# Sword fern die-off phenomenon:

## Evaluating the relationship between soil moisture and the withering ferns of Seward Park





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Figure 1: Example of a small patch of sword ferns that recently died in Seward Park.

### <u>Abstract</u>

Sword ferns (Polystichum munitum) are rapidly declining in Seward Park, one of the few remnant oldgrowth forests in the Puget Sound lowlands. Based on recent transmission experiments, we hypothesize that a pathogen may be responsible for the phenomenon. However, other studies suggest that the phenomenon is driven by seasonal drought stress, mediated by local site conditions that influence soil moisture. We compared areas of Seward Park containing symptomatic sword ferns with areas of Seward Park and Schmitz Preserve Park containing healthy ferns. In each experimental group, we surveyed the soils, site conditions, and plant communities. We also collected tissue samples to measure the foliar moisture content of sword ferns, and to compare their microbial endophyte communities using the Illumina MiSeq platform. Foliar moisture content was significantly lower in symptomatic sword ferns compared with healthy ferns. However, foliar moisture content was not correlated with soil moisture content, which was similar between all three experimental groups. Soil bulk density was also similar between the symptomatic and healthy areas of Seward Park. Widespread mortality was not observed among sword ferns in Schmitz Preserve Park, where the soils are more coarsely textured than in Seward Park. Although the microbiome component of our study is ongoing, our preliminary results remain consistent with the pathogen hypothesis. Our findings suggest that the sword fern decline is driven by a process independent of drought stress or the site conditions that influence soil moisture content.



### Introduction

Over the past decade, sword ferns (*Polystichum munitum*) have died in sudden mortality events at dozens of locations throughout the Puget Sound region (Shannon, 2020a). In 2013, a particularly large mortality event was observed in a remnant old-growth forest in Seward Park, an area that has since been the focus of much research into the phenomenon (Doughton, 2019). Although a range of possible explanations have been explored, including pathogens, herbivory by mountain beaver (*Aplodontia rufa*), and soil nutrient deficiencies (Kildisheva and Aghai, 2018), the sword fern decline remains unexplained.

Following the mortality event in Seward Park, the phenomenon appears to have spread outwardly through the sword fern population over multiple years. However, it does not appear to affect co-occurring plant species such as *Mahonia nervosa*, *Sambucus racemosa* and *Oemleria cerasiformis*. Although previous efforts using culture-based techniques and qPCR have been unable to identify a candidate organism (Kildisheva and Aghai, 2018), these observations nonetheless suggest that a host-specific pathogen may be responsible for the phenomenon. Two recent studies demonstrated that symptoms can be experimentally transmitted from symptomatic ferns to healthy ferns (Mendenhall et al., 2019; Shannon, 2020b). For example, Shannon (2020b) demonstrated the *in situ* transmission of symptoms after planting healthy sword ferns in a area of Seward Park where the phenomenon was actively spreading, resulting in decreased frond production after two years. These findings support the hypothesis that the phenomenon is driven by a pathogen.

However, symptoms of the phenomenon often resemble the effects of drought stress and can include crispate pinnae or stunted frond development. Some evidence suggests that local site conditions may influence the phenomenon by augmenting the available soil moisture. For example, in an area of Seward Park with high sword fern mortality, the proximity of sword ferns to large-diameter trees and coarse woody debris was correlated with a higher likelihood of survival (Shannon, 2019). These patterns could be partially explained by the mitigation of drought stress through hydraulic redistribution (Brooks et al., 2002) or common mycorrhizal networks shared with overstory trees (Teste et al., 2009). Another study found that soil bulk density was significantly higher in a die-off site in Seward Park (Masaoka, 2017), which may suggest that the water-holding capacity of the soil was limited due to soil compaction (Archer and Smith, 1972). Based on these findings, we consider the hypothesis that the phenomenon may be driven by seasonal drought stress, aggravated by local site conditions that influence soil moisture.

Our study explores these competing hypotheses by comparing areas of Seward Park containing dead and symptomatic ferns with areas containing predominantly healthy ferns. We included Schmitz Preserve Park as an additional control due to it being a remnant old-growth forest with a similar structure and composition as Seward Park. To evaluate the potential role of seasonal drought, we analyzed the foliar moisture content of the sword ferns and compared it with the moisture content and bulk density of the soil. To explore the pathogen hypothesis, we prepared tissue samples for highthroughput sequencing to screen the endophytic microbial communities of sword ferns, with a focus on bacteria, fungi, and oomycetes. The microbiome component of the study is ongoing.

This report provides a detailed description of our methods so they may be adapted for future research and monitoring. We also present some preliminary results from our analysis of the foliar moisture content and soil properties. Although symptomatic ferns show signs of drought stress in their foliage, the phenomenon cannot be explained solely by a deficit in soil moisture.



## <u>Methods</u>

#### Study Areas

Seward Park (47.556, -122.251) and Schmitz Preserve Park (47.574, -122.400), both located in Seattle, Washington, represent some of the few remaining low elevation old-growth forests in Puget Sound. Seward Park is located on the Bailey peninsula and is surrounded by Lake Washington. Although Seward Park encompasses an area of 121 ha (300 ac), its largest contiguous upland forest has an area of approximately 50 ha (120 ac). Schmitz Preserve Park is located on the Duwamish Peninsula, approximately 11 km (7 mi) to the west of Seward Park. With an area of 21 ha (53 ac), Schmitz Preserve Park represents the majority of a larger forest fragment that extends onto adjacent parcels, including private property, right-of-way, and other public property.

As these parks are both located in the City of Seattle, we assume them to have a nearly identical climate. Based on the 1981 – 2010 climate normals for Seattle (NOAA, 2021), the average annual precipitation is 96 cm (37.8 in) and the average annual temperature is 11.7 °C (53.0 °F), with a summer high of 23.4 °C (74.1 °F) and a winter low of 2.5 °C (36.5 °F). However, over the twentieth century, the climate of the Pacific Northwest has increased in temperature by 0.8 °C (1.5 °F), and models predict the annual temperature to increase by an additional 3.0 °C (5.3 °F) by the 2080s (Mote and Salathé, 2010). Although total annual precipitation is predicted to remain fairly stable, some models predict changes in its seasonal distribution, leading to wetter winters and drier summers (Mote and Salathé, 2010).

Located in the western hemlock (*Tsuga heterophylla*) zone of Western Washington (Franklin and Dyrness, 1988), the upland forests of Seward Park and Schmitz Preserve Park are similar to each other in their composition. The areas included in our survey are generally co-dominated by Douglas fir (*Pseudotsuga menziesii*), western red cedar (*Thuja plicata*) and/or western hemlock (*Tsuga heterophylla*). We also included mixed deciduous – conifer areas co-dominated by big leaf maple (*Acer macrophyllum*) and/or red alder (*Alnus rubra*). To a lesser extent, grand fir (*Abies grandis*), madrona (*Arbutus menziesii*), Pacific dogwood (*Cornus nuttallii*), shore pine (*Pinus contorta*), bitter cherry (*Prunus emarginata*), and Pacific yew (*Taxus brevifolia*) are also represented in these forests. Tree regeneration is dominated by western hemlock, western red cedar, and grand fir.

The understory communities in the surveyed areas vary in their composition. However, they are generally consistent with the TSHE-PSME/POMU-DREX, PSME-TSHE/VAOV/POMU, and PSME-TSHE/GASH/POMU plant associations, as defined by Chappell (2004). The understory is dominated by perennial shrubs and ferns, particularly sword fern (*Polystichum munitum*) and/or evergreen huckleberry (*Vaccinium ovatum*) and salal (*Gaultheria shallon*). Other frequently occurring species include beaked hazelnut (*Corylus cornuta*), spreading woodfern (*Dryopteris expansa*), oceanspray (*Holodiscus discolor*), dwarf Oregon grape (*Mahonia nervosa*), osoberry (*Oemleria cerasiformis*), baldhip rose (*Rosa gymnocarpa*), salmonberry (*Rubus spectabilis*), trailing blackberry (*Rubus ursinus*), red elderberry (*Sambucus racemosa*), and red huckleberry (*Vaccinium parvifolium*).

Seward Park and Schmitz Preserve Park contain an array of other distinct plant communities that were excluded from our survey, including forested wetlands, freshwater shoreline, shrubland, riparian corridors, Garry oak (*Quercus garryana*) forest, and mixed conifer – broadleaf evergreen forest (Hanson



and Davidson, 2005; Papanikolas, 1996). Seward Park also contains heavily landscaped areas maintained with non-native trees, shrubs or grass.

The forest structure of both parks is likely influenced by the historical resource management of the Duwamish Tribe. Both forests are located in close proximity to the sites of historical prairies (Thrush, 2007) which were maintained using prescribed burning to enhance food productivity and to prevent the encroachment of shrubs and trees (Turner and Peacock, 2005). Repeated disturbance by non-stand replacing fires may explain why both forests appear to contain multiple age-cohorts of Douglas fir, a pioneer species that regenerates in the open canopy conditions created by moderate-severity fires (Dunn et al., 2020). The presence of fire scars on the bark of large-diameter trees and stumps provides further evidence of the historical role of fire in these forests.

As a notable difference between the two parks, Schmitz Preserve Park shows evidence that some of the large-diameter Douglas fir and western red cedar trees were selectively harvested by Euro-American settlers in isolated areas of the forest. These harvests likely occurred prior to 1908, when the largest parcel was first established as a park (Sherwood, n.d.). In Seward Park, the upland forest does not show evidence of industrial logging, although some firewood cutting and removal of windthrown trees may have occurred during the 1930s (Hanson and Davidson, 2005).

The soils also differ between Seward Park and Schmitz Prserve Park. Based on our observations of the soil texture in the upper horizons, the upland areas of Schmitz Preserve Park are consistent with the Indianola series, which is characterized by deep loamy sands that developed from glacial outwash (Soil Survey Staff, 2009). The forested wetland areas of Schmitz Preserve Park, which were not sampled, have clayey soils that likely developed from glacial lacustrine deposits (Sherwood, n.d.), consistent with the Kitsap series (Soil Survey Staff, 2018a). In contrast, the soils of Seward Park are predominantly silty loams and gravelly loams that developed from glacial till (Hanson and Davidson, 2005). As the lake that surrounds Seward Park was formed by a receding glacier (Sherwood, 1973), lacustrine deposits may have also influenced the parent materials, which would explain the more finely textured soils that we encountered in some areas. We speculate that the soils of Seward Park may represent an intergrade between the Alderwood and Kitsap series (Soil Survey Staff, 2018a, 2018b), which are strongly associated with each other in King County (Snyder et al., 1973).

#### **Experimental Design**

#### Approach

Our study compared forested areas of Seward Park containing sword ferns (*Polystichum munitum*) that were predominantly healthy or predominantly symptomatic. We included upland forested areas of Schmitz Preserve Park containing healthy sword ferns as an additional control due to similarities in the structure and composition of their plant communities, as well as their close proximity to each other. Together, these sites represent three experimental groups: i) Schmitz–Healthy, ii) Seward–Healthy, and iii) Seward–Symptomatic (Figure 1). We considered using a blocked design, however ferns with symptoms consistent with the phenomenon were generally not observed in Schmitz Preserve Park. In each experimental group, we established 12 square plots, each with an area of 100 m<sup>2</sup> (0.025 acres).



#### Park



Figure 2: Experimental design.

#### Site Selection

Initially, we had intended on delineating a large area containing healthy sword ferns in Seward Park and comparing it with another area containing predominately symptomatic and/or dead ferns. However, during our preliminary reconnaissance, we observed dead and symptomatic ferns in nearly all areas of the park. The phenomenon has become so extensive in Seward Park that there is no longer a contiguous area with healthy sword ferns large enough for a survey like ours. Therefore, we adapted our protocol by developing a set of criteria for defining healthy and symptomatic areas.

Symptomatic plots were randomly placed in forested areas of Seward Park where: i) the understory plant community is dominated by native plants; ii) living sword ferns are present; and iii) more than 50% of the sword ferns have symptoms or have recently died. We defined symptomatic ferns as those with any combination of the following criteria: i) crispate and/or contorted pinnae; ii) pinnae with atypical hues; iii) fronds with aborted or undeveloped pinnae at the distal end; iv) fronds with partial die-back at the distal end; v) fronds with a contorted rachis; and/or vi) a ratio of young fronds to old fronds less than 0.5 (Figure 9). We defined healthy ferns as lacking the above symptoms.

Healthy plots were randomly placed in forested areas of Seward Park and Schmitz Preserve park where: i) the understory plant community is dominated by native plants; ii) living sword ferns are present; iii) more than 50% of the sword ferns are healthy; and iv) less than 25% of the sword ferns have recently died. As an exception, plot 21 was mistakenly placed in an area where 28% of the sword ferns were estimated to have recently died.

Thus, our plots in Seward Park are best interpreted as representing fragmented patches of healthy sword ferns, surrounded by a matrix of dead and symptomatic ferns.



#### Constraints

Our survey should not be construed as being proportionally representative of the plant communities in these parks. As a result of our strict criteria for comparing healthy and symptomatic plots, our survey omitted non-forested areas of the parks, areas dominated by non-native species, areas where living sword ferns are no longer present due to the die-off phenomenon (Figure 11), and areas where sword ferns were not a component of the understory in recent history. Our criteria also resulted in the omission of areas with ambiguous or intermediate effects of the phenomenon, such as isolated patches of healthy ferns in areas with high mortality or areas with low mortality and ambiguous symptoms. As additional constraints, we avoided 6 long-term monitoring plots established by EarthCorps and 20 long-term monitoring plots established by researchers at the University of Washington. For safety reasons, we avoided overhead hazards and areas with slopes over 100%. We also avoided sampling within 10 m of the trails due to the ongoing pandemic.

#### **Field Survey**

Plots were surveyed by Dylan Mendenhall and Andres Barrera between August 27, 2020 and September 10, 2020. At every plot, we recorded the survey date, latitude, longitude, slope, slope position, aspect, and canopy cover. We used a modified Braun-Blanquet cover-abundance scale to visually estimate the abundance of each vascular plant species observed within 10 m of the ground (Braun-Blanquet et al., 1932; Mueller-Dombois and Ellenberg, 1974). Trees and non-vascular plants were not included in the cover-abundance survey. The mid-point values of the cover classes were used in subsequent analysis. We measured the diameter at breast height (i.e. bole diameter at 1.4 m above the ground) on the uphill side of every tree. Trees below 1.4 m in height (i.e. "seedlings") were rarely observed and were not included in the survey.

Cover Class	Cover Range (%)	Cover Midpoint (%)
А	0 - 1	0.5
В	1 - 5	3
С	5 - 15	10
D	15 - 25	20
Е	25 - 50	37.5
F	50 - 75	62.5
G	75 - 95	85
Н	95 - 100	97.5

Table 1: Modified Braun-Blanquet cover-abundance scale.



#### Sample Collection and Preparation

In each plot, a single sword fern was selected for tissue sampling using the criteria described above for distinguishing healthy and symptomatic ferns. Prior to sampling, we photographed the fern and noted general observations about its health.

We counted the total number of young and old fronds as a measure of the fern's recent productivity. We defined young fronds as those that were produced during the most recent spring and summer, and old fronds as those that were produced during any previous year. We used morphological, developmental, and physiological traits to determine their relative age. Sword ferns generally produce fronds in discrete age-cohorts, with a new set of fronds produced each year. Each set of fronds is generally retained for 2 to 3 years. As fronds only produce sori during their first year aboveground, we primarily used the presence of sori on the underside of the pinnae for identifying young fronds. We also interpreted the relative age of the fronds based on the abundance and orientation of the scales at the base of their stipes. During their first spring and summer aboveground, the stipes are covered by a dense layer of scales. Over time, the scales become more appressed to the stipe and gradually detach, presumably through senescence, abrasive action, and rainfall. We interpreted fronds with more sparsely covered stipes and more appressed scales as having been produced during a previous year.

From each fern, we collected samples from the roots, rhizome, stipe, and pinnae. We collected 20 random pinnae and placed them in non-sterile plastic bags. From a random frond, we collected a single section of the stipe, approximately 2 cm in length, and immediately placed it in a sterile microcentrifuge tube. We then excavated one of the rhizomes. Mature sword ferns are almost universally comprised of rhizomes with multiple branches. Thus, we were generally able to sample a single branch while leaving the rest of the plant undisturbed. We cut an approximately 10 cm section from the terminal end of the rhizome and placed it in a non-sterile plastic bag. We collected approximately 1 g of the root tissue, which was immediately placed in a sterile microcentrifuge tube. Tissue samples were stored at 4°C for approximately 24 hours before being transported to the lab.

Soil samples were collected from the uppermost B horizon, approximately 5 to 10 cm in depth. For the soil texture sample, we aggregated five 200 ml sub-samples from random locations in the plot. From this aggregate sample, 200 ml was placed in a non-sterile plastic bag for subsequent analysis. We also collected a single bulk density sample using a core with a volume of 78.5 cm<sup>3</sup>, which was immediately placed in a sealed tin.

#### Laboratory Analysis

#### Foliar Moisture Content and Soil Properties

The soil texture samples were sieved to 2 mm and air-dried for long-term storage. Analysis of the soil texture is ongoing. Bulk density samples were oven-dried at 105°C to a constant weight. Gravimetric moisture content was determined using the bulk density samples. To determine foliar moisture content, 10 random pinnae from each fern were oven-dried at 95°C to a constant weight.



#### **High-Throughput Sequencing**

The stipe and root samples were stored in a freezer at -20°C without additional preparation. The rhizome samples and the remaining pinnae were stored at 4°C for another 2 weeks before further processing. Surface sterilization of the pinnae followed the methods of Younginger and Ballhorn (2017). Pinnae were then placed in sterile centrifuge tubes and stored in a freezer at -20°C.

Rhizomes were dissected in two stages. On an open bench, we used a razor to remove the remaining soil, stipes, and the outer layer of the rhizome. We removed a section approximately 5 cm long from the terminal end of the rhizome and transferred it to a laminar flow hood for further preparation. In the second stage, the rhizome was cut into smaller sections approximately 2 cm in diameter. The rhizome sections were surface sterilized in the same manner as the pinnae (Younginger and Ballhorn, 2017) with a final rinse in deionized water. The rhizome sections were then cut into small pieces approximately 5 mm in diameter and then aggregated together. Of these aggregated pieces, approximately 4 g were placed in a sterile centrifuge tube and stored in a freezer at -20°C until further use.

All 36 rhizome samples and 4 pinnae samples (#25, #28, #33, #34) were shipped on dry ice to MPG Ranch for DNA extraction, PCR, and MiSeq library preparation. These analyses are ongoing.

#### Statistical analysis

In our preliminary review of the data, we used analysis of variance (ANOVA) followed by a Tukey HSD test to compare the foliar moisture content, soil moisture content, and soil bulk density of the three experimental groups. We used Pearson's correlation coefficient to compare soil moisture content with foliar moisture content. For all statistical tests, we used an alpha level of 0.05. All analyses were conducted using the R programming language (R Core Team, 2019).



## <u>Results</u>

#### Anecdotal Observations

In Seward Park, we observed a wide range of morphological, developmental, and physiological traits associated with the die-off phenomenon, including: ferns with a low ratio of young to old fronds; fronds with a contorted rachis; fronds with crispate or contorted pinnae; fronds with an atypical hue; fronds that had not fully expanded at their distal end; and fronds that had partial die-back at their distal end after fully developing (Figure 9).

Some of the symptomatic ferns showed clear evidence of a gradual decline over multiple years. For example, we inferred that fern #33 produced an apparently healthy set of fronds in 2018, a set of discolored fronds in 2019, and no new fronds in 2020 (Figure 3).



*Figure 3: Example of a symptomatic fern (#33) showing evidence of it declining over multiple years. Arrows indicate two distinct cohorts of fronds that presumably emerged in 2018 and 2019. No fronds were produced in 2020.* 



To the best of our knowledge, there is no evidence of widespread mortality among the sword fern population in Schmitz Preserve Park. We observed only one sword fern that had recently died (plot #4), which was readily explained by competition for soil moisture with a large diameter tree growing in close proximity. All other sword ferns in the plot appeared healthy.

On a small subset of the sampled ferns, we conducted an impromptu dissection in the field by cutting a longitude section through a rhizome. We found a set of fully developed leaf buds clustered around the apical meristem of the rhizome of all healthy ferns. In contrast, the symptomatic ferns generally had no leaf buds present, or in some cases, leaf buds that appeared to have withered after their initial development. We confirmed this observation in the lab, where we made longitudinal sections of individual leaf buds. Leaf buds from healthy ferns had a fully developed spiral shape. In contrast, leaf buds from symptomatic ferns failed to develop past their primordial stage or appeared to have withered away after initially developing into a spiral shape.

As another pertinent observation, we found an Anderson's fern (*Polystichum andersonii*) in an area of Seward Park with over 50% mortality among sword ferns (*Polystichum munitum*). Despite being adjacent to several dead and symptomatic sword ferns, the Anderson's fern appeared relatively healthy. Its leaves did not have any symptoms associated with the die-off phenomenon, and there was a set of fully developed leaf buds at the base of its fronds.

#### **Preliminary Analysis**

Mean foliar moisture content (FMC) of the pinnae was significantly different between the three experimental groups based on ANOVA ( $F_{2,33} = 27.52$ , p < 0.001). A post-hoc Tukey HSD test revealed the mean FMC of the pinnae was significantly lower in the symptomatic ferns at Seward Park (0.98 g·g<sup>-1</sup> ± 0.11 SE) compared with the healthy ferns at both Seward Park (1.67 g·g<sup>-1</sup> ± 0.07 SE, p < 0.001) and Schmitz Preserve Park (2.11 g·g<sup>-1</sup> ± 0.14 SE, p < 0.001). The FMC of the healthy ferns in Seward Park was also significantly lower than the healthy ferns in Schmitz Preserve Park (p = 0.039).

The soil bulk density was significantly different between the experimental groups ( $F_{2,33} = 5.122$ , p = 0.012). The mean bulk density in the Schmitz–Healthy group was 0.96 g·g<sup>-1</sup> ± 0.04 SE, significantly lower than the bulk density of the Seward–Healthy group (1.11 g·g<sup>-1</sup> ± 0.04 SE, p = 0.081) and the Seward–Symptomatic group (1.17 g·g<sup>-1</sup> ± 0.03 SE, p = 0.011). However, the bulk density of the Seward–Healthy and Seward–Symptomatic groups were similar to each other (p = 0.658).

The gravimetric soil moisture content was similar between all three experimental groups ( $F_{2,33} = 0.663$ , p = 0.522), with means ranging from 0.10 g·g<sup>-1</sup> to 0.12 g·g<sup>-1</sup>. Furthermore, the foliar moisture content of the ferns was not correlated with soil moisture content (r = 0.07, p = 0.675).





*Figure 4: Longitudinal sections through the rhizomes of a healthy fern (left) and a symptomatic fern (right). Arrow indicates leaf buds that failed to develop on the symptomatic fern. Bars represent 5 cm.* 



*Figure 5: Longitudinal sections through the dormant leaf buds of a healthy fern (left) and a symptomatic fern (right). Bars represent 2 cm.* 



*Figure 6: Polystichum andersonii. This fern appeared relatively healthy despite being in close proximity to several dead and symptomatic sword ferns (Polystichum munitum). Arrow indicates a dead sword fern.* 



Figure 7: Foliar moisture content  $(g \cdot g^{-1})$  of sampled Polystichum munitum ferns. Data are displayed as Tukey-style boxplots; horizontal bar represents the median value; box represents the interquartile range (IQR); vertical lines represent minimum and maximum values up to 1.5 x IQR; points represent outliers.



Figure 8: Soil bulk density ( $g \cdot cm^{-3}$ ). Data are displayed as Tukey-style boxplots; horizontal bar represents the median value; box represents the interquartile range (IQR); vertical lines represent minimum and maximum values up to 1.5 x IQR; points represent outliers.



### **Discussion**

Under the drought hypothesis, we predicted that differences in foliar moisture content (FMC) would be correlated with differences in soil moisture content. We expected these patterns to be particularly evident during the dry season when we collected our samples. During this season, the chlorophyll fluorescence values ( $F_v/F_m$ ) of sword ferns decline in parallel with lower soil moisture content (Kildisheva et al., 2018). These changes in chlorophyll fluorescence are likely an indication of water stress (Maxwell and Johnson, 2000). In our study, FMC was indeed lower in symptomatic ferns compared with healthy ferns. However, this pattern was not correlated with differences in soil moisture content, which was similar between the three experimental groups. These results do not support the drought hypothesis, which is unsurprising given that sword ferns are well-adapted to surviving extended periods of drought (Baer et al., 2016).

Our results are consistent with the findings of two previous studies. In a greenhouse experiment, cut fronds that were exposed to symptomatic tissue developed significantly lower FMC compared to controls exposed to healthy sword fern tissue (Mendenhall et al., 2019), which suggests that a transmissible agent obstructed the vascular tissue. Likewise, in a field study based in Seward Park, symptomatic ferns had significantly lower midday xylem water potential compared with healthy ferns found elsewhere in the park (Goldstein-Miller et al., 2019). These patterns could potentially be explained by xylem cavitation or tylose formation induced by a vascular pathogen, which can result in symptoms that are similar in appearance to drought stress (Yadeta and Thomma, 2013).

The effect of this phenomenon on the FMC of *Polystichum munitum* has potential implications for wildfire management in the Western Cascades. The length of the active burning season and the behavior of wildfires are partially influenced by the foliar moisture content of understory plants (Agee et al., 2002). For example, the back rate of fire spread is controlled by the moisture content of surface fuels (Fernandes et al., 2009; Rossa, 2017). In the current study, many of the symptomatic ferns had FMC values well below 1.00 g·g<sup>-1</sup> (100%), a critical threshold for wildfire behavior in shrublands and forests with Mediterranean climates (Nolan et al., 2020, 2016). For areas affected by the sword fern die-off phenomenon, predictive models of fuel loading may need to be adjusted to account for unusually low moisture content in surface fuels during late summer and early autumn.

In our study, the soils associated with symptomatic ferns and healthy ferns in Seward Park had a similar bulk density, which conflicts with the results of a previous study in which soil bulk density was found to be significantly higher in an area of Seward Park affected by the phenomenon (Masaoka, 2017). This discrepancy is likely due to differences in our sampling approach. Masaoka (2017) focused on sampling soils from a relatively confined area of Seward Park where a mortality event occurred in 2013 (i.e. "ground zero"). In our study, the plots were dispersed over a wide area of Seward Park and are therefore more representative of the phenomenon. Nonetheless, we agree with Masaoka's overall interpretation; soils with bulk densities in the range observed are unlikely to restrict plant growth. Although we found significant differences in bulk density between Schmitz Preserve Park and Seward Park, these results are to be expected given that the soils developed from different parent materials (Soil Survey Staff, 2018b, 2009).

Because the soils of Schmitz Preserve Park are more coarsely textured, and because the bulk density is significantly lower, we would expect Schmitz Preserve Park to have better drainage than in Seward Park.



As a result, the effects of seasonal drought stress should be more pronounced in the upland areas of Schmitz Preserve Park compared with Seward Park. However, the FMC of the Schmitz–Healthy group was significantly higher than the Seward–Healthy group, despite the soil moisture content being similar. This observation leads us to believe that the die-off phenomenon may be so widespread in Seward Park that visual assessments may not be entirely reliable for identifying "healthy" ferns, except in terms of their relative differences. Future studies would be cautioned against using any areas of Seward Park as a reference site for healthy sword ferns.

Our most consequential finding may be the accidental discovery of a lone *Polystichum andersonii* fern in an area of Seward Park where the die-off phenomenon was especially severe. *Polystichum andersonii* is thought to be a tetraploid hybrid between *P. munitum* and *P. kwakiutlii* (Hitchcock and Cronquish, 2018). As these species are closely related to each other, this observation may suggest that the purported pathogen is host-specific to *Polystichum munitum*. Future research is needed to evaluate the susceptibility of other *Polystichum* species that are native to the Pacific Northwest, such as *P. imbricans* and *P. californicum*. Identifying a resistant *Polystichum* species would provide an option for partially restoring the understory structure in areas affected by the phenomenon.

Lastly, it was clear from our observations in Schmitz Preserve Park that mature stands of sword fern have a low mortality rate under normal circumstances. Therefore, the rapid decline of sword ferns in Seward Park cannot be attributed to the typical succession of old growth forests in our region.

### **Conclusions**

The sword fern decline at Seward Park is far from an isolated phenomenon; new die-offs continue to be observed at other locations throughout Western Washington. Although our study is ongoing, our preliminary analysis explored the hypothesis that the phenomenon is primarily caused by drought stress. More specifically, we predicted that foliar moisture content (FMC) would be lower in symptomatic ferns and that these differences would correlate with soil moisture content. We also expected differences in soil bulk density to be associated with the phenomenon by moderating the soil water holding capacity.

Although we did find significantly lower FMC in the symptomatic ferns, these differences could not be explained by differences in soil moisture content or soil bulk density. Our results suggest that the phenomenon is driven by a factor that is independent of soil water availability. Our preliminary results are consistent with the pathogen hypothesis.

Our study raises concerns about the potential effects of the phenomenon on wildfire behavior, particularly in areas of the Olympic Peninsula and the Western Cascades where the understory is dominated by *Polystichum munitum*. As another implication of our research, we identified *Polystichum andersonii* as a candidate species for restoring the understory structure of forests affected by the sword fern die-off phenomenon. However, further research is needed to confirm its resistance.

In the next stage of our study, we will use a metabarcoding approach to identify potential bacteria, fungi, or oomycetes that may be responsible for the phenomenon, building on previous research that identified fungal endophytes in the pinnae of sword ferns (Younginger and Ballhorn, 2017). Further



research would be needed to understand how the pathogen is transmitted and its mechanism for inducing water stress, among other symptoms. As a potential limitation, our amplicon sequencing approach will not be able to identify RNA viruses, which are known to affect other fern species (Valverde and Sabanadzovic, 2009). It also remains possible that the phenomenon is caused by changes in the life cycle or behavior of an invertebrate vector rather than through the introduction of a pathogen.

In the absence of a discrete die-off event, the phenomenon is difficult to detect, even while it continues to spread through the understory. Many of the symptoms are identical to the temporary effects of seasonal drought. However, repeated observations at the same location can reveal the rapid decline of sword ferns over multiple years. We recommend that land managers establish monitoring programs to evaluate potential changes in their local sword fern populations and to prioritize areas for management.

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### <u>Appendix</u>



Figure 9: Signs and symptoms of the sword fern die-off phenomenon: A) reduced production of new fronds; B) contorted rachis; C) crispate and contorted pinnae; D) discolored pinnae; E) underdeveloped and aborted fronds; F) die-back at distal end of a frond. In areas affected by the phenomenon, these signs and symptoms can occur in any combination and vary in severity.





Figure 10: Example of an individual sword fern that died in Upper Luther Burbank Park on Mercer Island. Some of the nearby sword ferns did not have symptoms, despite their close proximity.



Figure 11: Example of an area in Seward Park previously dominated by sword ferns. Areas with 100% mortality could not be included in our survey. Arrows indicate the remnants of dead sword ferns.



Figure 12: Frond from an apparently healthy fern (left) compared with a frond from a symptomatic fern (right) in Upper Luther Burbank Park on Mercer Island.