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Additional Information

1 **Cotton, cowpea and sesame are alternative crops to cucurbits in soils naturally infested**
2 **with *Monosporascus cannonballus***

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15

16 **ABSTRACT**

17 *Monosporascus cannonballus* is an important cucurbit root pathogen which has been
18 reported in the main production areas of melon and watermelon in Brazil and worldwide and
19 potentially capable to colonize roots of different species. Crop rotation is considered an
20 effective management strategy to prevent this disease. The aim of this study was to evaluate
21 the response of different crops, pumpkin, cotton, cowpea, sesame, watermelon, melon, corn,
22 cucumber, sorghum and tomato, to the infection of this pathogen. Seedlings were transplanted
23 into plastic containers with an inoculum concentration of 20 colony forming units (CFU) g⁻¹
24 of *M. cannonballus*. Fifty days after transplanting the variables analyzed were the degree of
25 disease severity on the root system and the frequency of reisolation. On cucurbits, the results

26 demonstrated different degrees of susceptibility among crops and cultivars, being melon and
27 watermelon the most sensitive species. In contrast, *Cucurbita* cultivars were the most tolerant.
28 Regarding non-cucurbit crops, maize, sorghum and tomato presented root discoloration and
29 *M. cannonballus* was reisolated from roots. Cotton, cowpea and sesame cultivars were not
30 affected by the pathogen, so they can be considered as alternative crops to be cultivated, or in
31 rotation with cucurbits, in *M. cannonballus* infested soils.

32

33 **KEYWORDS:** Crop rotation, host range, *Monosporascus* root rot and vine decline,
34 pathogenicity, soilborne pathogen.

35

36 1 INTRODUCTION

37 *Monosporascus cannonballus* Pollack & Uecker is an important cucurbit root
38 pathogen causing the disease known as "Monosporascus root rot and vine decline (MRRVD)"
39 (Martyn & Miller, 1996). This soilborne fungus has been reported in the main production
40 areas of melon (*Cucumis melo* L.) and watermelon [*Citrullus lanatus* (Thunb.) Matsum &
41 Nakai] cultivation in Brazil (Sales Júnior et al., 2004, 2010) and in other 21 countries
42 worldwide (Al-Mawaali, Al-Sadi, Al-Said, & Deadman, 2013; Cohen, Pivonia, Crosby, &
43 Martyn, 2012b; Yan, Zang, Huang, & Wang, 2016).

44 Root rot caused by *M. cannonballus* is part of a complex syndrome where this fungus
45 can be isolated alone or in association with other soilborne pathogens, such as *Acrocalymma*
46 *vagum* Crous & Trakunyingcharoen (Armengol, Vicent, Martínez-Culebras, Bruton, & García
47 Jiménez, 2003; Farr, Miller, & Bruton, 1998), *Fusarium oxysporum* f. sp. *melonis* Snyder &
48 Hansen, *Macrophomina phaseolina* (Tassi) Goid. (Cohen, Elkabetz, & Edelstein, 2016;
49 Cohen, Omari, Porat, & Edelstein, 2012a), *Fusarium solani* f. sp. *cucurbitae* Snyder &
50 Hansen, *Olpidium* spp. (Aleandri et al., 2017; Cara et al., 2008; Stanghellini & Misaghi, 2011;

51 Stanghellini, Mohammadi & Adaskaveg, 2014), *Pythium spinosum* Swada and *Rhizoctonia*
52 *solani* Kühn (Al-Sadi et al., 2011), and *Plectosphaerella melonis* (Watan & Sato) Phillips,
53 Carlucci & Raimondo (Armengol et al., 1998; Bruton, Davis, & Gordon, 1995). In Brazil, this
54 syndrome is considered an important limiting factor for cucurbits cultivation (Bezerra et al.,
55 2013).

56 *Monosporascus cannonballus* is a thermophilic fungus, which seems to be adapted to
57 Arid and Semi-arid climates, surviving in the soil in the absence of suitable hosts for long
58 periods in the form of ascospores (Medeiros, Silva, Oliveira, Ferreira, & Sales Júnior, 2008).
59 The symptoms associated with root rot caused by *M. cannonballus* can be easily observed on
60 melon plants close to harvest (Cohen, Pivonia, Crosby, & Martyn, 2012b), where severe vine
61 decline is observed. This is due to the rotting of the root system, which can no longer supply
62 the water needs of the plant, leading it frequently to its death. In addition, the affected root
63 system shows the presence of black perithecia from which abundant ascospores are produced,
64 being the main fungus reproduction structures (Louws, Rivarda, & Kubota, 2010).

65 Several studies have reported different management strategies to control MRRVD,
66 such as the use of green fertilization (Sales Júnior, Senhor, Michereff, & Medeiros, 2017),
67 application of fumigants (Stanghellini et al., 2003), destruction of postharvest plant residues
68 (Radewald, Ferrin, & Stanghellini, 2004), chemical control (Pivonia, Gerstl, Maduel, Levita,
69 & Cohen, 2010), application of essential oils (Fernandes et al., 2015; Awad, 2016), the use of
70 plant-growth promoting bacteria (Antonelli, xxx) and antagonistic biocontrol agents (Zhang,
71 1999; Júnior, 2007; Aleandri 2015), and grafting on *Cucurbita* hybrid rootstocks (Al-
72 Mawaali, Al-Sadi, Al-Said, & Deadman, 2016; Beltrán, Vicent, García-Jiménez, &
73 Armengol, 2008; Edelstein et al., 2017). However, some of these techniques are not sufficient
74 if applied alone, but can be effective when sustainable measures are integrated (Medeiros,

75 Silva, Oliveira, Ferreira, & Sales Júnior, 2008). In this sense, the practice of alternative
76 management techniques such as crop rotation needs to be elucidated.

77 Previous studies have demonstrated that, in addition to the Cucurbitaceae family, *M.*
78 *cannonballus* has been reported on roots of *Iris* sp., *Trifolium pratense* L. (red clover),
79 *Medicago sativa* L. (alfalfa) and *Sesamum indicum* L. (sesame) (Sivanesan, 1991), *Triticum*
80 sp. and *Achyranthes aspera* L. (Hawksworth, & Ciccarone, 1978) and *Lepidium lasiocarpum*
81 Nutt. (Stanghellini, Kim, & Rasmussen, 1996) in field samples. In addition, *M. cannonballus*
82 was isolated from artificially inoculated roots of *Zea mays* L. (corn), *Sorghum bicolor* L.
83 (Moench) (sorghum), *Beta vulgaris* L. (beet), *M. sativa*, *Triticum aestivum* L. (wheat) and
84 *Phaseolus vulgaris* L. (bean) (Mertely, Martyn, Miller, & Bruton, 1993). Therefore, it is
85 important to evaluate the response of different crop species to *M. cannonballus* in order to
86 determine which of them could be used in a crop rotation management program for this
87 disease in Brazil and also in other cucurbit growing areas severely affected by this pathogen.
88 In this sense, this study aims to assess the severity reaction on the root system of different
89 selected cucurbit and non-cucurbit crops after the inoculation with two isolates of *M.*
90 *cannonballus*.

91

92 **2 MATERIALS AND METHODS**

93 **2.1 *M. cannonballus* isolates and inoculum preparation**

94 Two *M. cannonballus* isolates were used in this study: CMM 2390 and CMM 3646,
95 obtained from melon and *Boerhavia diffusa* L., respectively, which were deposited in the
96 culture collection Prof. Maria Menezes of the Universidade Federal Rural de Pernambuco -
97 UFRPE: (Pernambuco, Brazil). Previous trials demonstrated that these isolates were
98 pathogenic to melon (Rodrigues, 2013).

99 Fungal inoculum was produced following the methodology described by Armengol
100 et al. (1998). Cultures were grown on potato-dextrose agar (PDA) at 26°C prior to
101 introduction to a sand-oat hull (*Avena sativa* L.) medium (0.5 L sand, 46 g ground oat hulls,
102 37.5 mL distilled water). The medium was mixed and transferred to 1 L flasks, autoclaved on
103 3 successive days, then inoculated with each fungal isolate. When the colonized material was
104 about 5 cm in diameter, the flasks were shaken to distribute the fungus evenly throughout the
105 mix and incubated at 25–30°C for 21–28 days. Following incubation, colony-forming units
106 (CFU) were quantified by serial dilution using 1% hydroxyethyl cellulose.

107

108 **2.2 Pathogenicity tests**

109 Pathogenicity tests were conducted in a greenhouse at Mossoró, State of Rio Grande
110 do Norte (RN); coordinates (5°11'15"S and 37°20'39" W, 18 m altitude).

111 The cultures and cultivars used in this experiment were melon: 'Goldex' and 'SF-69';
112 watermelon: 'Crimson Sweet' and 'Sugar Baby'; *Cucumis sativus* L. (cucumber): 'Aodai' and
113 'Marketer'; *Cucurbita* sp. (pumpkin): 'Bahiana' and 'Moranga'; *Solanum lycopersicum* L.
114 (tomato): 'Santa Clara' and 'Santa Cruz'; *Gossypium hirsutum* L. (cotton): 'BRS 286' and 'BRS
115 335'; sesame: 'Seda' and 'G4'; corn: 'BRS 205' and 'AG 7098'; sorghum: 'Ponta Negra' and
116 'Santa Elisa'; and *Vigna unguiculata* (L.) Waup. (cowpea): 'BRS Cauamé' and 'BRS Itaim'.
117 The non-cucurbit crops were selected because of their frequent use in the cucurbit off season
118 in the melon and watermelon producing region in the Brazilian states of RN and Ceará (CE).

119 Two separate experiments were carried out, one for each *M. cannonballus* isolate.
120 The experimental design was completely randomized with 20 treatments and four replicates
121 per experiment, being the experimental unit composed by a potted plant.

122 Seeds from each crop and cultivar were surface disinfected with sodium hypochlorite
123 (2.5% active chlorine) for 1 min and then seeded in expanded polystyrene trays containing

124 128 cells and filled with sterile Tropstrato® substrate. The seedlings were transplanted 9 days
125 after sowing into plastic containers with a capacity of 2 L, filled with a 1: 1: 1 sterile mixture
126 of soil, Tropstrato® substrate and washed sand, previously autoclaved at 120°C for 1 h.

127 In each of the replicates, before inoculation, an inoculum concentration of 20 colony
128 forming units (CFU) g⁻¹ of the respective *M. cannonballus* isolate was added to the soil (Sales
129 Júnior, Vicent, Armengol, García-Jiménez, & Kobori, 2002). Subsequently, the containers
130 were incubated in a greenhouse under controlled conditions of 30-35°C and relative humidity
131 70% ± 2.

132

133 **2.3 Disease severity evaluation**

134 Fifty days after transplanting, the entire plants were collected carefully and the roots
135 washed with running water to remove adhered soil remains. Then, the degree of severity
136 reaction on the root system was evaluated using the score scale from 0 to 4 described by
137 Armengol et al. (1998), where 0 = healthy roots; 1 = mild discoloration, 2 = moderate
138 discoloration with few lesions; 3 = severe discoloration with abundant lesions and 4 = totally
139 deteriorated. Then, resistance classes were assigned to the results of severity obtained, being:
140 0-1.0 = highly resistant; 1.01-2.0 = resistant; 2.01-3.0 = susceptible and 3.01-4.0 = highly
141 susceptible.

142

143 **2.4 Frequency of reisolation**

144 Fungal isolation was conducted after disease severity evaluation in PDA with the
145 addition of 500 ppm of streptomycin sulphate (PDAS). In each plant, seven small root
146 fragments were taken from affected areas, and then plated in one PDAS Petri dish. Plates
147 were incubated at 27-29°C in darkness for a five days period.

148 The growth of *M. cannonballus* colonies was assessed, and the frequency of isolation
149 per treatment was determined using the following formula: Frequency = (F x 100) / TF, being
150 F the number of fragments from which *M. cannonballus* was obtained and TF the total
151 fragments plated in culture medium.

152

153 **2.5 Statistical analysis**

154 Severity results were analyzed with to the non-parametric Kruskal-Wallis test at the
155 probability level of 5% ($p < 0.05$) using the software Assistat, version 7.7 (Silva & Azevedo,
156 2016).

157

158 **3 RESULTS**

159 **3.1 Disease severity**

160 Inoculation with *M. cannonballus* isolate CMM 2390 caused significant statistical
161 effect on root disease severity among the different cultivars ($\chi^2 = 68.38$; $p < 0.05$) (Table 1).

162 The cultivars tested were grouped in the four classes of severity reaction, 30% of
163 which were considered highly susceptible: melon: 'Goldex' (mean disease severity 3.50) and
164 'SF-69' (3.25); watermelon: 'Sugar Baby' (3.50); cucumber: 'Marketer' (3.75) and 'Aodai'
165 (4.00); and Pumpkin: 'Bahiana' (3.75). The same percentage (30%) was obtained for
166 watermelon: 'Crimson Sweet' (3.00); pumpkin: 'Moranga' (2.50); corn: 'BRS 205' (2.50) and
167 'AG 7098' (2.75); and tomato: 'Santa Cruz' (3.00) and 'Santa Clara' (3.00), which were
168 considered susceptible to inoculation with isolate CMM 2390. In total 60% of the cultivars
169 tested were classified as susceptible and highly susceptible to isolate CMM 2390. The
170 sorghum cultivars: 'BRS Ponta Negra' and 'BRS Santa Elisa' and the sesame cultivars: 'G4'
171 and 'Seda' were considered resistant to isolate CMM 2390, with mean disease severity values
172 of 1.75; 1.50; 1.50 and 1.50, respectively. The other cultivars tested, cowpea: 'BRS Cauamé'

173 and 'BRS Itaim' and cotton: 'BRS 335' and 'BRS 286' were considered highly resistant,
174 obtaining mean disease severity values of 0.00; 0.00; 1.00 and 1.00, respectively (Table 1).

175 Similar results were obtained when the same cultivars were inoculated with *M.*
176 *cannonballus* isolate CMM 3646. In this case, statistical analysis also confirmed significant
177 difference among the cultivars ($\chi^2 = 65.62, p < 0.05$) (Table 1).

178 Of the cultivars tested, 25% resulted highly susceptible, being: melon: 'Goldex' (3.25)
179 and 'SF-69' (mean disease severity 3.25); watermelon: 'Crimson Sweet' (3.25); cucumber:
180 'Aodai' (3.75); and pumpkin: 'Bahiana' (3.50). The following cultivars: watermelon 'Sugar
181 Baby' (3.00); cucumber: 'Aeketer' (2.75), pumpkin: 'Moranga' (2.50); corn: 'AG 7098' (2.50);
182 and tomato: 'Santa Cruz' (3.00) and 'Santa Clara' (3.00) were considered susceptible to *M.*
183 *cannonballus* isolate CMM 3646. In total, 55% of the cultivars tested were rated as
184 susceptible and highly susceptible to this isolate. In contrast, 45% of the cultivars tested were
185 considered resistant or highly resistant. Sorghum cultivars: 'BRS Ponta Negra' and 'BRS Santa
186 Elisa'; corn: 'BRS 205'; sesame: 'G4'; and cotton: 'BRS 335' were resistant and showed disease
187 severity values of 1.75; 1.50; 2.00; 1.25 and 1.50, respectively. The other cultivars tested
188 resulted highly resistant: cowpea: 'BRS Cauamé' (0.00) and 'BRS Itaim' (0.00); sesame: 'Seda'
189 (1.00); and cotton 'BRS 286' (1.00) (Table 1).

190

191 **3.2 Reisolation frequency**

192 Reisolation frequency of the *M. cannonballus* isolate CMM 2390 from the roots of the
193 inoculated cultivars presented the highest values for cucurbit cultivars: watermelon: 'Crimson
194 Sweet' (85.7%) and 'Sugar Baby' (53.6%); melon: 'Goldex' (53.6%) e 'SF-69' (42.8%);
195 cucumber: 'Marketer' (39.3%) and 'Aodai' (39.3%); pumpkin: 'Moranga' (39.3%). One
196 exception was the pumpkin cultivar 'Bahiana', which presented a low reisolation percentage
197 (10.7%).

198 In the group of non-cucurbit crops, cotton cultivars: 'BRS 335' (0.0%) and 'BRS 286'
199 (3.57%); sesame: 'G4' (0.0%) and 'Seda' (3.57%); cowpea: 'BRS Cauamé' (0.0%) and 'BRS
200 Itaim' (0.0%); and tomato: 'Santa Cruz' (3.57%) presented very low or null percentages of
201 reisolation. In contrast, corn cultivars: 'AG 7098' (53.6%) and 'BRS 205' (42.8%); sorghum
202 'BRS Ponta Negra' (28.6%); and tomato: 'BRS Santa Elisa' (14.3%) and 'Santa Clara' (21.4%),
203 showed variable colonization with this isolate of *M. cannonballus* (Table 2).

204 Results of reisolation frequency from plants inoculated with isolate CMM 3646 were
205 similar to those obtained with isolate CMM 2390. The highest reisolation values were
206 obtained for cucurbit cultivars watermelon: 'Crimson Sweet' (53.6%) and 'Sugar Baby'
207 (53.6%), melon: 'Goldex' (39.3%) and 'SF-69' (32.1%) and cucumber: 'Marketer' (42.9%) and
208 'Aodai' (42.9%). The exception were the pumpkin cultivars 'Bahiana' and Moranga, which
209 presented the same low reisolation percentage (7.1%). It was not possible to reisolate the
210 fungus from the non-cucurbit crops cowpea: 'BRS Cauamé' and 'BRS Itaim', sesame: 'G4' and
211 'Seda' and cotton: 'BRS 335' and 'BRS 286'. The reisolation percentage in sorghum: 'BRS
212 Santa Elisa' (3.57%), was lower than 5% and, in contrast, it was possible to reisolate *M.*
213 *cannonballus* from corn cultivars 'BRS 205' (28.6%) and 'AG 7098' (42.8%) and tomato:
214 'Santa Cruz' (7.14%) and 'Santa Clara' 10.7% (Table 2).

215

216 **4 DISCUSSION**

217 In this study, the Cucurbitaceae family showed the highest levels of root damage after
218 inoculation with *M. cannonballus*, being melon and watermelon the most sensitive species.
219 These results agree with previous research that already indicated melon and watermelon as the
220 most susceptible crops to this pathogen, although the cultivars used here are different from
221 those evaluated previously. In fact, pathogenicity studies conducted up to the present time
222 with commercial hybrids of melon and watermelon have not yet found any resistance to *M.*

223 *cannonballus* (Armengol et al., 1998; Davis et al., 2008; Martyn & Miller, 1996; Mertely,
224 Martyn, Miller, & Bruton, 1993; Sales Júnior, Vicent, Armengol, García-Jiménez, & Kobori,
225 2002; Wolff & Miller 1998). King, Davis, Zhang, & Crosby (2010) reported some melon
226 cultivars belonging to the types Conomom, Inodorus, Cantaloupensis and Agrestis as resistant
227 to *M. cannonballus*, but the fruits produced by them have no commercial value, presenting
228 low or no quality for the market.

229 Regarding reisolation frequency of *M. cannonballus*, also melon and watermelon
230 showed the highest values. In a similar work, Mertely, Martyn, Miller, & Bruton (1993)
231 obtained a reisolation percentage of *M. cannonballus* over 70% for the cultivars 'Black
232 Diamond' and 'Royal Sweet '(watermelon)', 'Magnum 45' and 'Honeydew Green Flesh'
233 (melon) and 'Poinsette 76 '(cucumber).

234 In our study the *Cucurbita* cultivars 'Bahiana' and 'Moranga' resulted susceptible to
235 both *M. cannonballus* isolates inoculated but, in contrast, the reisolation frequency was low.
236 Mertely, Martyn, Miller, & Bruton (1993) demonstrated the tolerance of *Cucurbita* cultivars
237 to *M. cannonballus*, because they presented relatively low values of isolation frequency, when
238 compared with those obtained with cucumber, melon and watermelon cultivars included in
239 their inoculation experiments. Although Alfaro-Fernández & García-Luis (2009)
240 demonstrated with histological studies that *M. cannonballus* is capable to infect *C. maxima*
241 tissues to some extent, subsequent studies have explored the good performance of *Cucurbita*
242 hybrid rootstocks for the management of MRRVD in field conditions (Al-Mawaali, Al-Sadi,
243 Al-Said, & Deadman, 2016; Beltrán, Vicent, García-Jiménez, & Armengol, 2008; Cohen,
244 Burger, Horev, Porat, & Edelstein, 2005; Demartelaere, Freitas, Soares, Queiroz, & Sales
245 Júnior, 2015; Edelstein, M., Cohen, R., Burger, Y., & Shriber, 1999; Kim et al., 2016; Louws,
246 Rivarda, & Kubota, 2010; Park et al., 2013).

247 Regarding non-cucurbit crops, our results were similar to that reported by Mertely,
248 Martyn, Miller, & Bruton (1993), who compared the susceptibility to *M. cannonballus* of
249 eight non-cucurbit crops: 'Pioneer 8358' (sorghum), 'Asgrow 405W' (corn), 'Rutgers' tomato,
250 'Paymaster 145' (cotton), 'Era' (wheat), 'Cimmaron' (alfalfa) and 'Improved Commodore'
251 (bean). Their results indicated that corn, wheat and tomato cultivars showed a slight
252 discoloration in the root system, as well as a slight reduction in the dry weight of tomato and
253 wheat roots. Perithecia of *M. cannonballus* were also observed in bean and sorghum roots,
254 although there was not a reduction in plant development. In a pathogenicity study with *M.*
255 *cannonballus* on Solanaceae species, Tsay & Tung (1997) reported a slight rot in the root
256 system in *S. lycopersicum*, *S. melongena* L. (eggplant), *Capsicum annuum* L. (pepper),
257 *Brassica oleracea* L. var. *italica* (broccoli) and *B. oleraceae* var. *capitata* L. (cabbage).

258 Other reports of *M. cannonballus* isolated from roots of non-cucurbit crops were
259 obtained from field surveys of *Iris* sp., *T. pratense*, *M. sativa*, *S. indicum* (Sivanesan, 1991),
260 *Triticum* sp. (Hawksworth & Ciccarone, 1978) and *L. lasiocarpum* (Stanghellini, Kim, &
261 Rasmussen, 1996). However, further pathogenicity studies with these hosts were not carried
262 out.

263 Regarding the isolation frequency for non-cucurbit crops, Mertely, Martyn, Miller, &
264 Bruton (1993) found the presence of *M. cannonballus* perithecia in roots of artificially
265 inoculated plants of corn and sorghum, with an isolation frequency of 33% and 40%,
266 respectively. These values were similar to that found here for maize, where re-isolation
267 percentages higher than 28% were observed for both cultivars and the two *M. cannonballus*
268 isolates. In the case of sorghum, *M. cannonballus* was re-isolated from roots, but with low
269 percentages. These results suggest the potential role of these crops as hosts of *M.*
270 *cannonballus*. On the other side, the same authors did not obtain re-isolation of *M.*
271 *cannonballus* from tomato and cotton. This fact, in the case of tomato, contradicts the results

272 obtained in our study, although the frequency of isolation of *M. cannonballus* did not exceed
273 25%. Later, Tsay & Tung (1997) studying the susceptibility of Solanaceae inoculated with *M.*
274 *cannonballus*, found reisolation percentages between 5 and 25% in tomato, eggplant, pepper,
275 broccoli and cabbage roots.

276 In our study the cultivars of cowpea, sesame and cotton were not affected by *M.*
277 *cannonballus*, being the results with cotton coincident to the results found by Mertely,
278 Martyn, Miller, & Bruton (1993).

279 The differences in pathogenicity exhibited by the two *M. cannonballus* isolates on
280 cucurbit species may be due to genetic variability, a factor that can configure specific and
281 differentiated degrees of virulence. According to Bruton (1998), there is considerable
282 variation in virulence among *M. cannonballus* isolates ranging from weakly virulent to highly
283 virulent. In Brazil, Andrade et al. (2005), classified *M. cannonballus* isolates obtained from
284 melon production areas of the states of Rio Grande do Norte and Ceará (CE), in three distinct
285 groups of similarity, based on mycelial compatibility grouping (MCG) study. In a similar
286 study, Bezerra et al (2013) assigned 58 isolates obtained from seven melon fields in three
287 municipalities of Northeastern Brazil into four MCGs. Subsequently, Correia et al. (2014)
288 investigated the fitness components of 57 isolates of *M. cannonballus* obtained from Brazilian
289 melon fields by evaluating their mycelial growth rate, perithecia and ascospore production,
290 sensitivity to the fungicide fluazinam and virulence to melon seedlings. A multivariate cluster
291 analysis allowed the separation of these isolates in 18 groups of similarity.

292 Our results present a great concern for the melon and watermelon producers in Brazil,
293 since corn and sorghum are the two main crops grown by them during the off-season, because
294 they profit from the remaining fertilization in the field left by melon and watermelon crops.
295 Thus, it is possible that maize and sorghum crops contribute to the *M. cannonballus* inoculum

296 build-up in the soil, but further research in field conditions is needed to confirm this
297 hypothesis.

298 The adoption of cultural practices such as crop rotation as a strategy contributing to
299 minimize the economic losses caused by the attack of *M. cannonballus* to melon and
300 watermelon crops should take into account the results here reported. Cotton, cowpea and
301 sesame cultivars were not affected by the pathogen, so they can be considered as the
302 recommended alternative crops to be cultivated, or in rotation with cucurbits, in *M.*
303 *cannonballus* infested soils. This technique can be effective when integrated with other
304 control measures for a sustainable MRRVD management.

305

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309

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479

480 TABLE 1. Reaction of cucurbit and non-cucurbit crops to *Monosporascus cannonballus*.

Host	Cultivar	Isolate CMM 2390 ^a			Isolate CMM 3646		
		Rank	Mean	RC ^b	Rank	Mean	RC
<i>Cucumis melo</i>	Goldex	61.75	3.50	HS	61.37	3.25	HS
	SF-69	56.62	3.25	HS	61.37	3.25	HS
<i>Citrullus lanatus</i>	Sugar Baby	61.75	3.50	HS	57.00	3.00	SU
	Crimson Sweet	51.50	3.00	SU	61.37	3.25	HS
<i>Cucumis sativus</i>	Markerter	66.87	3.75	HS	50.37	2.75	SU
	Aodai	72.00	4.00	HS	70.12	3.75	HS
<i>Cucurbita sp.</i>	Bahiana	66.87	3.75	HS	65.75	3.50	HS
	Moranga	41.62	2.50	SU	46.00	2.50	SU
<i>Vigna unguiculata</i>	BRS Cauamé	4.50	0.00	HR	4.50	0.00	HR
	BRS Itaim	4.50	0.00	HR	4.50	0.00	HR
<i>Sorghum bicolor</i>	BRS Ponta Negra	27.62	1.75	R	30.37	1.75	R
	BRS Santa Elisa	23.75	1.50	R	25.75	1.50	R
<i>Zea mays</i>	BRS 205	41.50	2.50	SU	35.00	2.00	R
	AG 7098	46.50	2.75	SU	46.00	2.50	SU
<i>Solanum lycopersicum</i>	Santa Cruz	51.50	3.00	SU	55.87	3.00	SU
	Santa Clara	51.62	3.00	SU	54.75	3.00	SU
<i>Sesamum indicum</i>	G4	23.75	1.50	R	21.12	1.25	R
	Seda	23.75	1.50	R	16.50	1.00	HR
<i>Gossypium hirsutum</i>	BRS 335	16.00	1.00	HR	25.75	1.50	R
	BRS 286	16.00	1.00	HR	16.50	1.00	HR
χ^2		68.38*			65.62*		

481 ^aisolates of *M. cannonballus*; ^bRC=reaction class to *M. cannonballus*: HR= highly resistant; R= resistant; SU=
482 susceptible; HS= highly susceptible (Armengol et al., 1998); χ^2 = chi-square value significant at 5% by Kruskal-
483 Wallis test.

484

485

486 TABLE 2. Frequency of isolation of *Monosporascus cannonballus* from 20 hosts inoculated
 487 with isolates CMM 2390 and CMM 3646.

Host	Cultivar	Isolate CMM 2390	Isolate CMM 3646
		% ^a	%
<i>Cucumis melo</i>	Goldex	53.6	39.3
	SF-69	42.8	32.1
<i>Citrullus lanatus</i>	Sugar Baby	53.6	53.6
	Crimson Sweet	85.7	53.6
<i>Cucumis sativus</i>	Markerter	39.3	42.9
	Aodai	39.3	42.9
<i>Cucurbita sp.</i>	Bahiana	10.7	7.10
	Moranga	39.3	7.10
<i>Vigna unguiculata</i>	BRS Cauamé	0.00	0.00
	BRS Itaim	0.00	0.00
<i>Sorghum bicolor</i>	BRS Ponta Negra	28.6	17.8
	BRS Santa Elisa	14.3	3.60
<i>Zea maiz</i>	BRS 205	42.8	28.6
	AG 7098	53.6	42.8
<i>Solanum lycopersicum</i>	Santa Cruz	3.57	7.14
	Santa Clara	21.4	10.7
<i>Sesamum indicum</i>	G4	0.00	0.00
	Seda	3.57	0.00
<i>Gossypium hirsutum</i>	BRS 335	0.00	0.00
	BRS 286	3.57	0.00

488 ^apercentage of 28 isolation points from which *M. cannonballus* was isolated.