

Tennessee State University

Digital Scholarship @ Tennessee State University

Agricultural and Environmental Sciences
Faculty Research

Department of Agricultural and Environmental
Sciences

5-20-2022

Efficacy and Timing of Application of Fungicides, Biofungicides, Host-Plant Defense Inducers, and Fertilizer to Control Phytophthora Root Rot of Flowering Dogwoods in Simulated Flooding Conditions in Container Production

Krishna Neupane

Bhawana Ghimire

Fulya Baysal-Gurel

Follow this and additional works at: <https://digitalscholarship.tnstate.edu/agricultural-and-environmental-sciences-faculty>



Part of the [Plant Pathology Commons](#)

1 **Efficacy and Timing of Application of Fungicides, Biofungicides, Host-Plant**
2 **Defense Inducers, and Fertilizer to Control Phytophthora Root Rot of**
3 **Flowering Dogwoods in Simulated Flooding Conditions in Container**
4 **Production**

5

6 Krishna Neupane, Bhawana Ghimire and Fulya Baysal-Gurel*

7

8

9

10 Department of Agricultural and Environmental Sciences, College of Agriculture, Tennessee State
11 University, Otis L. Floyd Nursery Research Center, 472 Cadillac Lane, McMinnville, TN, USA.

12 *Corresponding author: F. Baysal-Gurel, email: fbaysalg@tnstate.edu

13

14

15

16

17

18

19

20

21

22

23

24

25

26

27

28 **Abstract**

29 Phytophthora root rot caused by *Phytophthora cinnamomi* Rands is one of the major diseases of
30 flowering dogwoods (*Cornus florida* L.). The severity of root rot disease increases when the
31 plants are exposed to flooding conditions. A study was conducted to determine the efficacy and
32 timing of application of different fungicides, biofungicides, host plant defense inducers, and
33 fertilizer to manage Phytophthora root rot in month-old seedlings in simulated flooding events
34 for 1-, 3-, and 7- days. Preventative treatments were drench applied 3 weeks and 1 week before
35 flooding whereas curative treatments were applied 24 hrs. after flooding. Dogwood seedlings
36 were inoculated with *P. cinnamomi* 3 days before the flooding. Plant height and width were
37 recorded at the beginning and end of the study. At the end of the study, plant total weight and
38 root weight were recorded and disease severity in the root was assessed using a scale of 0-100%.
39 Root samples were plated using PARPH-V8 medium to determine the percentage recovery of the
40 pathogen. Empress Intrinsic, Pageant Intrinsic, Segovis, and Subdue MAXX, as preventative and
41 curative applications, were able to suppress the disease severity compared to the inoculated
42 control in all flooding durations. All treatments, with the exception of Stargus as preventative
43 application 3 weeks before flooding and Orkestra Intrinsic as curative application, were able to
44 suppress the disease severity compared to the inoculated control for 1-day flooding event. Aliette
45 and ON-Gard were effective in the first trial when applied preventatively in both 1 week and 3
46 weeks before flooding but not in the second trial. Signature Xtra was effective as preventative
47 application but not as a curative application. Interface was effective as curative application but
48 not as preventative application. The findings of this study will help nursery growers to
49 understand the performance of fungicides, biofungicides, host-plant defense inducers, and
50 fertilizer in different time intervals and repeated applications to manage Phytophthora root rot in
51 flooding conditions.

52

53

54

55

56 **Keywords:** biological control, chemical control, *Phytophthora cinnamomi*, soilborne pathogen,
57 woody ornamental

58 **Introduction**

59 Phytophthora root rot caused by *Phytophthora cinnamomi* Rands (family Peronosporaceae) is
60 one of the most important diseases of the ornamental crops (Brown et al. 2019a,b; Duan et al.
61 2008; Ferguson and Jeffers 1999; Hu et al. 2010; Neupane et al. 2021). *Phytophthora* causes
62 root rot diseases in more than 5,000 species of woody plants worldwide (Balci et al. 2007;
63 Erwin and Ribeiro 1996; Hardham and Blackman 2018; Shakya et al. 2021). The infestation of
64 *P. cinnamomi* has caused the extinction of a few native species of forest plants in Australia and
65 has posed a threat in the United States (Hardham 2005; Hu et al. 2010). Among the woody
66 ornamental crops, the major host for the notorious *P. cinnamomi* oomycete includes *Acer* spp.,
67 *Cornus* spp., *Juglans* spp., *Prunus* spp., *Rhododendron* spp. and *Quercus* spp. (Dai et al. 2020;
68 Erwin and Ribeiro 1996; Ribeiro and Linderman 1991; Zentmyer 1980). The economic and
69 scientific importance of this pathogen can be estimated as it has been ranked in the top 10
70 oomycete plant pathogens of the world (Kamoun et al. 2015) and addressed as a “key threatening
71 process to Australia’s biodiversity” in the Environmental Protection and Biodiversity
72 Conservation Act 1999 (Shakya et al. 2021). *P. cinnamomi* causes rotting of the fine and fibrous
73 roots limiting the supply of food and water to the shoot region. As a result, the plant suffers
74 wilting, dieback, and stem cankers (Hardham 2005; Hardham and Blackman 2018). It also
75 causes gradual tree decline, large basal cankers with bleeding spots, root rot, and death of the
76 forest trees (Robin et al. 2012).

77 Flooding has a direct impact on the severity of Phytophthora root rot. The susceptibility
78 of the plant to the pathogen increases when exposed to flooding (de Silva et al. 1999; Jacobs and
79 Johnson 1996). Trees infected with the root rot pathogen decline faster when they are exposed to
80 flooding conditions (Brown et al. 2019b; Ploetz and Schaffer 1989; Ploetz and Schaffer 1987;
81 Reeksting et al. 2016). In some cases, flooding becomes the only reason for destruction of trees
82 which otherwise would have survived with just a *P. cinnamomi* infection (Weste and Marks
83 1987). The destructive potentiality of the *Phytophthora* pathogens increases with flooding
84 because flooding creates favorable conditions for infection such as limited oxygen and increased
85 zoospores movement (Kawase 1981; Kozlowski 1997; Wilcox and Mircetich 1985). In a study
86 conducted by Brown et al. (2019b), there were 1.7%, 15.7%, and 34.8% increases in the

87 mortality of flowering dogwood seedlings for 1 day, 3 days, and 7 days of simulated flooding
88 conditions, respectively.

89 Management of Phytophthora root rot is challenging not only because of the wide host
90 range of the pathogen but for various other reasons: a) higher production of chlamydozoospores
91 (Zentmyer 1980), b) unexpressed pathogens in contaminated plants and growth media (Benson
92 and Grand 2000; Hardham and Blackman 2018; Kong et al. 2003), and c) high dispersal of
93 pathogen either by zoospores or running water (Duniway 1976; Erwin and Ribeiro 1996).

94 Different cultural, biological, and chemical practices have been applied for the
95 management of Phytophthora root rot (Baysal-Gurel et al. 2016; Brown et al. 2019b; Gould
96 2012; Neupane et al. 2021; Parajuli et al. 2021; Williams-Woodward and DeMott 2013).
97 However, most of these practices have been inconsistent. The application of mulches, composts,
98 and animal manure was found to restrict the growth of *P. cinnamomi* (Aryantha and Guest 2006;
99 Evidente et al. 2009; Richter et al. 2011). Hu et al. (2010) claimed that chemical control is the
100 most effective strategy to manage *P. cinnamomi* among other practices being used. Application
101 of fungicides like mefenoxam and fosetyl-Al have been effective in preventing the development
102 and spread of the pathogen Neupane et al. 2021; Ribeiro and Linderman 1991; Weiland et al.
103 2021). In the studies conducted by Benson and Blazich (1989) and Hu et al. (2010), systemic
104 mefenoxam alone successfully managed Phytophthora root rot in *Azalea*, *Rhododendron*, *Ilex*,
105 *Myrica*, *Berberis*, *Kalmia*, *Heliamphora*, *Rhododendron*, and *Viburnum*, respectively. Fosetyl-Al
106 alone, acibenzolar-S-methyl, and pyaclostrobin with boscalid were also able to manage *P.*
107 *cinnamomi* in avocado, redbuds, and tulip poplar trees (Addresso et al. 2018; Benson 1985;
108 Bezuidenhout et al. 1987).

109 Biofungicides and fertilizers were also found effective in the management of
110 Phytophthora root rot. *Trichoderma*, *Pseudomonas*, *Bacillus*, *Streptomyces*, and *Enterobacter*
111 resulted in increased plant resistance to disease and enhanced growth and productivity (Belimov
112 et al. 2001; Harman 2006; Harman et al. 2004; Pieterse et al. 2000; Pieterse et al. 2001; Wang
113 et al. 2000). Rootshield Plus (*Trichoderma harzianum* strain T-22, *T. virens* strain G-41) and
114 Stargus (*Bacillus amyloliquefaciens* F727) reduced the disease severity of *P. cinnamomi* in
115 flowering dogwoods for 1-day flooding when used preventatively (Brown et al. 2019b). Biogen
116 (*Azotobacter vinaudit* + *A. chroococum*), Nitrobein (*A. chroococum* + *Azospirillum lipoferum*),

117 and Bioarc (*B. megaterium*) were found to be effective against Phytophthora root rot of apple
118 and citrus (Sharkawy et al. 2014). Host plant defense inducers, strobilurin, paclobutrazol, and
119 triazole fungicides also helped in effective management of *P. cinnamomi* by increasing the
120 immune system of the plants against flooding, drought, extreme temperatures, and pollutants
121 (Han et al. 2012; Lin et al. 2006; Neupane et al. 2021).

122 Different fungicides, biofungicides, and host-plant defense inducers have been used for
123 the management of Phytophthora root rot but most of them have been inconsistent when used in
124 flooding conditions. Many research studies have already demonstrated that flooding exacerbates
125 the root rot infection caused by *Phytophthora*. In this study, selected products were used as two
126 different preventative applications (1 week and 3 weeks before flooding) and at least two or
127 more curative applications on flowering dogwood seedlings in simulated flooding conditions of
128 1-, 3- and 7- days to assess their effectiveness in successful management of Phytophthora root
129 rot. The major objective of this study is to evaluate products to provide potential management
130 strategies to protect woody ornamental crops against *P. cinnamomi* exposed to the flooding
131 conditions. This study assesses the performance of different fungicides, biofungicides, host-plant
132 defense inducers, and fertilizer with different application intervals or repeated applications to
133 combat Phytophthora root rot in flooding conditions.

134 **Materials and Methods**

135 **Study location and growing conditions:** This greenhouse study was conducted at Otis L. Floyd
136 Nursery Research Center, Tennessee State University, McMinnville, Tennessee. One-year old
137 seeds of flowering dogwoods ‘Cherokee Princess’ were stratified with the potting substrate
138 (Morton’s Grow Mix #2: Canadian sphagnum peat [60%], vermiculite [20%], and perlite [20%],
139 average substrate bulk density 144 kg/m³) (Morton’s Horticultural Products, McMinnville, TN,
140 USA) in a clear zip lock bag starting from August of 2019 (trial 1) and September of 2020 (trial
141 2) for four and three months, respectively (Reed 2005). As the seeds broke their dormancy and
142 radicles emerged out, they were transferred to the No. 72 tray (Morton’s Horticultural Products)
143 on 6 January 2020 (trial 1) and 10 January 2021 (trial 2) using the same potting substrate used
144 for stratification. Overhead irrigation was provided daily for 1 min twice a day. The seedlings
145 were transplanted to 10.2 cm pots containing potting substrate (Morton’s Nursery Mix:
146 processed pine bark [55%-65%], Canadian sphagnum peat and sand) (Morton’s Horticultural

147 Products) on 11 February 2020 (trial 1) and 21 February 2021 (trial 2). Seedlings were irrigated
148 daily for 2 min throughout the experimental period.

149 **Treatments:** A total of thirteen treatments (7 fungicides, 3 host plant defense inducers, 2
150 biofungicides, and 1 fertilizer) were used with two controls (inoculated and non-inoculated) as a
151 preventative application to assess their effectiveness against *P. cinnamomi* on flowering
152 dogwood seedlings in simulated flooding conditions. For the curative application, only nine
153 treatments (7 fungicides and 2 host plant defense inducers) were used with two controls similar
154 to the preventative application. The list of the treatments used in the study is provided in Table 1.
155 All treatments were drench applied in both preventative and curative applications. The
156 preventative application of the treatments was categorized into two different timings: 3 weeks
157 before the flooding and 1 week before the flooding. Preventative application of treatments 3
158 weeks before flooding was carried out on 2 April 2020 (trial 1) and 12 April 2021 (trial 2).
159 Preventative application 1 week before flooding was carried out on 16 April 2020 (trial 1) and 26
160 April 2021 (trial 2). Curative treatments were applied 24 hrs. after flooding for all 1-, 3-, and 7-
161 days. The first application of curative treatments for 1-day flooding was 25 April 2020 (trial 1)
162 and 5 May 2021 (trial 2), 3-days flooding was 27 April 2020 (trial 1) and 7 May 2021 (trial 2),
163 and 7 days flooding was 1 May 2020 (trial 1) and 11 May 2021 (trial 2). All curative applications
164 were made at least twice or more following standard protocol from the manufacturing company,
165 and their application interval is given in Table 1. TRIACT® 70 (clarified hydrophobic extract of
166 neem oil (70%), OHP, Inc., Mainland, PA, USA) was sprayed over the seedlings in trial 2 on 26
167 March and 2, 9, and 16 April 2021 to suppress the infection of powdery mildew. Powdery
168 mildew symptoms were not observed in the seedlings during trial 1.

169 **Pathogen inoculation and flooding:** Inoculum of the pathogen was prepared exactly two weeks
170 before the inoculation date using the isolate FBG201510 of *P. cinnamomi* (GenBank accession
171 no. MK099813) following standard protocol (Brown et al. 2019b; Neupane et al. 2021). For the
172 first trial, flowering dogwood seedlings were inoculated on 20 April 2020, and for the second
173 trial on 30 April 2021. After 3 days of pathogen inoculation, flooding was carried out. Flooding
174 was simulated by placing the pots containing seedlings in a zip-lock bag filled with water.
175 Flooding was done on the same day, 23 April 2020 (trial 1) and 3 May 2021 (trial 2) for all

176 seedlings from preventative and curative treatments. All the flowering dogwood seedlings were
177 inoculated with *P. cinnamomi* except for the non-inoculated treatments.

178 **Data recording:** Data were recorded on plant growth, severity of Phytophthora root rot, and
179 percentage recovery of the pathogen from the root samples. Plant growth data including plant
180 height and plant width was recorded at the beginning and end of the study. Total fresh plant
181 weight and total fresh root weight were recorded at the end of the study. Disease severity was
182 assessed after the completion of study by removing the soil and washing the roots. Disease
183 severity was ranked by visual observation using a scale of 0-100% based on the percentage of
184 root with symptoms where 0% = healthy and 100% = completely damaged.

185 For the percentage recovery of the pathogen, ten randomly selected root cuttings of
186 approx. 15 mm were plated on PARPH-V8 medium. After 7 days post plating, the plated roots
187 were assessed to determine the recovery of the pathogen exhibiting growth patterns similar to *P.*
188 *cinnamomi*. PARPH-V8 media was prepared following the standard protocol (Ferguson and
189 Jeffers 1999).

190 *Trichoderma* colonies were assessed for the treatment Rootshield Plus⁺ and non-
191 inoculated control using *Trichoderma* selective medium. *Trichoderma* selective medium was
192 prepared by mixing 0.1 g of MgSO₄·7H₂O (Sigma-Aldrich), 0.45 g of KH₂PO₄ (Sigma-
193 Aldrich), 0.5 g of NH₄NO₃ (Alfa Aesar, Tewksbury, MA, USA), 0.075 g of KCl (Sigma-
194 Aldrich), 1.5 g of dextrose (VWR, Radnor, PA, USA), 0.01 g of FeSO₄·7H₂O (VWR), 0.01 g of
195 MnSO₄·H₂O (Fisher Scientific, Pittsburgh, PA, USA), 0.01 g of ZnSO₄·7H₂O (Fisher
196 Scientific), 0.015 g of rose bengal (Fisher Scientific), 10.0 g of agar (Sigma-Aldrich) with 500
197 ml of deionized water and autoclaved for 15 minutes (Chung and Hoitink 1990). Additionally,
198 0.05 g of PCNB (99% [GC]), 5 µl of Subdue MAXX (Syngenta International AG), 0.025 g
199 chloramphenicol (Sigma-Aldrich), and 0.025 g streptomycin sulfate (Acros Organics) were
200 mixed to the medium. One-gram root sample of each plant were ultrasonicated with 10 ml of
201 sterilized deionized water (5.7 liters of Fisherbrand M-Series Mechanical Ultrasonic Cleaning
202 Bath; Thermo Fisher Scientific Inc., Waltham, MA, USA) for 3 min and then agitated with a
203 shaker (Fisherbrand Incubating Mini-Shaker; Thermo Fisher Scientific Inc.) at 250 rpm for 30
204 min at room temperature. The particles were then allowed to settle for 30 min. Serial dilutions of
205 10⁻² and 10⁻⁴ were prepared and spread-plated using glass beads (3-mm solid glass beads; Walter

206 Stern Inc., Manorhaven, NY, USA), as well as 100 μ l of the undiluted sample.
207 *Trichoderma* colonies on plates were recorded after incubation for 10-days in a dark container at
208 room temperature. The number of colony forming units (CFU) per gram of root sample was
209 calculated using the plate counts, the dilution factor, and the plated volume [CFU= (number of
210 colonies x dilution factor)/volume plated (ml)].

211 **Data analysis:** The study was designed in a completely randomized design with six replications
212 of each treatment. Data analysis for the plant growth data were done using one-way analysis of
213 variance (ANOVA) with SAS statistical software and means separated by Fisher's Least
214 Significant Difference test ($\alpha = 0.05$) (Proc GLM). Welch *t*-test ANOVA was used for data with
215 unequal variance (Welch 1947; Zheng et al. 2013). Percentage data on root rot disease severity
216 and pathogen recovery from root samples were analyzed using Generalized Linear Mixed Model
217 (GLMM) with beta distribution (PROC GLIMMIX). Data with the values 0% and 100% were
218 changed to 0.01 and 0.99 to meet the assumption of GLMM. Data on the *Trichoderma* colonies
219 were analyzed using PROC GENMOD with negative binomial distribution (Agresti 2003).

220 **Results**

221 Performance of the treatments varied in preventative and curative applications except for
222 Empress Intrinsic, Pageant Intrinsic, Segovis, and Subdue MAXX. They were consistent in
223 suppressing *Phytophthora* root rot disease severity compared to the inoculated control in both
224 trials.

225 Signature Xtra performed better in the first trial with both preventative and curative
226 applications for 1-day, 3-days and 7-days flooding in suppressing disease severity. Efficacy was
227 also demonstrated in preventative treatments for all the flooding conditions in trial 2 including 1-
228 day flooding preventative treatment. However, it was not effective in suppressing disease
229 severity in 3-days and 7-days flooding for the curative application in trial 2. Actigard suppressed
230 *Phytophthora* root rot disease severity significantly in all flooding days when applied
231 preventatively 3 weeks before flooding. But it was ineffective in 7-days flooding in trial 2 when
232 applied 1 week before flooding. Interface was effective as a curative application compared to the
233 preventative application. Stargus and ON-Gard did not perform well when applied preventatively
234 3 weeks or 1 week before flooding. Rootshield Plus was effective against *Phytophthora* root rot

235 in 1-day flooding and 3-days flooding. The highest pathogen recovery was obtained from the
236 inoculated control and the least from the non-inoculated control. *Trichoderma* colonies were
237 higher in 1-day and 3-days flooding periods than at 7-days flooding.

238 ***Preventative treatments - 3 weeks before flooding***

239 ***1-day flooding:*** In trial 1, no significant differences were observed among the treatments for
240 plant height increase and plant width increase compared to the inoculated control (Height: $F =$
241 1.69, $P = 0.0746$; Width: $F = 0.97$, $P = 0.4931$). Seedlings treated with ON-Gard, Aliette,
242 Signature Xtra, Interface, and the non-inoculated control had significantly higher total fresh plant
243 weight when compared to the inoculated control ($F = 3.27$, $P = 0.0036$). For the fresh root
244 weight, all treatments had significantly higher total fresh root weight except for Actigard,
245 Empress Intrinsic, Stargus, RootShield Plus, Segovis, Subdue MAXX, and Tartan as compared
246 to the inoculated control ($F = 3.39$, $P = 0.0028$). All treatments significantly lowered the disease
247 severity except Stargus when compared to the inoculated control ($F = 7.29$, $P = 0.0004$, Fig. 1).
248 Percentage recovery of the pathogen from the plated roots was significantly lower for the
249 treatments Orkestra Intrinsic, Segovis, Subdue MAXX, Signature Xtra, Tartan, and Interface
250 among the preventative treatments compared to the inoculated control ($F = 2.37$, $P = 0.0089$,
251 Table 2).

252 In trial 2, Empress Intrinsic, Subdue MAXX, and Signature Xtra had significantly higher
253 plant height increase among the treated seedlings compared to the inoculated control ($F = 6.94$, P
254 ≤ 0.0001). No significant differences were observed among the treated seedlings for average
255 plant width increase, total fresh plant weight, and total fresh root weight (Width: $F = 0.84$, $P =$
256 0.6223; Total plant weight: $F = 0.47$, $P = 0.9397$; Total root weight: $F = 0.54$, $P = 0.9019$). All
257 treatments significantly suppressed *Phytophthora* root rot disease severity compared to the
258 inoculated control ($F = 5.16$, $P \leq 0.0001$, Fig. 1). Pathogen recovery was significantly lower for
259 all treatments except for Actigard, Aliette, RootShield Plus and Tartan ($F = 2.59$, $P = 0.0042$,
260 Table 2).

261 ***3-days flooding:*** In trial 1, no significant differences were observed among the treatments for
262 plant height increase compared to the inoculated control ($F = 1.52$, $P = 0.1230$). Average plant
263 width increase was higher for all treatments except for Actigard and RootShield Plus as
264 compared to the inoculated control ($F = 2.32$, $P = 0.0279$). Flowering dogwood seedlings treated

265 with Signature Xtra had significant higher total fresh plant weight and total fresh root weight
 266 compared to the inoculated control (Total plant weight: $F = 4.45$, $P = 0.0004$; Total root weight:
 267 $F = 2.83$, $P = 0.0092$). All treatments significantly suppressed Phytophthora root rot disease
 268 severity except for Stargus and Interface compared to the inoculated control ($F = 8.31$, $P \leq$
 269 0.0001 , Fig. 1). Percentage recovery of the pathogen was significantly lower for Subdue MAXX,
 270 Signature Xtra and Tartan compared to the inoculated control ($F = 1.84$, $P = 0.0477$, Table 2).

271 In trial 2, plant height increase was significantly higher for all treatments except for
 272 Actigard, Tartan, ON-Gard, and Stargus ($F = 3.28$, $P = 0.0035$). No significant differences were
 273 observed among the treated seedlings for average plant width increase ($F = 0.72$, $P = 0.7446$).
 274 Fresh plant weight was significantly higher for all treatments except for Stargus, ON-Gard, and
 275 Tartan among the treated seedlings compared to the inoculated control ($F = 1.87$, $P = 0.0434$).
 276 No significant differences were observed among the treatments for fresh root weight compared
 277 to the inoculated control ($F = 1.60$, $P = 0.0980$). Seedlings treated with all other treatments
 278 except for Stargus and ON-Gard had significantly suppressed disease severity compared to the
 279 inoculated control ($F = 5.80$, $P \leq 0.0001$, Fig. 1). All treatments had significantly lower
 280 percentage recovery of the pathogen compared to the inoculated control ($F = 5.11$, $P \leq 0.0001$,
 281 Table 2).

282 **7-days flooding:** In trial 1, there were significant differences among the treatments for plant
 283 height increase, average plant width increase, total fresh plant weight, total fresh root weight,
 284 Phytophthora root rot disease severity, and pathogen recovery. Aliette, Stargus, ON-Gard,
 285 Pageant Intrinsic, and Interface had significantly higher plant height increase as compared to the
 286 inoculated control ($F = 2.17$, $P = 0.0383$). Average plant width increase was higher for seedlings
 287 treated with Aliette, ON-Gard, and Signature Xtra compared to the inoculated control ($F = 6.44$,
 288 $P \leq 0.0001$). Seedlings treated with Aliette, ON-Gard, Pageant Intrinsic, Signature Xtra and
 289 Interface had significantly higher total fresh plant weight and those treated with only ON-Gard
 290 and Pageant Intrinsic had significantly higher total fresh root weight as compared to the
 291 inoculated control (Total plant weight: $F = 2.20$, $P = 0.0366$; Total root weight: $F = 1.96$, $P =$
 292 0.0324). All treatments significantly suppressed Phytophthora root rot disease severity compared
 293 to the inoculated control except Stargus and Interface ($F = 9.23$, $P \leq 0.0001$, Fig. 1). Aliette, ON-

294 Gard and Subdue MAXX had significantly lower percentages of pathogen recovery compared to
295 the inoculated control among the treated seedlings ($F = 2.19$, $P = 0.0157$, Table 2).

296 In trial 2, no significant differences were observed among the treatments for plant height
297 increase, average plant width increase, total fresh plant weight, and total root weight compared to
298 the inoculated control (Height: $F = 1.56$, $P = 0.1110$; Width: $F = 1.64$, $P = 0.0875$; Total plant
299 weight: $F = 0.80$, $P = 0.6703$; Total root weight: $F = 0.62$, $P = 0.8364$). All treatments except
300 Aliette, Stargus, ON-Gard, RootShield Plus, and Interface significantly suppressed Phytophthora
301 root rot disease severity compared to the inoculated control ($F = 3.97$, $P \leq 0.0001$, Fig. 1).
302 Percentage recovery of the pathogen was significantly lower for all treatments except Stargus,
303 RootShield Plus, Tartan and Interface compared to the inoculated control ($F = 5.41$, $P \leq 0.0001$,
304 Table 2).

305 In trial 1, *Trichoderma* colonies of RootShield Plus were significantly higher for 1-day
306 flooding and 3-days flooding compared to 7-days flooding ($\chi^2 = 11.57$, $P = 0.0031$) whereas in
307 trial 2, *Trichoderma* colonies in 1-day flooding were significantly higher compared to 7-days
308 flooding but those in 3-days flooding were not significantly different than 7-days flooding ($\chi^2 =$
309 7.41 , $P = 0.0246$, Fig. 2).

310 ***Preventative application - 1 week before flooding***

311 ***1-day flooding:*** In trial 1, flowering dogwood seedlings treated with Signature Xtra and Aliette
312 had significantly higher plant height increase, and those treated with Signature Xtra and Interface
313 had significantly higher total fresh plant weight compared to the inoculated control (Height: $F =$
314 3.02 , $P = 0.0010$; Total plant weight: $F = 4.57$, $P = 0.0003$). No significant differences were
315 observed among the treatments for average plant width increase compared to the inoculated
316 control ($F = 1.19$, $P = 0.3029$). Total fresh root weight was significantly higher for the seedlings
317 treated with Interface when compared to the inoculated control ($F = 3.99$, $P = 0.0009$). Disease
318 severity was significantly suppressed by all treatments compared to the inoculated control ($F =$
319 7.06 , $P \leq 0.0001$, Fig. 3). All treatments had significantly lower percentages of pathogen
320 recovery compared to the inoculated control ($F = 4.89$, $P \leq 0.0001$, Table 3).

321 In trial 2, no significant differences were observed among the treatments for plant height
322 increase, plant width increase, total plant weight and total root weight compared to the inoculated

323 control (Height: $F = 1.45$, $P = 0.1521$; Width: $F = 1.04$, $P = 0.4278$; Total plant weight: $F = 0.42$,
 324 $P = 0.9654$; Total root weight: $F = 0.35$, $P = 0.9832$). All treatments significantly suppressed
 325 Phytophthora root rot disease severity compared to the inoculated control ($F = 5.48$, $P \leq 0.0001$,
 326 Fig. 3). Percentage recovery of the pathogen was significantly lower for all treatments except for
 327 Actigard and Aliette compared to the inoculated control ($F = 2.64$, $P = 0.0036$, Table 3).

328 **3-days flooding:** In trial 1, no significant differences were observed among the treatments for
 329 plant height increase, total plant weight and total root weight (Height: $F = 1.72$, $P = 0.0684$;
 330 Total plant weight: $F = 1.52$, $P = 0.1260$; Total root weight: $F = 1.68$, $P = 0.0778$) whereas
 331 Actigard and RootShield Plus had significantly higher average plant width increase compared to
 332 the inoculated control ($F = 2.20$, $P = 0.0365$). Disease severity was significantly lowered by all
 333 treatments compared to the inoculated control ($F = 6.03$, $P \leq 0.0001$, Fig. 3). All treatments had
 334 significantly lower percentage of pathogen recovery compared to the inoculated control except
 335 Stargus ($F = 3.06$, $P = 0.0009$, Table 3).

336 In trial 2, Empress Intrinsic, Segovis, Subdue MAXX, and Signature Xtra had
 337 significantly higher plant height increase compared to the inoculated control ($F = 2.58$, $P =$
 338 0.0156). Seedlings treated with Subdue MAXX and Signature Xtra had significantly higher
 339 average plant width increase among the treated seedlings compared to the inoculated control ($F =$
 340 3.22 , $P = 0.0040$). ON-Gard, Segovis, Subdue MAXX, Signature Xtra, and Stargus had
 341 significantly higher total fresh plant weight ($F = 3.36$, $P = 0.0030$) whereas only Segovis,
 342 Subdue MAXX, and Signature Xtra had significantly higher total fresh root weight ($F = 3.04$, P
 343 $= 0.0058$) among the treated dogwood seedlings compared to the inoculated control. All
 344 treatments significantly suppressed Phytophthora root rot disease severity except Stargus and
 345 Interface compared to the inoculated control ($F = 11.10$, $P \leq 0.0001$, Fig. 3). Percentage recovery
 346 of the pathogen was significantly lower in all treatments compared to inoculated control except
 347 Stargus and Interface ($F = 4.24$, $P \leq 0.0001$, Table 3).

348 **7-days flooding:** In trial 1, no treated seedlings had significantly higher plant height increase and
 349 average plant width increase compared to the inoculated control (Height: $F = 0.83$, $P = 0.6303$;
 350 Width: $F = 0.74$, $P = 0.7299$). Aliette, Stargus, Orkestra Intrinsic, RootShield Plus, Subdue
 351 MAXX, and Signature Xtra treatments had significantly higher total fresh plant weight compared
 352 to the inoculated control ($F = 15.31$, $P \leq 0.0001$). However, all these treatments resulted in

353 significantly higher total fresh plant weight and higher total fresh root weight except for Subdue
354 MAXX compared to the inoculated control ($F = 6.69, P \leq 0.0001$). For the disease severity, all
355 treatments except for Interface significantly suppressed Phytophthora root rot compared to the
356 inoculated control ($F = 8.06, P \leq 0.0001$, Fig. 3). Treatments Stargus, ON-Gard, Segovis, Tartan,
357 and Interface had similar pathogen recovery as inoculated control ($F = 9.05, P \leq 0.0001$, Table
358 3).

359 In trial 2, no significant differences were observed among the treatments for plant height
360 increase and average plant width increase (Height: $F = 0.92, P = 0.5454$; Width: $F = 0.88, P =$
361 0.5840). Total fresh plant weight and total fresh root weight were significantly higher in
362 seedlings treated with Segovis and Subdue MAXX compared to the inoculated control (Total
363 plant weight: $F = 3.69, P = 0.0015$; Total root weight: $F = 2.90, P = 0.0077$). All treatments
364 except for Actigard, Aliette, Stargus, ON-Gard, Tartan, and Interface significantly suppressed
365 disease severity compared to the inoculated control ($F = 7.10, P \leq 0.0001$, Fig. 3). All treatments
366 had significantly lower pathogen recovery compared to the inoculated control except Actigard,
367 Aliette, Stargus, ON-Gard, Tartan, and Interface ($F = 6.25, P \leq 0.0001$, Table 3).

368 *Trichoderma* colonies were significantly higher for 1-day flooding and 3-days flooding
369 compared to 7-days flooding in both trials (Trial 1: $\chi^2 = 10.22, P = 0.0006$; Trial 2: $\chi^2 = 12.3, P$
370 $= 0.0021$, Fig. 2).

371 ***Curative application***

372 ***1-day flooding:*** In trial 1, flowering dogwood seedlings treated with Signature Xtra and Interface
373 had significantly higher plant height increase, total fresh plant weight, and total fresh root weight
374 compared to the inoculated control (Height: $F = 2.40, P = 0.0419$; Total plant weight: $F = 3.19,$
375 $P = 0.0113$; Total root weight: $F = 3.47, P = 0.0073$). All fungicidal treatments except Subdue
376 MAXX had higher average plant width increase compared to the inoculated control ($F = 4.35, P$
377 $= 0.0020$). All treatments significantly suppressed Phytophthora root rot disease severity
378 compared to the inoculated control ($F = 1.94, P \leq 0.0001$, Fig. 4). All treatments except Interface
379 had significantly lower pathogen recovery compared to the inoculated control ($F = 2.18, P =$
380 0.0332 , Table 4).

381 In trial 2, no significant differences were observed among the treatments for plant height
 382 increase ($F = 1.88$, $P = 0.0686$) and total fresh root weight ($F = 1.90$, $P = 0.0645$) as compared to
 383 the inoculated control. Empress Intrinsic and Interface treated seedlings had significantly higher
 384 total fresh plant weight ($F = 3.15$, $P = 0.0030$), compared to the inoculated control among the
 385 fungicidal treatments. All treatments had significantly higher average plant width increase except
 386 Pageant Intrinsic and Tartan compared to the inoculated control ($F = 2.85$, $P = 0.0063$). All
 387 treatments except Orkestra Intrinsic significantly suppressed Phytophthora root rot disease
 388 severity compared to the inoculated control ($F = 5.19$, $P \leq 0.0001$, Fig. 4). All treatments had
 389 significantly lower percentage of pathogen recovery compared to the inoculated control ($F =$
 390 2.30 , $P = 0.0246$, Table 4).

391 **3-days flooding:** In trial 1, seedlings treated with Signature Xtra and Interface had significantly
 392 higher plant height increase, average plant width increase, total fresh plant weight, and total fresh
 393 root weight as compared to the inoculated control (Height: $F = 11.95$, $P \leq 0.0001$; Width: $F =$
 394 28.50 , $P \leq 0.0001$; Total plant weight: $F = 7.74$, $P \leq 0.0001$; Total root weight: $F = 9.23$ $P \leq$
 395 0.0001). All treatments significantly suppressed Phytophthora root rot disease severity compared
 396 to the inoculated control ($F = 14.48$, $P \leq 0.0001$, Fig. 4). No differences were observed among
 397 the fungicidal treatments for the percentage of pathogen recovery compared to the inoculated
 398 control ($F = 1.25$, $P = 0.0028$, Table 4).

399 In trial 2, no significant differences were observed among the treatments for the plant
 400 height increase, average plant width increase, total fresh plant weight, and total fresh root weight
 401 (Height: $F = 0.90$, $P = 0.5401$; Width: $F = 0.59$, $P = 0.8145$; Total plant weight: $F = 0.31$, $P =$
 402 0.9757 ; Total root weight: $F = 0.19$, $P = 0.9960$). All treatments except Orkestra Intrinsic and
 403 Signature Xtra significantly suppressed Phytophthora root rot disease severity compared to the
 404 inoculated control ($F = 6.18$, $P \leq 0.0001$, Fig. 4). All treatments except Orkestra Intrinsic had
 405 significantly lower pathogen recovery compared to the inoculated control ($F = 2.83$, $P = 0.0065$,
 406 Table 4).

407 **7-days flooding:** In trial 1, no treated seedlings had significantly higher plant height increase,
 408 average plant width increase, total plant weight and total fresh root weight compared to the
 409 inoculated control (Height: $F = 1.87$, $P = 0.0699$; Width: $F = 2.04$, $P = 0.0460$; Total plant
 410 weight: $F = 1.54$, $P = 0.1512$; Total root weight: $F = 1.92$, $P = 0.0623$). Similar to 1-day and 3-

411 days flooding, all treatments significantly suppressed Phytophthora root rot disease severity
412 compared to the inoculated control ($F = 7.35$, $P \leq 0.0001$, Fig. 4). All treatments except Aliette,
413 Orkestra Intrinsic, and Interface significantly lower pathogen recovery compared to the
414 inoculated control ($F = 4.14$, $P = 0.0003$, Table 4).

415 In trial 2, Subdue MAXX and Interface had significantly higher plant height increase
416 compared to the inoculated control (Height: $F = 5.41$, $P = 0.0005$). No significant differences
417 were observed among the treated dogwood seedlings for average plant width increase, total fresh
418 weight and total fresh root weight compared to the inoculated control (Width: $F = 6.15$, $P =$
419 0.0002 ; Total plant weight: $F = 5.97$, $P = 0.0002$; Total root weight: $F = 6.66$, $P = 0.0001$). All
420 treatments except for Orkestra Intrinsic and Signature Xtra significantly suppressed the disease
421 severity compared to the inoculated control ($F = 9.60$, $P \leq 0.0001$, Fig. 4); however, only
422 Empress Intrinsic, Segovis, Subdue MAXX, and Interface were the fungicidal treatments with
423 significantly lower pathogen recovery compared to the inoculated control ($F = 3.90$, $P = 0.0005$,
424 Table 4).

425 Discussion

426 Fungicides, biofungicides, host-plant defense inducers, and fertilizer had varied results in their
427 ability to suppress Phytophthora root rot severity during flooding conditions. Some of the
428 treatments were effective when used preventatively 3 weeks before the flooding whereas some of
429 them were effective when used 1 week before flooding. Most of the treatments used as curative
430 applications, except for Orkestra Intrinsic and Signature Xtra, successfully suppressed
431 Phytophthora root rot disease severity. The effectiveness of some of the treatments also varied
432 between the trials. Irrespective of the treatments being able to suppress root rot disease
433 symptoms caused by *P. cinnamomi*, all inoculated seedlings exhibited the disease symptoms.
434 The highest disease severity was observed in the inoculated control while the lowest disease
435 severity was observed in the non-inoculated control. The physical appearances (shedding of
436 leaves, plants vigor) of the non-inoculated control plants were better than those of the inoculated
437 control plants (visual observation).

438 The results displayed that the severity of the disease increased with increased flooding
439 duration, which is consistent with the results of Brown et al. (2019b, 2019a) using *Cornus florida*

440 L. seedlings. The severity of *Phytophthora* root rot in dogwood seedlings was higher in 7-days
441 flooding than 3-days or 1-day flooding in both preventative and curative treatments. Wilcox and
442 Mircetich (1985) and de Silva et al. (1999) observed increased root rot severity in increased
443 flooding conditions in two species of *Prunus* and *Vaccinium corymbosum* L., respectively. In a
444 study conducted by Ploetz and Schaffer (1987), a contrast between the disease severity in the
445 fine loamy soil and potting mixture was observed in the avocado seedlings where the fine loamy
446 soil had increased root rot severity but not in potting mixture. The author explained the
447 differences were because of the water holding capacity of the two soil types. In our study, as the
448 simulated flooding conditions were created for 1-, 3-, and 7-days using zip-lock bags, the disease
449 severity increased with increased flooding duration. This shows that proper drainage is important
450 to manage the *Phytophthora* infections. The moisture condition in pine bark substrate is less
451 compared to the usual field soil where nursery crops are grown. In a study conducted by Ownley
452 et al. (1990), the moisture retained in pine bark was comparatively less than that of pine bark
453 mixed with peat or soil. But in another study conducted by Wisdom et al. (2017), the total
454 porosity of the pine bark substrate alone was comparatively less than the combination of pine
455 bark and vermiculite. The increased porosity in the potting substrate helps to increase oxygen
456 amount in the root zone and delays root rot disease development. Increased soil moisture creates
457 favorable conditions for the production, dispersal, and germination of zoospores. Also, the
458 saturated soil conditions make root and collar regions more susceptible for *Phytophthora*
459 infestation (Newhook and Podger 1972; Weste and Marks 1987). In the absence of soil
460 moisture, chlamydospores are formed and once the environmental conditions become favorable,
461 they germinate to form either mycelium or sporangium with zoospores to start re-infection
462 (Weste and Vithanage 1978, 1979). Thus, appropriate cultural measures like soil reclamation,
463 specifically proper drainage, are important for managing *Phytophthora* root rot.

464 Several systemic fungicides were effective against *P. cinnamomi* during flooding
465 conditions. Mefenoxam (Subdue MAXX) fungicide has been used preventatively and curatively
466 to manage the oomycetes in nurseries, turf, and landscape. In our study, mefenoxam was one of
467 the few fungicides that was effective in suppressing *Phytophthora* root rot disease severity in
468 both preventative and curative applications for all 1-, 3-, and 7-days flooding. In other studies,
469 conducted by Baysal-Gurel et al. (2016) and Gray et al. (2020), mefenoxam was effective in
470 managing *Phytophthora* root rot in dogwood, hydrangea, and citrus seedlings, respectively. Soil

471 drench application of mefenoxam fungicide was able to control root rot caused by *P. cinnamomi*
472 in ornamental crops (Hu et al. 2010). Mefenoxam acts against both mycelial growth and
473 sporulation, and checks the synthesis of ribosomal RNA to suppress the pathogenic activity of *P.*
474 *cinnamomi* (Cohen and Coffey 1986; Davidse et al. 1983). However, excessive use of
475 fungicides can lead to the emergence of a fungicide insensitive pathogen population.

476 Along with Subdue MAXX, Intrinsic brand fungicides like Empress Intrinsic and Pageant
477 Intrinsic suppressed Phytophthora root rot severity in both preventative and curative applications.
478 Intrinsic brand fungicides are a strobilurin class of fungicides, which are found to act as host-
479 plant defense inducers (Chandrasekhar et al. 2017). Host-plant defense inducers act to improve
480 the plant's immune system and enhance the defensive/response mechanism of plants in biotic
481 and abiotic stress conditions (Chandrasekhar et al. 2017; Brown et al. 2019b). Brown et al. (2019
482 a,b) and Baysal-Gurel et al. (2016) evaluated the efficacy of pyroclostrobin (strobilurin) to
483 manage Phytophthora root rot in flowering dogwoods and hydrangeas, respectively, and found it
484 to be effective. Similarly, a higher rate of pyroclostrobin in comparison to its lower rate was
485 found to be effective as seed treatment of soybean and corn against Phytophthora root rot (Li et
486 al. 2020; Radmer et al. 2017). Strobilurin inhibits mitochondrial respiration of the oomycetes
487 along with enhancing physiological changes to embrace plant vigor, stress tolerance, and disease
488 resistance (Han et al. 2012; Herms et al. 2002; Jaleel et al. 2008; Venancio et al. 2003).

489 In this study, Segovis (piperidinyl-thiazole isooxazoline) showed promising results in
490 managing root rot disease caused by *P. cinnamomi* in all flooding durations as both preventative
491 and curative applications. The drench application of the fungicide significantly reduced
492 Phytophthora root rot in impatiens (*Impatiens walleriana* Hooker) even after 120 days of
493 application (Harlan and Hausbeck 2019). In another study conducted by Gray et al. (2020), a
494 lower dose of oxathiapiprolin (piperidinyl-thiazole isooxazoline) was effective in management of
495 root rot disease in citrus seedlings. Piperidinyl-thiazole isooxazoline targets the oxysterol binding
496 protein to provide resistance against the oomycete pathogens (Mboup et al. 2021; Miao et al.
497 2016; Pasteris et al. 2016). This fungicide highly inhibits mycelial growth, sporangial
498 production, and cystospore germination, especially in *Phytophthora* spp. (Lin et al. 2020).

499 Some of the fungicides were inconsistent in their efficacy against *P. cinnamomi* during
500 the flooding conditions. Treatments such as Actigard (acibenzolar-S-methyl) were effective in

501 both trials of preventative application 3 weeks before flooding. But they were ineffective in the
502 second trial of preventative application 1 week before flooding. The preventative application of
503 acibenzolar-S-methyl and mfenoxam was previously found to be effective against *Phytophthora*
504 root rot of pepper in the flooding conditions (Matheron and Porchas 2002). Similarly, a curative
505 application of acibenzolar-S-methyl was able to significantly reduce *P. cinnamomi* infection in
506 *Pinus radiata* D. Don, *Banksia integrifolia* L., and *Isopogon cuneatus* R. Br. (Ali et al. 2000). In
507 recent years, increasing cases of acibenzolar-S-methyl consistency against fungal pathogens have
508 been observed in many studies (Bokshi et al. 2006; Darras et al. 2006). One of the reasons might
509 be different environmental conditions or seasonal timing between the trials. Brown et al. (2019b)
510 and Darvas and Becker (1984) mentioned that extensive application of mfenoxam in nurseries
511 could lead to fungicide resistance affecting the effectiveness of other fungicides.

512 Phosphoric acid compound of fosetyl-al (Aliette, Signature Xtra) is a recommended
513 chemical control for *P. cinnamomi*. However, the inconsistency appeared with curative and
514 preventative applications in this study. Aliette was effective in trial 1 but failed to suppress
515 *Phytophthora* root rot severity compared to the inoculated control in trial 2 for both preventative
516 and curative applications. Unlike Aliette, Signature Xtra was effective in preventative
517 application but performed poorly in trial 2 as a curative application. Previously, Coffey (1987),
518 Duvenhage (1994), and Darvas et al. (1983) found that fosetyl-al was effective against
519 *Phytophthora* root rot of avocado caused by *P. cinnamomi*. Similarly, fosetyl-al was
520 recommended for the long-term management of crown rot and root rot disease in apples
521 (Utkhede and Smith 1993).

522 The effect of Stargus (*Bacillus amyloliquefaciens* strain F727) to suppress *Phytophthora*
523 root rot severity was negligible as it was not effective in all 1-, 3- and 7-days flooding when
524 applied 3 weeks before flooding. However, it was effective in 1-day flooding when applied 1
525 week before flooding. A similar result was observed by Brown et al. (2019b) where Stargus was
526 effective in 1-day flooding but not for a longer duration. *Bacillus amyloliquefaciens* was found to
527 be effective in controlling *Phytophthora* root rot disease severity in soybean in non-flooding
528 conditions (Liu et al. 2019). Another biofungicide, Rootshield Plus (*Trichoderma harzianum*
529 Rifai strain T-22 and *T. virens* strain G-41), was effective in all flooding durations when applied
530 1 week before flooding; but was effective only in 1-day flooding when applied 3 weeks before

531 flooding. The number of *Trichoderma* colonies decreased with increased flooding duration. This
532 suggests that the application of biofungicides should be done closer to the flooding duration to
533 ensure maximum protection. The efficacy of the microorganisms against the pathogen may
534 decrease with time and flooding duration as the number of colonies are reduced. Similar
535 suggestions were recommended by Knudsen and Li (1990) as an increase in soil moisture could
536 decrease hyphal proliferation.

537 Similarly, effectiveness of the Stressgard formulated fungicides (multiple modes of
538 action), Tartan and Interface, varied. Tartan significantly suppressed Phytophthora root rot
539 disease severity in all 1-, 3-, and 7-days flooding in curative and preventative applications but
540 failed in the second trial of preventative application 1 week before flooding. Interface could not
541 suppress the disease severity in all flooding durations when applied 3 weeks before flooding.
542 However, it suppressed disease severity during 1-day flooding duration when applied 1 week
543 before flooding. Intrinsic brand fungicide Orkestra was effective as a preventative application in
544 both the trials but was ineffective in trial 2 of curative application.

545 In all these inconsistencies, we observed that most of the treatments failed in the second
546 trial, irrespective of the application method. This might be because of the increased disease
547 pressure in the second trial. The possible reasons for increased disease severity in trial 2 may be
548 a) the difference in environmental conditions between trial 1 and trial 2, and b) the occurrence of
549 powdery mildew in trial 2, c) TRIACT® 70 might have performed poorly against powdery
550 mildew and the effect of root rot and powdery mildew might have caused the treatments to
551 perform poorly. However, the foliar spray of TRIACT® 70 should have had uniform effects
552 across all treatments.

553 In this study, comparatively less disease severity was observed in curative treatments
554 than the preventative treatments, which might be because of the repeated application of the
555 curative treatments. The differences in root rot severity of the inoculated controls in preventative
556 application 3 weeks before flooding and 1 week before flooding were minimal. However, some
557 of the fungicides performed differently in two application timings. Actigard was more effective
558 when applied 3 weeks before flooding than 1 week before flooding. Contrastingly, RootShield
559 Plus was more effective when applied 1 week before flooding than 3 weeks before flooding. ON-

560 Gard was effective in 1-day and 3-days flooding when applied 1 week before flooding but was
561 effective only for 1-day flooding when applied 3 weeks before flooding.

562 The efficacy of the fungicides, biofungicides, host-plant defense inducers, and fertilizer
563 have varied in different application timings, application methods, flooding durations, and
564 between the trials. The growers should take these factors into consideration when looking for
565 management practices for their crops. As severity of infection by *P. cinnamomi* increases with
566 increasing soil moisture, cultural practices to improve drainage are highly recommended. Based
567 on this study, both the preventative and curative applications of Empress Intrinsic, Pageant
568 Intrinsic, Segovis, and Subdue MAXX can be used as chemical control in nurseries. As flooding
569 conditions cannot be accurately predicted, curative application of these fungicides might be
570 helpful in managing Phytophthora root rot disease. The strategic repeated application of curative
571 fungicides may provide longer protection to the crop and prevent economic loss. Some of the
572 fungicides work best when applied preventatively while some work best as curative application.
573 In this study, Actigard and Tartan were effective when applied preventatively 3 weeks before the
574 flooding whereas RootShield Plus performed well when applied 1 week before the flooding.
575 Similarly, Signature Xtra and Orkestra Intrinsic were effective as preventative application
576 measure but not as curative application measure. This shows that effectiveness of the chemicals
577 depends on the application timing. Thus, selection of treatments based on its efficacy in different
578 application methods and application timing will help in effective management of *P. cinnamomi*
579 in flooding conditions.

580 **Acknowledgements**

581 This work was supported by the United States Department of Agriculture (USDA) National
582 Institute of Food and Agriculture (NIFA) Evans-Allen funding under award numbers TENX-
583 1926-CCOCP and TENX-S-1083, and USDA-NIFA, Southern Sustainable Agriculture Research
584 and Education program under award number GS20-228. Any mention of product names is for
585 informational purposes only and does not imply an endorsement by the authors or their
586 institution.

587 **References**

- 588 Addesso, K., Baysal-Gurel, F., Oliver, J., Ranger, C., and O'Neal, P. 2018. Interaction of a
589 preventative fungicide treatment and root rot pathogen on ambrosia beetle attacks during
590 a simulated flood event. *Insects* 9:83.
- 591 Agresti, A. 2003. *Categorical data analysis*. John Wiley & Sons. 2nd ed. Wiley, New York, NY.
- 592 Ali, Z., Smith, I., and Guest, D. I. 2000. Combinations of potassium phosphonate and Bion
593 (acibenzolar-S-methyl) reduce root infection and dieback of *Pinus radiata*, *Banksia*
594 *integrifolia* and *Isopogon cuneatus* caused by *Phytophthora cinnamomi*. *Australasian*
595 *Plant Pathology* 29:59-63.
- 596 Aryantha, I. N. P., and Guest, D. I. 2006. Mycoparasitic and antagonistic inhibition on
597 *Phytophthora cinnamomi* rands by microbial agents isolated from manure composts.
598 *Plant Pathology Journal* 5:297-298.
- 599 Balci, Y., Balci, S., Eggers, J., MacDonald, W., Juzwik, J., Long, R., and Gottschalk, K. 2007.
600 *Phytophthora* spp. associated with forest soils in eastern and north-central US oak
601 ecosystems. *Plant Disease* 91:705-710.
- 602 Baysal-Gurel, F., Kabir, N., and Blalock, A. 2016. Root Diseases of Hydrangea. TSU-16-
603 0237(A)-15-61065.
- 604 Belimov, A. A., Safronova, V. I., Sergeyeva, T. A., Egorova, T. N., Matveyeva, V. A.,
605 Tsyganov, V. E., Borisov, A. Y., Tikhonovich, I. A., Kluge, C., and Preisfeld, A. 2001.
606 Characterization of plant growth promoting rhizobacteria isolated from polluted soils and
607 containing 1-aminocyclopropane-1-carboxylate deaminase. *Canadian Journal of*
608 *Microbiology* 47:642-652.
- 609 Belisle, R. J., Hao, W., McKee, B., Arpaia, M. L., Manosalva, P., and Adaskaveg, J. E. 2019.
610 New Oomycota fungicides with activity against *Phytophthora cinnamomi* and their
611 potential use for managing avocado root rot in California. *Plant disease* 103:2024-2032.
- 612 Benson, D. 1985. Fungicides for control of *Phytophthora* root of azalea in landscapes beds. *Plant*
613 *Disease* 69:697-699.
- 614 Benson, D., and Blazich, F. A. 1989. Control of *Phytophthora* root rot of *Rhododendron*
615 *chapmanii* A. Gray with Subdue. *Journal of Environmental Horticulture* 7:73-75.

- 616 Benson, D., and Grand, L. 2000. Incidence of Phytophthora root rot of Fraser fir in North
617 Carolina and sensitivity of isolates of *Phytophthora cinnamomi* to metalaxyl. Plant
618 Disease 84:661-664.
- 619 Benson, D., Sidebottom, J., and Moody, J. 2006. Control of phytophthora root rot in field
620 plantings of fraser fir with fosetyl-al and mefenoxam. Plant Health Progress 7:25.
- 621 Bezuidenhout, J., Darvas, J., and Toerien, J. 1987. Chemical control of *Phytophthora*
622 *cinnamomi*. South African Avocado Growers' Association Yearbook 10:106-108.
- 623 Bokshi, A., Morris, S., Deverall, B., and Stephens, B. 2000. Induction of systemic acquired
624 resistance in potato. In: Proceedings of the Australian Potato Research, Development
625 And Technology Transfer Conference. Adelaide, Australia.
- 626 Bokshi, A., Morris, S., McConchie, R., and Deverall, B. 2006. Pre-harvest application of 2, 6-
627 dichloroisonicotinic acid, -aminobutyric acid or benzothiadiazole to control post-harvest
628 storage diseases of melons by including systemic acquired resistance (SAR). The Journal
629 of Horticultural Science and Biotechnology 81: 700-706.
- 630 Brown, M. S., Baysal-Gurel, F., Oliver, J. B., and Adesso, K. M. 2019a. Evaluation of
631 fungicides and biofungicide to control Phytophthora root rot (*Phytophthora cinnamomi*
632 Rands) and ambrosia beetles (Coleoptera: Curculionidae: Scolytinae) on flowering
633 dogwoods exposed to simulated flood events. Crop Protection 124:104834.
- 634 Brown, M. S., Baysal-Gurel, F., Oliver, J. B., and Adesso, K. M. 2019b. Comparative
635 performance of fungicides, biofungicides, and host plant defense inducers in suppression
636 of Phytophthora root rot in flowering dogwood during simulated root flooding events.
637 Plant Disease 103:1703-1711.
- 638 Chandrasekhar, B., Umesha, S., and Kumar, H. N. 2017. Proteomic analysis of salicylic acid
639 enhanced disease resistance in bacterial wilt affected chilli (*Capsicum annuum*) crop.
640 Physiological and Molecular Plant Pathology 98:85-96.
- 641 Chung, Y., and Hoitink, H. 1990. Interactions between thermophilic fungi and *Trichoderma*
642 *hamatum* in suppression of Rhizoctonia damping-off in a bark compost-amended
643 container medium. Phytopathology 80:73-77.
- 644 Coffey, M. D. 1987. Phytophthora root rot of avocado: an integrated approach to control in
645 California. Plant Disease 71:1046-1053.

- 646 Cohen, Y., and Coffey, M. D. 1986. Systemic fungicides and the control of oomycetes. Annual
647 Review of Phytopathology 24:311-338.
- 648 Dai, T., Wang, A., Yang, X., Yu, X., Tian, W., Xu, Y., and Hu, T. 2020. PHYCI_587572: an
649 RxLR effector gene and new biomarker in a recombinase polymerase amplification assay
650 for rapid detection of *Phytophthora cinnamomi*. Forests 11:306.
- 651 Darras, A., Joyce, D., and Terry, L. 2006. Acibenzolar-S-methyl and methyl jasmonate
652 treatments of glasshouse-grown freesias suppress post-harvest petal specking caused by
653 *Botrytis cinerea*. The journal of Horticultural Science and Biotechnology 81:1043-1051.
- 654 Darvas, J., Toerien, J., and Milne, D. 1983. Injection of established avocado trees for the
655 effective control of *Phytophthora* root rot. Citrus and Subtropical Fruit Journal 591:7-10.
- 656 Davidse, L. C., Hofman, A. E., and Velthuis, G. C. 1983. Specific interference of metalaxyl with
657 endogenous RNA polymerase activity in isolated nuclei from *Phytophthora megasperma*
658 f. sp. *medicaginis*. Experimental Mycology 7:344-361.
- 659 de Silva, A., Patterson, K., Rothrock, C., and McNew, R. 1999. *Phytophthora* root rot of
660 blueberry increases with frequency of flooding. HortScience 34:693-695.
- 661 Duan, C.-H., Riley, M., and Jeffers, S. 2008. Characterization of *Phytophthora cinnamomi*
662 populations from ornamental plants in South Carolina, USA. Archives of Phytopathology
663 and Plant Protection 41:14-30.
- 664 Duniway, J. 1976. Movement of zoospores of *Phytophthora cryptogea* in soils. Phytopathology
665 66:877-882.
- 666 Duvenhage, J. 1994. Monitoring the resistance of *Phytophthora cinnamomi* to fosetyl-Al and
667 H₃PO₃. South African Avocado Growers' Association Yearbook 17:35-37.
- 668 Erwin, D. C., and Ribeiro, O. K. 1996. *Phytophthora* diseases worldwide. American
669 Phytopathological Society (APS Press), St. Paul, MN.
- 670 Evidente, A., Cristinzio, G., Punzo, B., Andolfi, A., Testa, A., and Melck, D. 2009. Flufuran, an
671 antifungal 3, 5-disubstituted furan produced by *Aspergillus flavus* Link. Chemistry &
672 Biodiversity 6:328-334.
- 673 Ferguson, A., and Jeffers, S. 1999. Detecting multiple species of *Phytophthora* in container
674 mixes from ornamental crop nurseries. Plant Disease 83:1129-1136.
- 675 Gould, A. 2012. Disease control recommendations for ornamental crops. New Jersey
676 Agricultural Experiment Station Bulletin E036. New Jersey.

- 677 Gray, M. A., Nguyen, K. A., Hao, W., Belisle, R. J., Förster, H., and Adaskaveg, J. E. 2020.
 678 Mobility of oxathiapiprolin and mefenoxam in citrus seedlings after root application and
 679 implications for managing *Phytophthora* root rot. *Plant Disease* 104:3159-3165.
- 680 Han, S.-H., Kang, B.-R., Lee, J.-H., Lee, S.-H., Kim, I.-S., Kim, C.-H., and Kim, Y.-C. 2012. A
 681 trifloxystrobin fungicide induces systemic tolerance to abiotic stresses. *The Plant*
 682 *Pathology Journal* 28:101-106.
- 683 Hardham, A. R. 2005. *Phytophthora cinnamomi*. *Molecular Plant Pathology* 6:589-604.
- 684 Hardham, A. R., and Blackman, L. M. 2018. *Phytophthora cinnamomi*. *Molecular Plant*
 685 *Pathology* 19:260-285.
- 686 Harlan, B., and Hausbeck, M. 2019. Extended control of downy mildew and *Phytophthora* root
 687 rot on ornamentals with the novel fungicide oxathiapiprolin. Pages 235-240 in:
 688 International Symposium on Advanced Technologies and Management for Innovative
 689 Greenhouses: GreenSys2019 1296.
- 690 Harman, G. E., Howell, C. R., Viterbo, A., Chet, I., and Lorito, M. 2004. *Trichoderma* species—
 691 opportunistic, avirulent plant symbionts. *Nature Reviews Microbiology* 2:43-56.
- 692 Harman, G. E. 2006. Overview of mechanisms and uses of *Trichoderma* spp. *Phytopathology*
 693 96:190-194.
- 694 Herms, S., Seehaus, K., Koehle, H., and Conrath, U. 2002. A strobilurin fungicide enhances the
 695 resistance of tobacco against tobacco mosaic virus and *Pseudomonas syringae* pv. *tabaci*.
 696 *Plant Physiology* 130:120-127.
- 697 Hu, J., Hong, C., Stromberg, E. L., and Moorman, G. W. 2010. Mefenoxam sensitivity in
 698 *Phytophthora cinnamomi* isolates. *Plant Disease* 94:39-44.
- 699 Jacobs, K., and Johnson, G. 1996. Ornamental cherry tolerance of flooding and *Phytophthora*
 700 root rot. *HortScience* 31:988-991.
- 701 Kamoun, S., Furzer, O., Jones, J. D., Judelson, H. S., Ali, G. S., Dalio, R. J., Roy, S. G., Schena,
 702 L., Zambounis, A., and Panabières, F. 2015. The Top 10 oomycete pathogens in
 703 molecular plant pathology. *Molecular Plant Pathology* 16:413-434.
- 704 Kawase, M. 1981. Anatomical and morphological adaptation of plants to waterlogging.
 705 *HortScience*. 16:8-12.
- 706 Knudsen, G., and Li, B. 1990. Effects of temperature, soil moisture, and wheat bran on growth of
 707 *Trichoderma harzianum* from alginate pellets. *Phytopathology* 80:724-727.

- 708 Kong, P., Hong, C., and Richardson, P. 2003. Rapid detection of *Phytophthora cinnamomi* using
709 PCR with primers derived from the Lpv putative storage protein genes. *Plant Pathology*
710 52:681-693.
- 711 Kozłowski, T. 1997. Responses of woody plants to flooding and salinity. *Tree physiology*
712 17:490-490.
- 713 Li, P., Sun, P., Li, D., Li, D., Li, B., and Dong, X. 2020. Evaluation of Pyraclostrobin as an
714 ingredient for soybean seed treatment by analyzing its accumulation–dissipation kinetics,
715 plant-growth activation, and protection against *Phytophthora sojae*. *Journal of*
716 *Agricultural and Food Chemistry* 68:11928-11938.
- 717 Lin, D., Xue, Z., Miao, J., Huang, Z., and Liu, X. 2020. Activity and resistance assessment of a
718 new osbp inhibitor, r034-1, in *Phytophthora capsici* and the detection of point mutations
719 in pcorp1 that confer resistance. *Journal of Agricultural and Food Chemistry* 68:13651-
720 13660.
- 721 Lin, K.-H. R., Tsou, C.-C., Hwang, S.-Y., Chen, L.-F. O., and Lo, H.-F. 2006. Paclobutrazol pre-
722 treatment enhanced flooding tolerance of sweet potato. *Journal of Plant Physiology*
723 163:750-760.
- 724 Liu, D., Li, K., Hu, J., Wang, W., Liu, X., and Gao, Z. 2019. Biocontrol and action mechanism
725 of *Bacillus amyloliquefaciens* and *Bacillus subtilis* in soybean phytophthora blight.
726 *International Journal of Molecular Sciences* 20:2908.
- 727 Matheron, M., and Porchas, M. 2002. Suppression of *Phytophthora* root and crown rot on pepper
728 plants treated with acibenzolar-S-methyl. *Plant Disease* 86:292-297.
- 729 Mboup, M. K., Sweigard, J. W., Carroll, A., Jaworska, G., and Genet, J. L. 2021. Genetic
730 mechanism, baseline sensitivity and risk of resistance to oxathiapiprolin in oomycetes.
731 *Pest Management Science*. <https://doi.org/10.1002/ps.6700>.
- 732 Miao, J., Dong, X., Lin, D., Wang, Q., Liu, P., Chen, F., Du, Y., and Liu, X. 2016. Activity of
733 the novel fungicide oxathiapiprolin against plant-pathogenic oomycetes. *Pest*
734 *Management Science* 72:1572-1577.
- 735 Neupane, K., Alexander, L., and Baysal-Gurel, F. 2021. Management of *Phytophthora*
736 *cinnamomi* using fungicides and host plant defense inducers under drought conditions: A
737 case-study of flowering dogwood. *Plant Disease*. [https://doi.org/10.1094/PDIS-04-21-](https://doi.org/10.1094/PDIS-04-21-0789-RE)
738 [0789-RE](https://doi.org/10.1094/PDIS-04-21-0789-RE)

- 739 Newhook, F., and Podger, F. 1972. The role of *Phytophthora cinnamomi* in Australian and New
740 Zealand forests. Annual Review of Phytopathology 10:299-326.
- 741 Ownley, B., Benson, D., and Bilderback, T. 1990. Physical properties of container media and
742 relation to severity of Phytophthora root rot of Rhododendron. Journal of the American
743 Society for Horticultural Science 115: 564-570.
- 744 Parajuli, M., Panth, M., Gonzalez, A., Adesso, K. M., Witcher, A., Simmons, T., and Baysal-
745 Gurel, F. 2021. Cover crop usage for the sustainable management of soilborne diseases in
746 woody ornamental nursery production system. Canadian Journal of Plant Pathology.
747 <https://doi.org/10.1080/07060661.2021.2020336>
- 748 Pasteris, R. J., Hanagan, M. A., Bisaha, J. J., Finkelstein, B. L., Hoffman, L. E., Gregory, V.,
749 Andreassi, J. L., Sweigard, J. A., Klyashchitsky, B. A., and Henry, Y. T. 2016. Discovery
750 of oxathiapiprolin, a new oomycete fungicide that targets an oxysterol binding protein.
751 Bioorganic & Medicinal Chemistry 24:354-361.
- 752 Pieterse, C. M., Van Pelt, J. A., Ton, J., Parchmann, S., Mueller, M. J., Buchala, A. J., Métraux,
753 J.-P., and Van Loon, L. C. 2000. Rhizobacteria-mediated induced systemic resistance
754 (ISR) in *Arabidopsis* requires sensitivity to jasmonate and ethylene but is not
755 accompanied by an increase in their production. Physiological and Molecular Plant
756 Pathology 57:123-134.
- 757 Pieterse, C. M., Van Pelt, J. A., Van Wees, S. C., Ton, J., Léon-Kloosterziel, K. M., Keurentjes,
758 J. J., Verhagen, B. W., Knoester, M., Van der Sluis, I., and Bakker, P. A. 2001.
759 Rhizobacteria-mediated induced systemic resistance: triggering, signalling and
760 expression. European Journal of Plant Pathology 107:51-61.
- 761 Ploetz, R. C., and Schaffer, B. 1987. Effects of flooding and Phytophthora root rot on
762 photosynthetic characteristics of avocado. Pages 290-294 in: Proceedings of Florida State
763 Horticultural Society 100:290-294.
- 764 Ploetz, R. C., and Schaffer, B. 1989. Effects of flooding and Phytophthora root rot on net gas
765 exchange and growth of avocado. Phytopathology 79:204-208.
- 766 Radmer, L., Anderson, G., Malvick, D., Kurle, J., Rendahl, A., and Mallik, A. 2017. *Pythium*,
767 *Phytophthora*, and *Phytophthora* spp. associated with soybean in Minnesota, their
768 relative aggressiveness on soybean and corn, and their sensitivity to seed treatment
769 fungicides. Plant Disease 101:62-72.

- 770 Reed, S. M. 2005. Effect of storage temperature and seed moisture on germination of stored
771 flowering dogwood seed. *Journal of Environmental Horticulture* 23:29-32.
- 772 Reeksting, B. J., Olivier, N. A., and Van den Berg, N. 2016. Transcriptome responses of an
773 ungrafted *Phytophthora* root rot tolerant avocado (*Persea americana*) rootstock to
774 flooding and *Phytophthora cinnamomi*. *BMC Plant Biology* 16:1-19.
- 775 Ribeiro, O., and Linderman, R. 1991. Chemical and biological control of *Phytophthora* species
776 in woody plants. Pages 399-410 Cambridge University Press, Sydney.
- 777 Richter, B. S., Ivors, K., Shi, W., and Benson, D. 2011. Cellulase activity as a mechanism for
778 suppression of *Phytophthora* root rot in mulches. *Phytopathology* 101:223-230.
- 779 Robin, C., Smith, I., and Hansen, E. 2012. *Phytophthora cinnamomi*. *Forest Phytophthoras* 2.
- 780 Shakya, S. K., Grünwald, N. J., Fieland, V. J., Knaus, B. J., Weiland, J. E., Maia, C., Drenth, A.,
781 Guest, D. I., Liew, E. C., and Crane, C. 2021. Phylogeography of the wide-host range
782 panglobal plant pathogen *Phytophthora cinnamomi*. *Molecular Ecology* 30:5164-5178.
- 783 Sharkawy, S., Hilal, A., Mounir, G., and Mahmoud, N. 2014. Evaluation the efficacy of some
784 biofertilizers and biofungicides for controlling *Phytophthora* root rot of apple and
785 mandarin trees in egypt. *Egyptian Journal of Phytopathology* 42:43-52.
- 786 Utkhede, R., and Smith, E. 1993. Long-term effects of chemical and biological treatments on
787 crown and root rot of apple trees caused by *Phytophthora cactorum*. *Soil Biology and*
788 *Biochemistry* 25:383-386.
- 789 Wang, C., Knill, E., Glick, B. R., and Défago, G. 2000. Effect of transferring 1-
790 aminocyclopropane-1-carboxylic acid (ACC) deaminase genes into *Pseudomonas*
791 *fluorescens* strain CHA0 and its *gac A* derivative CHA96 on their growth-promoting and
792 disease-suppressive capacities. *Canadian Journal of Microbiology* 46:898-907.
- 793 Weiland, J. E., Scagel, C. F., Grünwald, N. J., Davis, E. A., and Beck, B. R. 2021. *Phytophthora*
794 species differ in response to phosphorous acid and mefenoxam for the management of
795 *Phytophthora* root rot in rhododendron. *Plant Disease*:PDIS-09-20-1960-RE.
- 796 Welch, B. L. 1947. The generalization of 'STUDENT'S' problem when several different
797 population variances are involved. *Biometrika* 34:28-35.
- 798 Weste, G., and Vithanage, K. 1978. Seasonal variation in numbers of chlamydospores in
799 Victorian forest soils infected with *Phytophthora cinnamomi*. *Australian Journal of*
800 *Botany* 26:657-662.

- 801 Weste, G., and Vithanage, K. 1979. Survival of chlamydospores of *Phytophthora cinnamomi* in
802 several non-sterile, host-free forest soils and gravels at different soil water potentials.
803 Australian Journal of Botany 27:1-9.
- 804 Weste, G., and Marks, G. 1987. The biology of *Phytophthora cinnamomi* in Australasian forests.
805 Annual Review of Phytopathology 25:207-229.
- 806 Wilcox, W., and Mircetich, S. 1985. Effects of flooding duration on the development of
807 *Phytophthora* root and crown rots of cherry. Phytopathology 75:1451-1455.
- 808 Williams-Woodward, J. L., and DeMott, M. E. 2013. Fungicide rResistance in *Pythium* and
809 *Phytophthora* from ornamentals in Georgia. Proceedings of the International Plant
810 Propagators Society-2013 1055:453-456.
- 811 Wisdom, B., Nyembezi, M., and Agathar, K. 2017. Effect of different vermiculite and pine bark
812 media substrates mixtures on physical properties and spiral rooting of radish (*Raphanus*
813 *sativus* L.) in float tray system. Rhizosphere 3: 67-74.
- 814 Zentmyer, G. A. 1980. *Phytophthora cinnamomi* and the diseases it causes. American
815 Phytopathological Society, St. Paul, MN.
- 816 Zheng, L., Diamond, J. M., and Denton, D. L. 2013. Evaluation of whole effluent toxicity data
817 characteristics and use of Welch's T-test in the test of significant toxicity analysis.
818 Environmental Toxicology and Chemistry 32:468-474.

819

820

821

822

823

824

825 **Tables**826 **Table 1.** Fungicides, biofungicides, host plant defense inducers and fertilizer used in this study.

Treatment ^a	Application type	Application rate		Application interval	Product group	Manufacturer ^b
		ml/liter	g/liter			
Actigard 50 WG	Preventative		0.30	1 week	Host plant defense inducer	Syngenta
Aliette 80 WDG	Preventative, Curative		3.74	6 weeks	Host plant defense inducer	Bayer
Empress Intrinsic	Preventative, Curative	0.47		3 weeks	Strobilurin	BASF
Interface Stressgard	Preventative, Curative	6.25		3 weeks	Strobilurin + dicarboximide	Bayer
ON-Gard [®] 5-0-0	Preventative	25		3 weeks	Fertilizer	BioWorks
Orkestra Intrinsic	Preventative, Curative	0.78		3 weeks	Strobilurin + succinate dehydrogenase inhibitor	BASF
Pageant Intrinsic	Preventative, Curative		1.35	3 weeks	Strobilurin + succinate dehydrogenase inhibitor	BASF
RootShield Plus ⁺ WP	Preventative		0.60	6 weeks	Biofungicide	BioWorks
Segovis	Preventative, Curative	0.25		3 weeks	Piperidinyl-thiazole isoxazoline	Syngenta
Signature Xtra Stressgard	Preventative, Curative		5.99	2 weeks	Host plant defense inducer	Bayer
Stargus	Preventative	10.00		1 week	Biofungicide	Marrone
Subdue MAXX	Preventative, Curative	0.16		10 weeks	Phenylamide	Syngenta
Tartan Stressgard	Preventative Curative	3.12		3 weeks	Strobilurin + triazole	Bayer

827 ^a Active ingredients (% A.I.): Actigard = acibenzolar-S-methyl (50%); Aliette = aluminum tris (0-ethyl phosphanate) (80%); Empress
828 Intrinsic = pyraclostrobin (23.3%); Interface Stressgard = trifloxystrobin (1.44%) + iprodione (23.1%); Stargus = *Bacillus*
829 *amyloliquefaciens* strain F727; ON-Gard[®] 5-0-0 = Total Nitrogen (5%); Orkestra Intrinsic = pyraclostrobin (21.26%) + fluxapyroxad
830 (21.26%); Pageant Intrinsic = pyraclostrobin (12.8%) + boscalid (25.2%); RootShield Plus⁺ = *Trichoderma harzianum* Rifai strain T-

831 22 (1.15%) + *T. virens* strain G-41 (0.61%); Segovis = oxathiapiprolin (18.7%); Signature Xtra Stressgard = aluminum tris (0-ethyl
832 phosphanate) (60%); Subdue MAXX = mefenoxam (22%); Tartan Stressgard = trifloxystrobin (4.17%) + triadimefon (20.86%).

833 ^b BASF=BASF Corporation, Florham Park, NJ; Bayer=Bayer AG, Monheim an Rhein, Germany; BioWorks=BioWorks Inc., Victor,
834 NY; Marrone =Marrone Bio Innovations, Inc., Davis, CA; Syngenta=Syngenta International AG, Basel, Switzerland

835 **Table 2.** Percent recovery of *Phytophthora cinnamomi* from root samples (mean ± SE) treated with preventively applied fungicides,
 836 biofungicides, fertilizer or host plant defense inducers 3 weeks before flooding.

Treatment ^a	Average (Mean ± SE) percent recovery of <i>P. cinnamomi</i> from root samples of preventative application 3 weeks before flooding at different flood durations					
	Trial 1			Trial 2		
	1 day	3 days	7 days	1 day	3 days	7 days
Actigard	60.00 ± 8.94 ab	43.33 ± 8.03 abc	43.33 ± 9.55 ab	60.00 ± 13.66 ab	46.67 ± 9.89 bcd	53.33 ± 9.89 bcd
Aliette	46.67 ± 6.67 a-d	40.00 ± 7.30 abc	26.67 ± 4.22 b	50.00 ± 12.38 ab	23.33 ± 8.03 cde	36.67 ± 8.03 de
Empress	43.33 ± 12.02 a-e	46.67 ± 16.06 abc	36.67 ± 10.85 ab	23.33 ± 15.85 cd	23.33 ± 13.08 ef	56.67 ± 6.15 cd
Interface	26.67 ± 4.22 de	53.33 ± 11.16 a	43.33 ± 6.15 ab	36.67 ± 9.55 bc	46.67 ± 8.43 b	70.00 ± 11.25 abc
ON-Gard	40.00 ± 11.55 a-e	50.00 ± 12.38 ab	33.33 ± 6.67 b	43.33 ± 14.06 bc	41.67 ± 9.1 bcd	60.00 ± 10.33 bcd
Orkestra	36.67 ± 6.15 b-e	46.67 ± 11.16 abc	40.00 ± 11.55 ab	30.00 ± 6.83 bc	23.33 ± 6.15 cde	63.33 ± 8.03 bcd
Pageant	40.00 ± 5.16 a-e	50.00 ± 11.25 abc	43.33 ± 6.15 ab	30.00 ± 4.47 bc	43.33 ± 9.55 bc	53.33 ± 11.16 bcd
RootShield Plus ⁺	46.67 ± 4.22 a-d	53.33 ± 16.87 a	46.67 ± 9.89 ab	66.67 ± 8.43 ab	33.33 ± 8.43 b-e	76.67 ± 10.85 ab
Segovis	36.67 ± 8.03 b-e	26.67 ± 6.67 a-d	46.67 ± 8.43 ab	36.67 ± 14.98 bc	36.67 ± 6.15 bcd	36.67 ± 6.15 de
Signature Xtra	33.33 ± 9.89 cde	30.00 ± 8.56 bcd	46.67 ± 12.29 ab	43.33 ± 13.08 bc	23.33 ± 9.55 def	53.33 ± 12.29 bcd
Stargus	56.67 ± 8.03 abc	40.00 ± 5.16 abc	36.67 ± 8.03 ab	40.00 ± 16.93 bc	46.67 ± 8.43 b	73.33 ± 9.89 abc
Subdue MAXX	23.33 ± 3.33 de	26.67 ± 6.67 cd	33.33 ± 8.43 b	23.33 ± 9.55 cd	16.67 ± 6.15 ef	26.67 ± 9.89 ef
Tartan	33.33 ± 4.22 cde	30.00 ± 6.83 bcd	46.67 ± 6.67 ab	43.33 ± 12.02 abc	40.00 ± 7.3 bcd	66.67 ± 11.16 abc
Non-inoculated control	20.00 ± 12.65 e	10.00 ± 4.47 d	3.33 ± 3.33 c	0.00 ± 0.00 d	0.00 ± 0.00 f	0.00 ± 0.00 f
Inoculated control	63.33 ± 10.85 a	60.00 ± 5.16 ab	56.67 ± 6.15 a	76.67 ± 8.03 a	80.00 ± 8.94 a	93.33 ± 4.22 a
<i>F</i>	2.37	1.84	2.19	2.59	5.11	5.41
df	14	14	14	14	14	14
<i>P</i>	0.0089	0.0477	0.0157	0.0042	<0.0001	<0.0001

837 ^a For each plant (six replications per treatment), ten randomly selected flowering dogwood root samples were plated on PARPH-V8

838 *Phytophthora*-selective medium to determine the percent recovery of *P. cinnamomi* from root samples.

839

840 **Table 3.** Percent recovery of *Phytophthora cinnamomi* from root samples (mean \pm SE) treated with preventively applied fungicides,
841 biofungicides, fertilizer or host plant defense inducers 1 week before flooding.

Treatment ^a	Average (Mean \pm SE) percent recovery of <i>P. cinnamomi</i> from root samples of preventative application 1 week before flooding at different flood durations					
	Trial 1			Trial 2		
	1 day	3 days	7 days	1 day	3 days	7 days
Actigard	26.67 \pm 8.43 bc	26.67 \pm 4.22 bc	63.33 \pm 12.02 b-f	66.67 \pm 6.67 ab	56.67 \pm 8.03 bc	86.67 \pm 6.67 ab
Aliette	13.33 \pm 4.22 bcd	23.33 \pm 6.15 cd	40.00 \pm 10.33 f	53.33 \pm 11.16 abc	53.33 \pm 6.67 bc	73.33 \pm 9.89 a-d
Empress	10.00 \pm 4.47 cd	36.67 \pm 6.15 bc	56.67 \pm 10.85 def	30.00 \pm 15.28 cd	36.67 \pm 10.85 cde	63.33 \pm 14.98 c-f
Interface	23.33 \pm 6.15 bc	30.00 \pm 8.56 bcd	80.00 \pm 12.65 abc	43.33 \pm 12.02 bcd	53.33 \pm 11.16 ab	86.67 \pm 6.67 ab
ON-Gard	26.67 \pm 8.43 bc	36.67 \pm 14.98 bcd	73.33 \pm 8.43 a-e	26.67 \pm 12.29 de	53.33 \pm 13.33 bc	66.67 \pm 16.06 ab
Orkestra	10.00 \pm 4.47 cd	26.67 \pm 12.29 cde	50.00 \pm 11.25 ef	33.33 \pm 13.33 bcd	56.67 \pm 12.02 bc	60.00 \pm 7.3 c-f
Pageant	16.67 \pm 8.03 bcd	26.67 \pm 12.29 cde	66.67 \pm 6.67 c-f	46.67 \pm 11.16 bcd	50.00 \pm 15.28 bcd	50.00 \pm 11.25 def
RootShield Plus ⁺	26.67 \pm 6.67 bc	36.67 \pm 12.02 bc	13.33 \pm 4.22 g	40.00 \pm 13.66 bcd	50.00 \pm 16.12 bcd	46.67 \pm 9.89 ef
Segovis	16.67 \pm 6.15 bcd	20.00 \pm 8.94 cde	83.33 \pm 9.55 ab	40.00 \pm 10.33 bcd	36.67 \pm 8.03 b-e	26.67 \pm 13.33 gh
Signature Xtra	16.67 \pm 6.15 bcd	6.67 \pm 4.22 de	16.67 \pm 6.15 g	36.67 \pm 8.03 bcd	26.67 \pm 9.89 def	60.00 \pm 11.55 b-f
Stargus	30.00 \pm 6.83 b	46.67 \pm 12.29 ab	73.33 \pm 9.89 a-d	33.33 \pm 14.3 bcd	60.00 \pm 5.16 ab	76.67 \pm 13.08 abc
Subdue MAXX	10.00 \pm 4.47 cd	10.00 \pm 4.47 cde	6.67 \pm 4.22 g	30.00 \pm 11.25 cd	20.00 \pm 8.94 ef	40.00 \pm 12.65 fg
Tartan	20.00 \pm 10.33 bc	27.5 \pm 10.47 cde	70.00 \pm 10 a-d	46.67 \pm 8.43 bcd	43.33 \pm 12.02 b-e	73.33 \pm 8.43 a-e
Non-inoculated control	0.00 \pm 0.00 d	0.00 \pm 0.00 e	0.00 \pm 0.00 g	0.00 \pm 0.00 e	0.00 \pm 0.00 f	0.00 \pm 0.00 h
Inoculated control	63.33 \pm 6.15 a	70.00 \pm 6.83 a	90.00 \pm 6.83 a	83.33 \pm 6.15 a	90.00 \pm 4.47 a	93.33 \pm 4.22 a
<i>F</i>	4.89	3.06	9.05	2.64	4.24	6.25
df	14	14	14	14	14	14
<i>P</i>	<0.0001	0.0009	<0.0001	0.0036	<0.0001	<0.0001

842 ^a For each plant (six replications per treatment), ten randomly selected flowering dogwood root samples were plated on PARPH-V8

843 *Phytophthora*-selective medium to determine the percent recovery of *P. cinnamomi* from root samples.

844

845 **Table 4.** Percent recovery of *Phytophthora cinnamomi* from root samples (mean ± SE) treated with curatively applied fungicides or
 846 host plant defense inducers.

Treatment ^a	Average (Mean ± SE) percent recovery of <i>P. cinnamomi</i> from root samples of curative application at different flood durations					
	Trial 1			Trial 2		
	1 day	3 days	7 days	1 day	3 days	7 days
Aliette	10.00 ± 6.83 bc	33.33 ± 8.43 ab	50.00 ± 13.42 a-d	30.00 ± 11.25 bc	23.33 ± 8.03 bcd	46.67 ± 12.29 abc
Empress	20.00 ± 10.33 bc	23.33 ± 12.02 ab	63.33 ± 12.02 c-f	23.33 ± 9.55 bc	23.33 ± 9.55 bcd	33.33 ± 8.43 bc
Interface	30.00 ± 19.15 ab	36.67 ± 14.06 a	53.33 ± 12.29 abc	16.67 ± 8.03 bc	20.00 ± 8.94 bcd	36.67 ± 10.85 bc
Orkestra	10.00 ± 6.83 bc	43.33 ± 17.45 a	60.00 ± 8.94 ab	26.67 ± 9.89 bc	53.33 ± 14.3 ab	50.00 ± 11.25 ab
Pageant	10.00 ± 4.47 bc	30.00 ± 15.28 ab	30.00 ± 11.25 def	20.00 ± 5.16 bc	36.67 ± 14.06 bc	43.33 ± 12.02 ab
Segovis	16.67 ± 6.15 bc	20.00 ± 10.33 ab	20.00 ± 7.3 efg	23.33 ± 8.03 bc	20.00 ± 7.3 bcd	36.67 ± 9.55 bc
Signature Xtra	6.67 ± 4.22 bc	36.67 ± 15.85 a	23.33 ± 8.03 efg	20.00 ± 10.33 bc	36.67 ± 12.02 bc	53.33 ± 8.43 ab
Subdue MAXX	23.33 ± 6.15 bc	13.33 ± 6.67 ab	26.67 ± 16.06 fg	15.00 ± 8.06 bc	13.33 ± 6.67 cd	20.00 ± 5.16 cd
Tartan	10.00 ± 4.47 bc	40.00 ± 10.33 a	36.67 ± 6.15 b-e	26.67 ± 8.43 b	36.67 ± 14.06 bc	40.00 ± 12.65 abc
Non-inoculated control	0.00 ± 0.00 c	0.00 ± 0.00 b	0.00 ± 0.00 g	0.00 ± 0.00 c	0.00 ± 0.00 d	0.00 ± 0.00 d
Inoculated control	50.00 ± 16.12 a	40.00 ± 10.33 a	66.67 ± 9.89 a	53.33 ± 11.16 a	73.33 ± 9.89 a	66.67 ± 8.43 a
<i>F</i>	2.18	1.25	4.14	2.30	2.83	3.90
df	10	10	10	14	14	10
<i>P</i>	0.0332	0.0028	0.0003	0.0246	0.0065	0.0005

847 ^a For each plant (six replications per treatment), ten randomly selected flowering dogwood root samples were plated on PARPH-V8

848 *Phytophthora*-selective medium to determine the percent recovery of *P. cinnamomi* from root samples.

849

850

851

852 **Figure Captions**

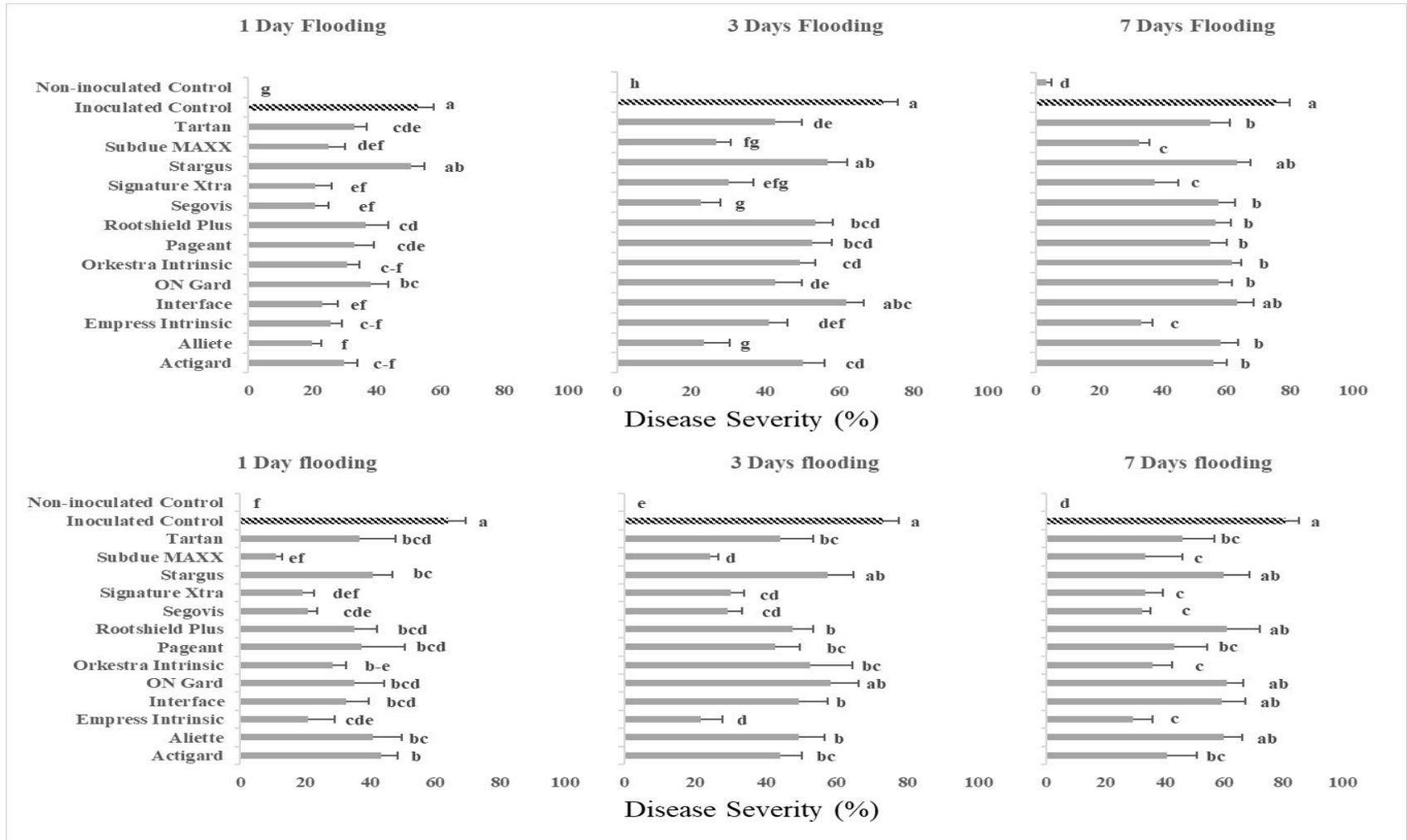
853 **Fig. 1.** Disease severity (mean \pm SE) of plants treated preventively 3 weeks before flooding with fungicides, biofungicides, fertilizer
854 or host plant defense inducers at 1-, 3-, or 7-days of flooding in trials 1 (top) and 2 (bottom). For root rot disease severity, each plant
855 was evaluated using a scale of 0–100% roots affected. Control treatments included the non-treated, inoculated and non-treated, non-
856 inoculated plants. Letters above the error bars represent significant differences in disease severity ($\alpha=0.05$, Fisher's LSD test).

857 **Fig. 2.** *Trichoderma* (mean \pm SE) colony counts on *Trichoderma*-selective medium at 1-, 3-, or 7-days of flooding in preventative
858 application 1 week (top) and 3 weeks (bottom) before flooding. RootShield Plus⁺-treated plants, undiluted root samples, as well as
859 dilutions of 10^{-2} and 10^{-4} , were plated on *Trichoderma*-selective medium, and the colonies were counted after 10 days of incubation.
860 Letters above bars represent significant differences in the number of *Trichoderma* colonies within flooding durations and trials
861 ($\alpha=0.05$, Least Squares Means).

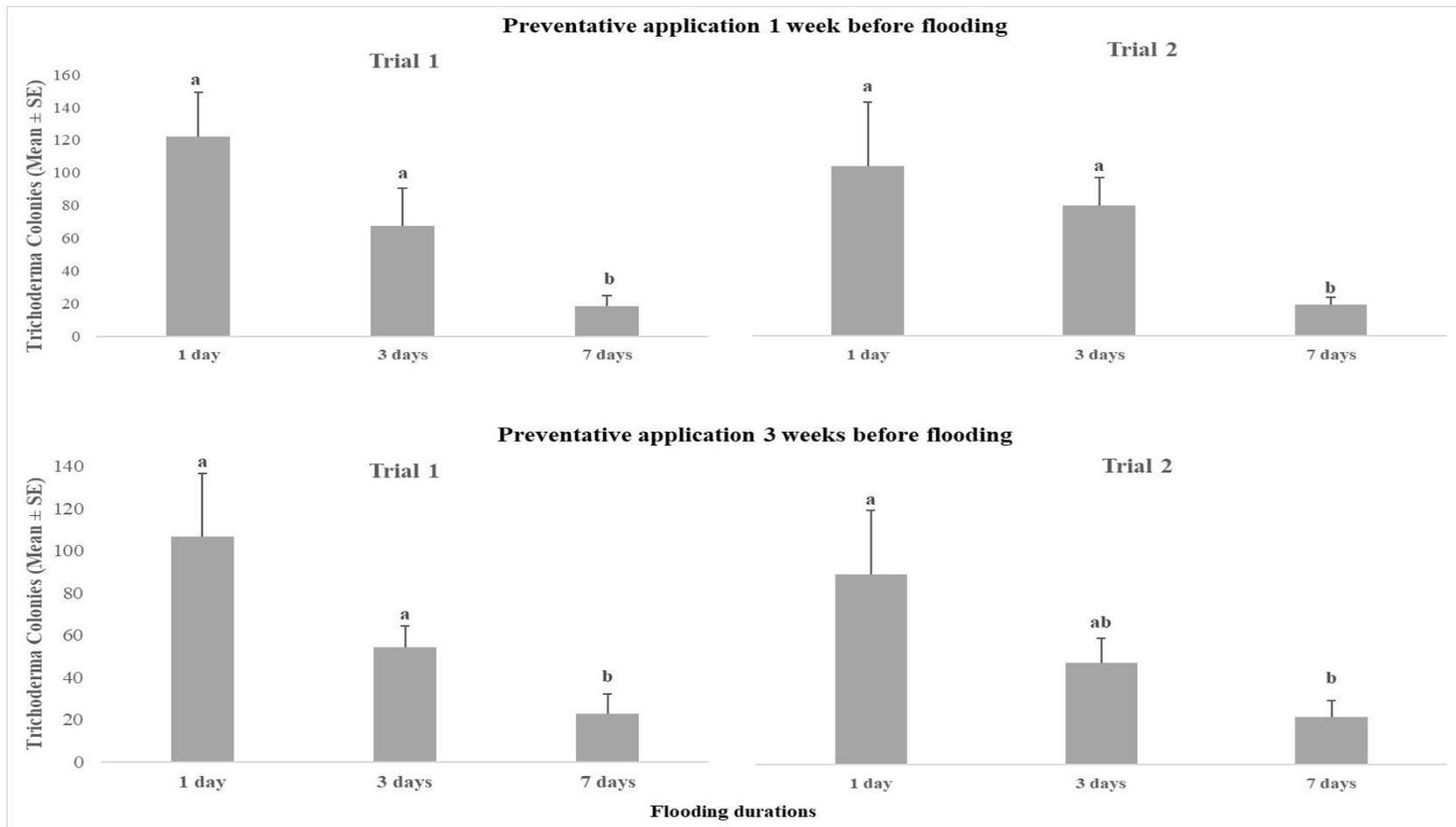
862 **Fig. 3.** Disease severity (mean \pm SE) of plants treated preventively 1 week before flooding with fungicides, biofungicides, fertilizer or
863 host plant defense inducers at 1-, 3-, or 7-days of flooding in trials 1 (top) and 2 (bottom). For root rot disease severity, each plant was
864 evaluated using a scale of 0–100% roots affected. Control treatments included the non-treated, inoculated, and non-treated, non-
865 inoculated plants. Letters above the error bars represent significant differences in disease severity ($\alpha=0.05$, Fisher's LSD test).

866 **Fig. 4.** Disease severity (mean \pm SE) of plants treated curatively with fungicides and host plant defense inducers at 1-, 3-, or 7-days of
867 flooding in trials 1 (top) and 2 (bottom). For root rot disease severity, each plant was evaluated using a scale of 0–100% roots affected.
868 Control treatments included the non-treated, inoculated, and non-treated, non-inoculated plants. Letters above the error bars represent
869 significant differences in disease severity ($\alpha=0.05$, Fisher's LSD test).

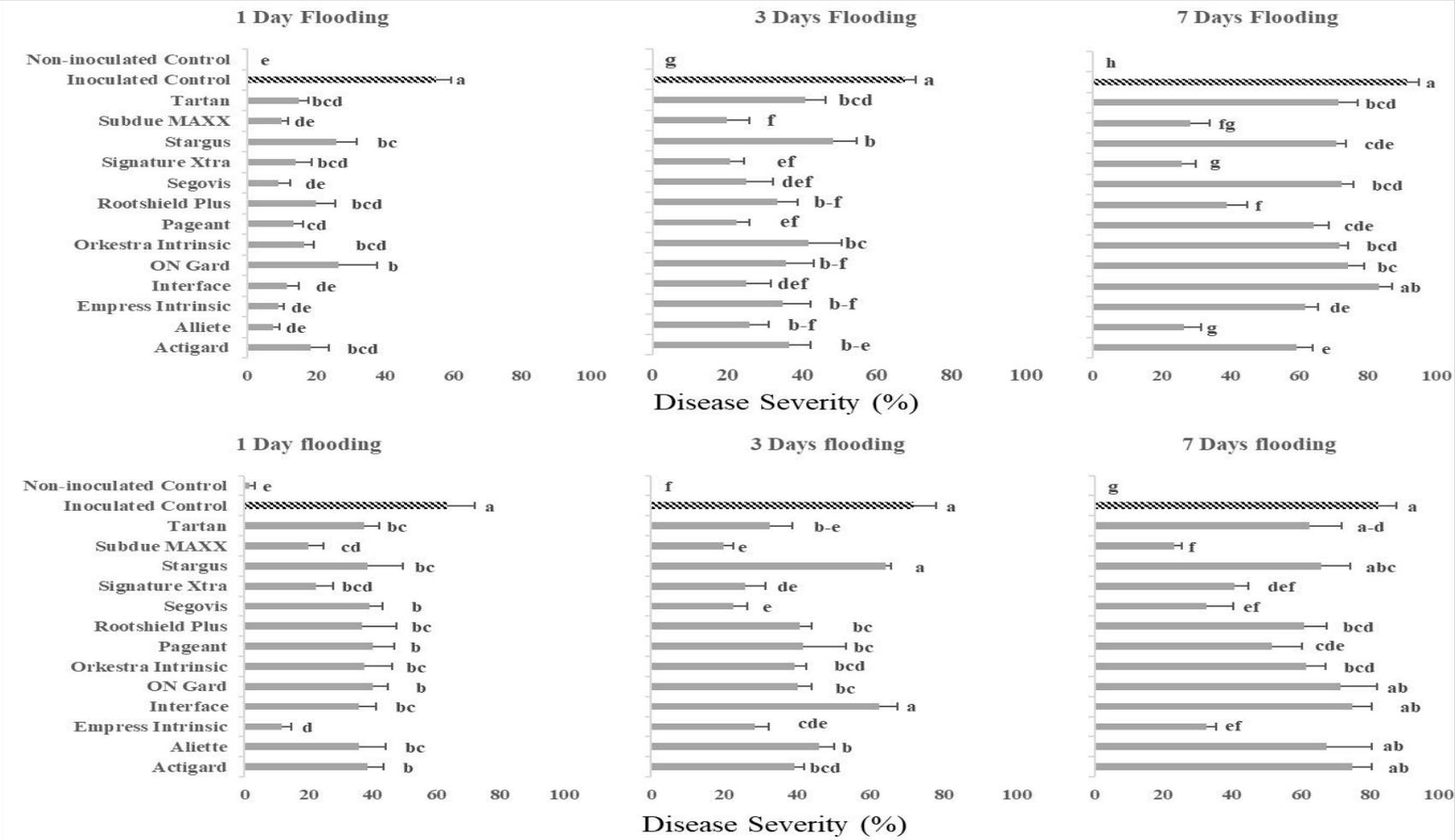
870



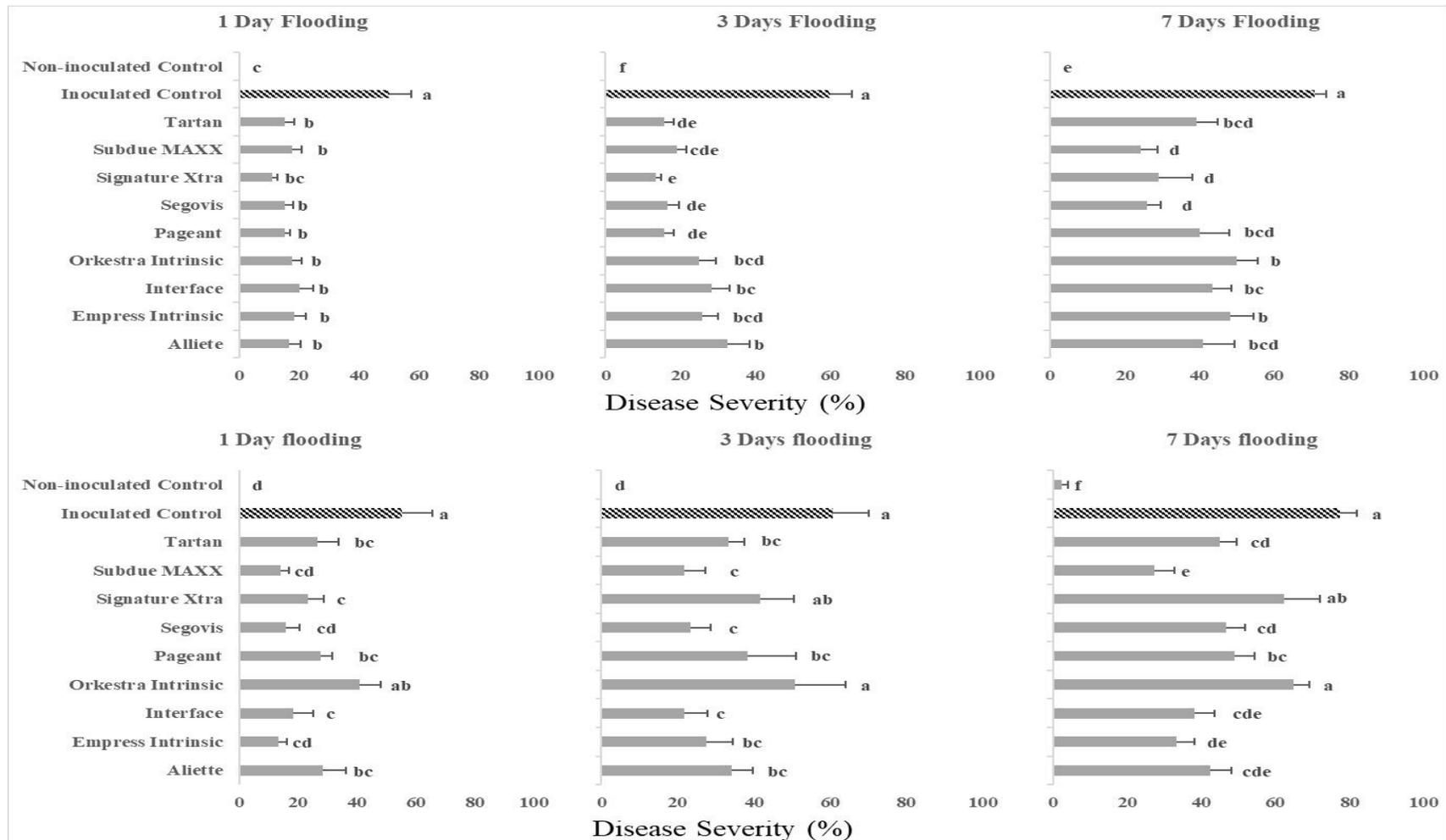
871 **Fig. 1.** Disease severity (mean ± SE) of plants treated preventively 3 weeks before flooding with fungicides, biofungicides, fertilizer
 872 or host plant defense inducers at 1-, 3-, or 7-days of flooding in trials 1 (top) and 2 (bottom). For root rot disease severity, each plant
 873 was evaluated using a scale of 0–100% roots affected. Control treatments included the non-treated, inoculated and non-treated, non-
 874 inoculated plants. Letters above the error bars represent significant differences in disease severity ($\alpha=0.05$, Fisher’s LSD test).



875 **Fig. 2.** *Trichoderma* (mean \pm SE) colony counts on *Trichoderma*-selective medium at 1-, 3-, or 7-days of flooding in preventative
 876 application 1 week (top) and 3 weeks (bottom) before flooding. RootShield Plus⁺-treated plants, undiluted root samples, as well as
 877 dilutions of 10^{-2} and 10^{-4} , were plated on *Trichoderma*-selective medium, and the colonies were counted after 10 days of incubation.
 878 Letters above bars represent significant differences in the number of *Trichoderma* colonies within flooding durations and trials
 879 ($\alpha=0.05$, Least Squares Means).



880 **Fig. 3.** Disease severity (mean ± SE) of plants treated preventively 1 week before flooding with fungicides, biofungicides, fertilizer or
 881 host plant defense inducers at 1-, 3-, or 7-days of flooding in trials 1 (top) and 2 (bottom). For root rot disease severity, each plant was
 882 evaluated using a scale of 0–100% roots affected. Control treatments included the non-treated, inoculated and non-treated, non-
 883 inoculated plants. Letters above the error bars represent significant differences in disease severity ($\alpha=0.05$, Fisher’s LSD test).



884 **Fig. 4.** Disease severity (mean \pm SE) of plants treated curatively with fungicides and host plant defense inducers at 1-, 3-, or 7-days of
 885 flooding in trials 1 (top) and 2 (bottom). For root rot disease severity, each plant was evaluated using a scale of 0–100% roots
 886 affected. Control treatments included the non-treated, inoculated and non-treated, non-inoculated plants. Letters above the error bars
 887 represent significant differences in disease severity ($\alpha=0.05$, Fisher's LSD test).