrone laboratory determined that these vines exhibited fewer tyloses than those inoculated with wild type *Xf* (Fig. 3). Therefore, our hypothesis is that enzymatic degradation of the plant cell wall by *Xf* CWDEs is generating cell wall fragments that elicit DAMP signaling defense pathways, which leads to downstream tylose production and PD symptom development in certain grape cultivars.

**Figure 3:** Images of grapevine xylem vessels obtained using microCT. Vines inoculated with wild-type Temecula 1 *Xf* had substantial vascular occlusions, whereas vines inoculated with Δ*engXCA1* had few tyloses similar to the PBS negative control (not shown). Top panels are cross-sectional views and bottom panels are longitudinal views. White brackets highlight occluded vessels and black bracket highlights open vessel.

**OBJECTIVES**

1) Qualitative analysis of the effect of cell wall degradation on the grapevine response to *Xf*.
2) Quantitative analysis of plant defense pathways induced by *Xf* cell wall degrading enzyme activity: biochemical and transcriptional studies.
3) Inhibition of the Type II secretion system using natural products produced by grapevine microbial endophytes.

**DESCRIPTION OF ACTIVITIES**

**Qualitative analysis of the effect of cell wall degradation on the grapevine response to *Xf***.

In the context of plant cell wall degradation, we are examining the effects that different *Xf* mutants (Δ*engXCA1*, Δ*engXCA2*, *egl* (all EGases and EGase/expansin hybrid) and *pglA* (a PG)) have on integrity and carbohydrate composition of grapevine pit membranes in different varieties using both microscopic and immunological techniques coupled with fluorescence (Sun et al., 2011) and/or electron (Sun et al., unpublished) microscopy. We are coupling these microscopic observations with macroscopic studies of the spatial distribution of tyloses and other vascular occlusions, such as plant-derived gels and bacterial aggregates using high resolution micro-computed tomography (microCT). This non-destructive method technique uses x-rays to create cross-sections of an object that can be used to re-create a virtual model (3D model). These experiments will allow us to match degradation of specific host cell wall carbohydrates with spatiotemporal patterns of production of tyloses in 3 dimensions.
Xf wildtype and mutant strains (ΔengXCA1, ΔengXCA1, ΔengXCA1/ΔengXCA2, ΔpglA and Δegl) have been used to inoculate Cabernet Sauvignon and Chardonnay grapevines in the greenhouse. PBS-inoculated vines were used as negative controls. Each Xf strain was inoculated into 27 plants (3 biological replicates with 9 technical replicates each) and PD symptoms were rated each week using the 0 – 5 PD rating index (Guilhabert and Kirkpatrick, 2005). Vine tissue samples are currently being collected for each of the three experiments: stem and petiole tissue for RNAseq, stem tissue for microCT analysis, and stem explants for EM analysis. Samples from three biological replications (consisting of three technical replications) per treatment have been collected at two time-points covering early and mid-infection based on the PD rating index (Early infection = 1 – 2, Mid-infection = 2 – 3) and are currently being analyzed or awaiting analysis.

**Modifications of different Xf strains on xylem structures of Chardonnay vines**—Early time-point samples from Chardonnay were analyzed using scanning electron microscopy to study vascular occlusion, pit membrane integrity and presence/absence of Xf in the xylem tissue after inoculation with wild-type Temecula 1, the PBS negative control, the ΔengXCA1 mutant, or the ΔpglA mutant. Our results indicate that no vascular occlusions have been observed in the vines inoculated with PBS (Fig. 4), wild-type Temecula 1 (Fig. 5A, B) and ΔpglA (Fig. 7A), respectively at the early time point. Tyloses were found in very few vessels of the vine inoculated with ΔengXCA1 (Fig. 6A, B), but were at the early developmental stages and did not occlude the vessels where they occurred. Xf cells were not observed in all the vines except that inoculated with ΔengXCA1 (Fig. 6D). Vessel-parenchyma pit membranes (PMs) were intact in the vines with the four different inoculums (Figs. 4B and 5C). Some broken (degraded) intervessel PMs were observed in the vine inoculated with either wild type (Fig. 5C, D) or ΔengXCA1 (Fig. 6C, D), but were rare or absent in the vine inoculated with either PBS or ΔpglA. (Fig. 7C). EM data is still being collected and analyzed for the remaining Chardonnay samples from the mid time point and the late time point.

**Figure 4:** Early time point observations in Chardonnay xylem: Vessel structural features in a Chardonnay stem inoculated with the PBS negative control. A. Transverse section of secondary xylem showing absence of vascular occlusion in the vessels. B. Tangential longitudinal section of secondary xylem, showing three transected vessels that have intact vessel-parenchyma PMs and do not contain vascular occlusions.
Figure 5: Early time point observations in Chardonnay xylem: Vessel structural features in a Chardonnay stem inoculated with wild-type Temecula 1. A. Transverse section of secondary xylem showing absence of vascular occlusion in the vessels. B. Tangential longitudinal section of secondary xylem. Vessels do not contain vascular occlusions. C. A transected vessel, showing oval vessel-parenchyma pit pairs and intact PMs (short arrows) and scalariform intervessel pit pairs (long arrows). D. Broken intervessel PMs.
Figure 6: Early time point observations in Chardonnay xylem: Vessel structural features in a Chardonnay stem inoculated with ΔengXCA1. A. Transverse section of secondary xylem. Most vessels have empty lumens but few vessels are filled with tyloses (arrow). B. Tangential longitudinal section of secondary xylem, showing a transected vessel with developing tyloses inside. C. Scalariform intervessel pit pairs in a vessel lateral wall. D. Enlargement of several intervessel pit pairs in a vessel lateral wall. Broken intervessel PMs (long arrows) are seen from a pit aperture and ΔengXCA1 cells (short arrows) are present on the lateral wall.

Figure 7: Vessel structural features in a Chardonnay stem inoculated with ΔpglA. A. Transverse section of secondary xylem. All the vessels are free of vascular occlusions. B. Tangential longitudinal section of secondary xylem, showing several transected vessels without vascular occlusion. C. Surface view of a vessel’s lateral wall. Whole intervessel PMs are visible after removal of secondary wall borders of intervessel pits. Intervessel PMs are intact and they are horizontally elongated and have a ladder-like arrangement along the vessel axial direction.
**Modifications of different Xf strains on xylem structures of Cabernet Sauvignon vines** - We examined some late time-point stem samples of Cabernet Sauvignon vines that were inoculated with PBS, wild-type Xf strains (Temecula 1 and Fetzer) and mutant Xf strains (ΔengXCA1, ΔengXCA2, ΔengXCA1/engXCA2, Δegl and ΔpglA), respectively. We found that at the late time-point of PD symptom development, certain Xf strains display differences in vascular occlusion, intervessel PM integrity and Xf existence at the third internode above the point of inoculation. Some preliminary data from the middle time-point are also included here to explore the process of xylem structure modifications during PD symptom progression. The remaining samples from the mid and late time points are still being analyzed as well as the early time points.

In the vines inoculated with PBS, vascular occlusion and Xf cells were not observed in the secondary xylem at both middle (Fig. 8) and late (Fig. 9) time-point. Intervessel PMs observed remained intact at the middle time-point but broken PMs were seen in a few vessels in the samples of the late-time point.

The inoculation with wild-type Xf also caused significant xylem structural modifications in Cabernet Sauvignon vines. At the middle time-point, about 30% of the vessels in the transverse section of secondary xylem contained tyloses (Fig. 10A), which partially or completely occluded the vessels (Fig. 10B). Wild-type Xf cells were also present in many vessels and occurred individually (Fig. 11C) or as small clusters (Fig. 11D). At the late time-point, over 50% of the vessels in the transverse section of a stem were occluded by tyloses and Xf cells occurred as large clusters in addition to individual occurrence or small clusters (Fig. 12A). Moderate amounts of degraded intervessel PMs were observed in both the middle and late time-point samples (Figs. 10C and 11B).

In ΔengXCA2-inoculated Cabernet Sauvignon vines, vascular occlusion occurred in a small extent at the middle time-point (Fig. 12A and B) and increased in quantity at the late time-point with the occluded vessels making up about 30% of the total vessels (Fig. 13A and B). ΔengXCA2 cells were present in the samples of both time-points (Fig. 12C and 13C, D). There are more broken intervessel PMs at the late time-point (Fig. 13C and D) than at the middle time-point (Fig. 12C). However, the inoculation with ΔengXCA1 resulted in little or no vascular occlusion in the samples of the late time-point (Fig. 14A). Some degrading intervessel PMs with different porosities were also seen in the ΔengXCA1-infected samples at the late time-point (Fig. 14B and C). Interestingly, in the late time-point samples inoculated with the ΔengXCA1/engXCA2 double mutant, tyloses were absent or occurred in very few vessels (Fig. 15A and B) and ΔengXCA1/engXCA2 cells were not observed (Fig. 15C). Intervessel PMs were mostly intact despite the existence of degrading intervessel PMs in few vessels (Fig. 15D).

The inoculation with Δegl also caused occlusion of a moderate number of vessels in the infected Cabernet Sauvignon vines at the late time-point (Fig. 16A and B). Δegl cells were abundantly present in some vessels and broken intervessel PMs in few vessels (Fig. 16C).

In the Fetzer-inoculated late time-point samples, most vessels were free of occlusions (Fig. 18A) and tyloses present in few vessels were at early developmental stages and did not occlude the vessels (Fig. 18D). Fetzer cells and broken intervessel PMs were present only in few vessels (Fig. 18C). Similarly, in the ΔpglA-inoculated samples, vessels were almost free of occlusions (Figs. 17A, B and 18B) and broken intervessel PMs (Fig. 17C) were seen in few vessels at both middle time-point and late time-point. (Fig. 18B). (Fig. 7C). ΔpglA cells were not observed in either of the time-points.
Figure 8: Xylem structural features in PBS-inoculated Cabernet Sauvignon vine at the middle time-point. A. Transverse section of secondary xylem. All the vessels are free of vascular occlusions. B. Tangential longitudinal section of secondary xylem showing two transected vessels without vascular occlusions.

Figure 9: Xylem structural features in PBS-inoculated Cabernet Sauvignon vine at the late time point. A. Transverse section of stem secondary xylem, showing absence of occluded vessels. B. Longitudinal section of stem secondary xylem, showing vessels free of tyloses.
Figure 10: Xylem structural features in wild-type Temecula 1-inoculated Cabernet Sauvignon vine at the middle time-point. A. Transverse section of stem secondary xylem, showing a large number of vessels occluded (arrows). B. Tangential longitudinal section of secondary xylem, showing one empty vessel and three vessels completely occluded by tyloses (arrowed). C. A longitudinally transected vessel. Intervessel PMs are partially degraded (arrows) and wild-type Temecula 1 cells occur mostly individually. D. wild-type Temecula 1 cells occur as small clusters.
Figure 11: Xylem structural features in wild-type Temecula 1-inoculated Cabernet Sauvignon vine at the late time-point. A. Longitudinal section of stem secondary xylem, showing abundant presence of wild-type cells in a vessel. B. A longitudinally transected vessels, showing that intervessel PMs have completely disappeared.

Figure 12: Xylem structural features in ΔengXCA2-inoculated Cabernet Sauvignon vine at the middle time-point of PD symptom development. A. Transverse section of stem secondary xylem. Few vessels are occluded (arrows). B. Longitudinal section of stem secondary xylem, showing two transected vessels fully occluded by tyloses. C. Broken intervessel PMs (arrows) and ΔengXCA2 cells in a longitudinally transected vessel.
Figure 13: Xylem structural features in ΔengXCA2-inoculated Cabernet Sauvignon vine at the late time-point of PD symptom development. A. Transverse section of secondary xylem, showing occlusion in some vessels. B. Longitudinal section of secondary xylem, show two transected vessels fully occluded by tyloses. C. A longitudinally transected vessel, showing an abundant presence of ΔengXCA2 cells. D. ΔengXCA2 cells on partially some degraded intervessel PMs (arrows indicate pores or cracks in the PMs).
Figure 14: Xylem structural features in ΔengXCA1-inoculated Cabernet Sauvignon vine at the late time point of PD symptom development. A. Longitudinal section of stem secondary xylem, showing open vessels. B and C. Longitudinally transected vessels, showing intervessel PMs with large (B) and small (C) pores, respectively.

Figure 15: Xylem structural features in ΔengXCA1/ΔengXCA2-inoculated Cabernet Sauvignon vine at the late time point of PD symptom development. A and B. Transverse section of stem secondary xylem, showing vessels free of occlusions. C. Longitudinal section of secondary xylem, showing empty vessels with mostly intact PMs. D. A longitudinally transected vessel, showing pores of different sizes in intervessel PMs.
Figure 16: Xylem structural features in Δegl-inoculated Cabernet Sauvignon vine at the late time-point of PD symptom development. A and B. Transverse section of stem secondary xylem, showing occurrence of vascular occlusion in some vessels (A) and fully occluded vessels (B). C. A longitudinally transected vessel, showing Δegl cells (short arrows) and broken PMs (long arrows).

Figure 17: Xylem structural features in ΔpglA-inoculated Cabernet Sauvignon vines at the middle time-point of PD symptom development. A. Transverse section of secondary xylem, showing vessels are free of vascular occlusions. B. Tangential longitudinal section of secondary xylem, showing two transected vessels without tyloses. C. The surface view of intervessel PMs, showing that small pores and cracks on several PMs (arrows).
Figure 18: Xylem structural features in wild-type Fetzer (A, C, D) and ΔpglA (B)-inoculated Cabernet Sauvignon vines at the late time-point of PD symptom development. **A and B.** Transverse section of stem secondary xylem, showing open vessels. **C.** A longitudinally transected vessel, showing broken intervessel PMs (long arrows) and Fetzer wild-type cells (short arrows). **D.** Small tyloses (arrows) are present in a longitudinal transected vessel.

In addition to samples imaged via electron microscopy, samples from the early and middle time-points in both Chardonnay and Cabernet Sauvignon have also been analyzed by microCT. This technique is particularly resource-intensive and thus, imaging all nine samples per treatment was not feasible. Instead, three samples per treatment were chosen randomly, and singular midslice images were analyzed to determine if tyloses formed in the xylem in response to *Xf* infection. Cabernet Sauvignon vines inoculated with wild-type Temecula 1 or ΔengXCA2 exhibited the most blocked vessels by tyloses at all time points, whereas vines inoculated with ΔengXCA1 exhibited fewer tyloses (Fig. 19). Additionally, vines inoculated with the wild type Fetzer strain and the *pglA* mutant exhibited very few tyloses, and vines inoculated with PBS (negative control) displayed no tyloses.

Transverse and longitudinal images slices of the selected samples from the early time-point in Chardonnay were also performed to visualize tylose formation, and machine learning algorithms were used to detect and quantify tyloses within vessels (Fig. 20). Several vessels from vines inoculated with wild-type Temecula 1 displayed tyloses, while fewer vessels were occluded in vines inoculated with the ΔengXCA1 mutant (Fig. 21). Vessels from vines inoculated with the PBS negative control were occlusion-free and displayed no tylose formation. Analysis of transverse and longitudinal images slices from the other time-points in both varieties are currently being analyzed.
The McElrone lab recently developed a method to measure starch content in ray and axial parenchyma (RAP) in vivo using microCT and machine learning algorithms (Earles 2018). In microCT images, x-ray absorption corresponds to the distinct molecular structure of air, water, starch and cell wall material, which enables the visualization of RAP, which are located in xylem tissue between radial files of vessels. While microCT images pictured here are of dried stems, patterns of full/empty RAP reflect those found in vivo in grapevine rootstocks and the method has implications for tracking starch utilization over the course of Xf infection. RAP in Cabernet Sauvignon vines inoculated with wild-type Temecula 1 show patterns of starch depletion at an early timepoint with significant depletion at a late timepoint, while RAP in ΔengXCA1-inoculated vines show RAP tissue full of starch at an early timepoint and moderate depletion at a late timepoint (Fig 22). RAP in PBS-inoculated vines remain full of starch at all timepoints. This technique is currently being used to analyze all samples for starch depletion.

**Tylose formation in Cabernet Sauvignon**

![Bar chart showing tylose formation in Cabernet Sauvignon](image)

**Figure 19:** Manual midslice analysis of %-tyloses (occluded vessels/total vessels) per treatment in Cabernet Sauvignon. Vessels with tyloses were manually counted on midslices of microCT scans. Wild-type Temecula 1 treatment showed high tylose formation relative to wild-type Fetzer. ΔengXCA1 treatments exhibited less tyloses than wild-type Temecula 1 overall, and a reduction of occluded vessels from early to late timepoints. ΔengXCA2 treatments show tylose formation comparable to wild-type Temecula 1.
**Figure 20:** Improved tyloses detection/quantification. Colored outlines in A (xy-axis) and B (yz-axis) correspond with C to help orient the viewer. Tyloses (highlighted in yellow) are small and rare features relative to empty vessels on the xy-axis, and can easily be confused with interconnected vessels, yet appear more distinctly in the yz-axis. D, longitudinal slices on the yz-axis can be used to train machine learning algorithms to automatically classify vessels containing tyloses for high throughput analysis.

**Figure 21:** ImageJ orthogonal views of tyloses in Chardonnay (early timepoint) vines inoculated with wild-type Temecula 1, ΔengXCA1, or PBS (negative control). Colored arrows on tranverse image slices (top) correspond to highlighted vessels of same color on the longitudinal image slices (bottom), cut from the vertical green line in the transverse image. Empty vessels appear dark gray, while tyloses appear as highly branched membranes within vessel elements. PBS buffer treatment exhibits no tylose formation.
**Figure 22:** Visual classification of ray and axial parenchyma (RAP) regions as full (magenta) or empty (yellow) in Cabernet Sauvignon vines inoculated with either wild-type Temecula 1, ΔengXCA1, or PBS (negative control). Longitudinal slices of outlined, late-timepoint RAP emphasize a spatial pattern of starch depletion with empty cells (dark airspace and light cell walls indicated with corresponding triangles) near the periphery bark (Ba) layer progressing towards the pith (Pi).

**Quantitative analysis of plant defense pathways induced by Xf cell wall degrading enzyme activity: biochemical and transcriptional studies.**

Pit membrane degradation by Xf CWDEs likely results in the release of small chain carbohydrates into the xylem. These oligosaccharides have been known to act as elicitors of plant immunity (i.e. damage-associated molecular patterns). It is possible that oligosaccharides released from pit membrane degradation are being recognized by associated parenchyma cells, triggering defense responses such as tylose production. To test this hypothesis, we are using RNAseq to analyze the grapevine transcriptome to determine if pit membrane degradation products act as elicitors of plant immunity and trigger tylose production. All tissue samples used for RNA extraction have been collected from the same plants and time-points as used for the qualitative experiments so that we can determine correlations between defense gene expression, pit membrane degradation, and/or tylose production. As these samples come from the same plants used in the qualitative experiments, all treatments, grapevine varieties, sample sizes, and time-points used are the same as in the previous section. Currently, stem and petiole tissue for all treatments from each time-point and variety have been collected, and are being prepared for RNAseq analysis.

**Inhibition of the Type II secretion system using natural products produced by grapevine microbial endophytes.**

Given that Xf CWDEs are important for the degradation of pit membranes (thus allowing systemic colonization), and their potential role in inducing tylose formation, it is imperative that these virulence factors are targeted for inhibition. However, inhibiting each CWDE individually as a commercial strategy for controlling Xf is both impractical and costly. Interestingly, these CWDEs are predicted (using SignalP software) to be secreted via the Type II secretion system (T2SS). The T2SS is a molecular nanomachine that transports pre-folded proteins from the periplasm across a dedicated channel in the outer membrane (Cianciotto, 2005, Korotkov et al., 2012). The T2SS systems of many plant and animal pathogens are either known or predicted to secrete proteins, namely
polymer degrading enzymes, which are involved in nutrient acquisition (Jha et al., 2005). The Xf CWDEs being studied in this proposal are predicted (using SignalP software) to be secreted through the T2SS. Proteins destined for secretion by the T2SS are first delivered to the periplasm via the Sec or Tat-dependent secretion pathway where they are folded (Slonczewski, 2014). Xf appears to only possess the Sec-dependent secretion pathway. Because of our interest in host CWDEs and their mechanism of secretion, we created a mutation in the xpsE gene, which encodes the putative ATPase that powers the T2SS. Grapevines inoculated with the xpsE mutant never developed PD symptoms and remained healthy, a phenotype similar to the grapevine response to the Xf pglA mutant (Fig. 23).

![Figure 23](image)

**Figure 23:** The Xf T2SS is necessary for PD development in grapevine. The ΔxpsE mutant does not induce PD symptoms in *V. vinifera* grapevines. Disease severity was based on a visual disease scale from 0 (no disease) to 5 (dead). Vines inoculated with 1X PBS (negative control) did not develop PD symptoms.

We hypothesize that this is due to the pathogen’s inability to secrete the CWDEs necessary for xylem colonization. In addition, we have indirect experimental evidence that *Xf* utilizes the T2SS to secrete PG. We observed that the ΔxpsE mutant produces visibly less EPS on XFM minimal medium containing pectin as the sole carbon source, resulting in a much less mucoid phenotype (data not shown). However, when wild type *Xf* and ΔxpsE are grown on XFM+galacturonic acid (i.e., the monomeric sugar that makes up the pectin polymer) or on XFM+glucose, both strains produce similar amounts of EPS. We infer from this that, indeed, breakdown of the pectin substrate is necessary to produce EPS and when the T2SS is disrupted this prevents secretion of PG and the subsequent breakdown of pectin.

Thus, we have compelling *in planta* and *in vitro* preliminary data indicating that *Xf* has a functional T2SS system and the proteins secreted by T2SS are critical for the infection process. From this we reason that the T2SS represents an excellent target for disease control because disrupting this system would provide comprehensive inhibition of secretion of PG (the major pathogenicity factor for *Xf*) and the other auxiliary CWDEs (Roper et al. 2007 and recent results discussed above). Therefore, identifying molecules that can inhibit T2SS function is an excellent avenue of research to pursue to develop strategies that mitigate PD by preventing pathogen ingress. This work is ongoing.

**PUBLICATIONS AND PRESENTATIONS**

**Poster Presentations**
Brian M. Ingel, Dario Cantu, Andrew McElrone, Qiang Sun, John Labavitch, and M. Caroline Roper. Characterization of *Xylella fastidiosa* plant cell wall degradation and inhibition of the Type II secretion machinery (2016). Pierce’s Disease Research Symposium, San Diego, CA.
Xf is the causal agent of PD of grapevine, a serious and often lethal disease of grapevines (Hopkins and Purcell, 2002, Chatterjee et al., 2008, Purcell and Hopkins, 1996). This xylem-limited bacterial pathogen colonizes the xylem and in doing so must be able to move efficiently from one xylem vessel element to adjacent vessels (Roper et al. 2007). Xylem conduits are separated by pit membranes (PMs) that are composed of primary cell wall that serve to prevent movement of air embolisms and pathogens within the xylem (Buchanan, 2000). More specifically, PMs are composed of cellulose microfibrils embedded in a meshwork of pectin and hemicellulose (Buchanan, 2000). The pore sizes within that meshwork range from 5 to 20 nM, which will not allow passive passage of Xf cells whose size is 250-500 x 1,000-4,000 nM (Perez-Donoso et al., 2010, Mollenhauer & Hopkins, 1974). Based on functional genomics and in planta experimental evidence, Xf utilizes host CWDEs to actively digest the polymers within the PMs, thereby facilitating its movement throughout the xylem network (Roper et al., 2007, Perez-Donoso et al., 2010). This previous work demonstrated that a polygalacturonase (PG), PglA, was required for movement and pathogenicity in grape (Roper et al. 2007). In addition, an EGase (EngXCA2) worked in concert with PG to breach pit membranes (Perez-Donoso et al. 2010). Based on these findings, inhibition of PG has been identified as a top research priority by the PD research board. Several other research groups are working towards inhibiting PG in planta as a means of PD control. In our currently supported project (project # 14-0144-SA), we have outlined objectives designed to complement and augment these current research efforts that are aimed at inhibiting PG. Our central hypothesis is that Xf utilizes other CWDEs in concert with PG to breach the pit membranes and that the majority of these are secreted by a common mechanism, the Type II Secretion System (T2SS). Our project is composed of two broad goals: 1) Elucidation of how the plant perceives host cell wall damage inflicted by the suite of Xf CWDEs during the infection process and 2) Utilization of natural products produced by grapevine microbial endophytes to inhibit the T2SS that delivers PG, and other CWDEs (EGases) to the xylem. We view this as a comprehensive approach to achieving disease control with the potential impact being to effectively disrupt systemic spread of Xf and vascular occlusions in the xylem and, therefore, PD development.

LAYPERSON SUMMARY

Xylella fastidiosa (Xf) relies on degradation of the plant cell wall to move within the grapevine, which occurs through cooperation between at least two classes of enzymes that target different carbohydrate components of the complex scaffold of the plant cell wall. A major goal of this project is to elucidate the mechanisms that lead to disassembly of the plant cell wall that eventually leads to systemic colonization of Xf in grapevines. Here we are performing experiments designed to better understand what facilitates movement of the bacterium and the subsequent clogging of the water-conducting cells that worsens Pierce’s Disease severity. In addition, we also are designing experiments to inhibit the secretion machinery responsible for delivering the Xf enzymes that are involved in Xf movement throughout the plant, thus, providing a comprehensive approach to restriction of Xf and disease development rather than targeting individual enzymes.

STATUS OF FUNDS

The funding for this project is largely going towards supporting a Ph.D. graduate student, Mr. Brian Ingel. This project is the main focus of his Ph.D. dissertation. We anticipate spending the remainder of the salary, supply, services and greenhouse recharge money associated with this project as it progresses.

SUMMARY AND STATUS OF INTELLECTUAL PROPERTY

There is no intellectual property associated with this project to date.

REFERENCES


FUNDING AGENCIES
Funding for this project was provided by the CDFA Pierce’s Disease and Glassy-winged Sharpshooter Board.