

24 **Summary**

- 25 1. Induced changes in plant quality can mediate indirect interactions between herbivores.
26 Although the sequence of attack by has been shown to influence plant responses, little
27 is known about how this affects the herbivores themselves.
- 28 2. We therefore investigated how induction by the leaf-herbivore *Spodoptera frugiperda*
29 influences resistance of teosinte (*Zea mays mexicana*) and cultivated maize (*Zea mays*
30 *mays*) against root-feeding larvae of *Diabrotica virgifera*. The importance of the
31 sequence of arrival was tested in the field and laboratory.
- 32 3. *S. frugiperda* infestation had a significant negative effect on colonization by *D. virgifera*
33 larvae in the field and weight gain in the laboratory, but only when *S. frugiperda* arrived
34 on the plant before the root herbivore. When *S. frugiperda* arrived after the root
35 herbivore had established, no negative effects on larval performance were detected.
36 Yet, adult emergence of *D. virgifera* was reduced even when the root feeder had
37 established first, indicating that the negative effects were not entirely absent in this
38 treatment.
- 39 4. The extent of defoliation of the plants was not a decisive factor for the negative effects
40 on root herbivore development, as both minor and major leaf damage resulted in an
41 increase in root resistance and the extent of biomass removal was not correlated with
42 root-herbivore growth. We propose that leaf-herbivore induced increases in feeding-
43 deterrent and/or toxic secondary metabolites may account for the sequence-specific
44 reduction in root-herbivore performance.

45 5. **Synthesis:** Our results demonstrate that the sequence of arrival can be an important
46 determinant of plant-mediated interactions between insect herbivores in both wild and
47 cultivated plants. Arriving early on a plant may be an important strategy of insects to
48 avoid competition with other herbivores. To fully understand plant-mediated
49 interactions between insect herbivores, the sequence of arrival should be taken into
50 account.

51

52 Key-words: Above-BG interactions, *Diabrotica virgifera*, induced resistance, plant-mediated
53 effects, plant quality, *Spodoptera frugiperda*, systemic signalling, *Zea mays*, teosinte.

54

55 **Introduction**

56 The metabolism of plants is remarkably adaptable to environmental stress: Upon attack
57 by insects and pathogens, dedicated signal transduction cascades are activated that help plants
58 to withstand and tolerate the ensuing threats (Howe and Jander, 2008, Dangl and Jones, 2001;
59 Rasmann *et al.*, this issue). Such changes do not only happen locally, but involve non-attacked
60 tissues as well (Schwachtje and Baldwin, 2008, Orians, 2005, Erb *et al.*, 2009c, Heil and Ton,
61 2008). Systemic effects following herbivory can have fitness consequences for temporally or
62 spatially separated organisms (van Loon *et al.*, 1998, Erb *et al.*, 2009a, Sticher *et al.*, 1997,
63 Poelman *et al.*, 2008a, Viswanathan *et al.*, 2005). Interestingly, it is becoming more and more
64 evident that changes in plant quality may even be more important than direct interference or
65 biomass removal in shaping competitive interactions between herbivores and future attacker
66 communities (Kaplan and Denno, 2007, van Zandt and Agrawal, 2004, Poelman *et al.*, 2010).
67 Some of the most dramatic examples in this context come from studies investigating plant-
68 mediated interactions between root- and leaf- feeding herbivores (Erb *et al.*, 2008):
69 Belowground (BG) herbivores have been shown to profoundly change leaf physiology, thereby
70 affecting aboveground (AG) attackers, and even higher trophic levels (Steinger and Müller-
71 Schärer, 1992, van Dam *et al.*, 2005, Soler *et al.*, 2005, Rasmann and Turlings, 2007) and *vice*
72 *versa*, AG herbivores can change root physiology and resistance (Moran and Whitham, 1990,
73 Masters, 1995, Soler *et al.*, 2007, Kaplan *et al.*, 2008).

74 In recent years, it has been hypothesized that plant-quality mediated interactions
75 between herbivores may not only depend on the combination of attackers, but also on their
76 sequence of arrival or timing (Blossey and Hunt-Joshi, 2003). Evidence for this concept comes

77 for example from a gene-expression study in *Nicotiana attenuata*, where it was found that the
78 order of attack of a sap-feeder and a chewing herbivore is an important determinant explaining
79 the ensuing transcriptional response (Voelckel and Baldwin, 2004). In *Solanum dulcamara*,
80 changes in polyphenol oxidase and peroxidase activity following tortoise and flea beetle attack
81 were determined by the first attacker, but not significantly modified after sequential feeding by
82 either species (Viswanathan *et al.*, 2007). Yet, despite the increasing evidence for the sequential
83 dependence of changes in plant-quality following attack, we are not aware of any study that
84 has tested the effect of an herbivore arriving before or after a second feeder on the
85 performance of the latter. Such experiments are especially difficult to conduct in the AG parts
86 of plants, as simultaneously occurring herbivores may interact directly with each other
87 compared to their sequential presence, thereby confounding direct and plant-mediated effects.
88 As root- and leaf-herbivores are spatially separated and do not have any physical contact during
89 their development, they represent an ideal model to study the effects of the sequence of
90 arrival.

91 We tested the effect of the sequence of arrival on the impact of leaf-herbivory on root
92 herbivore resistance using leaf-feeding larvae of the specialist noctuid moth *Spodoptera*
93 *frugiperda* (J.E. Smith) and root feeding larvae of the specialist beetle *Diabrotica virgifera*
94 *virgifera* (LeConte). These species co-occur in maize (*Zea mays* L.) agroecosystems in North
95 America and natural ecosystems in Mexico. *D. virgifera* passes the winter and/or dry periods
96 as eggs in the soil, from where the larvae hatch, locate their hosts and start feeding. Larvae can
97 cross distances up to 1m to find or switch host plants (Short and Luedtke, 1970, Suttle *et al.*,
98 1967). *S. frugiperda* on the other hand overwinters as pupa in tropical regions and the southern

99 US (Foster and Cherry, 1987), from where adults disperse and oviposit on growing plants. In the
100 main maize growing regions of North America, *S. frugiperda* therefore establishes later on the
101 host than *D. virgifera* (O'Day, 1998). In Mexico, where teosinte (the wild ancestor of maize) and
102 *D. virgifera* are believed to have evolved together (Branson and Krysan, 1981), it can be
103 expected that plants may be attacked first by either herbivore, depending on which species is
104 faster in colonizing its host at the beginning of the growing season. Furthermore, as *D. virgifera*
105 displays an enormous phenotypic plasticity in its diapause behavior (Branson, 1976), late
106 emerging or second generation *D. virgifera* larvae may encounter plants that have already been
107 attacked by both *D. virgifera* and *S. frugiperda*.

108 A combination of field and laboratory experiments was used to gain insight into the leaf-
109 herbivore induced changes in root resistance and the importance of sequential colonization. In
110 the field, we simulated a natural situation whereby early emerging *D. virgifera* larvae arrived on
111 the plant first, followed by *S. frugiperda* in the leaves and a subsequent second wave of root
112 herbivores. In the laboratory, we explicitly tested if the sequence of arrival influences leaf-
113 herbivore induced changes by adding and removing *S. frugiperda* larvae either before or after
114 the onset of *D. virgifera* feeding. In the laboratory, we not only tested cultivated maize (*Zea*
115 *mays mays*), but also its wild ancestor teosinte (*Zea mays mexicana*). The complementary
116 assays presented here provide clear evidence for the importance of the sequence of arrival of
117 different insect herbivores for plant-mediated interactions between them.

118

119 **Material and Methods**

120 *Field plants and insects*

121 For the field experiments, maize seeds (var. Delprim) were sown in 16 plots (3.05 m ×
122 3.05 m). Plots were arranged in a 2 x 8 rectangular pattern. All plants were sown on the 1st of
123 June 2009. Because of low initial germination, most plots did not reach the envisaged density of
124 64 plants per plot. Therefore, new seeds were sown or seedlings were transplanted two weeks
125 later to fill the gaps. To insure that western corn rootworm larvae would not move between
126 plots, a 3.05 m buffer containing no vegetation was maintained between each plot within rows
127 and four rows of commercial buffer maize were planted between the two blocks of eight plots.
128 Four additional rows of buffer maize were also planted at both sides of the study site to
129 minimize wind damage to the screen tents. Eight plots suffered from flooding (2 times for
130 approx. 48 h) during the early stage of the experiment. A block factor (flooding) was added to
131 the statistical model to account for this potential source of variability (see below). All the plots
132 were infested with *D. virgifera* eggs (600 WCR actual eggs every 30.5 cm of maize row) on the
133 18th of June. A diapausing strain was used for this infestation. Viability of these eggs averaged
134 83%, so viable egg numbers were close to 500 per 30.5 cm of maize row. On the 3rd of July,
135 when the plants had reached a height of approx. 50 cm and had developed 6 leaves, screen
136 tents (3.35 m × 3.96 m Insta-Clip, The Coleman Company, Inc., Wichita, KS) were placed over
137 the plots to reduce the natural colonization of herbivores. The tents were dug into the soil to a
138 depth of 15 cm to help secure the tents from wind damage. On the 10th of July, half of the plots
139 were infested with 20 neonate *S. frugiperda* larvae/plant using a bazooka/corn grit applicator
140 system (Wiseman et al. 1980). Control plants received the same volume of corn grit without

141 larvae. Because of the high mortality of the neonates after the first application, another 20 *S.*
142 *frugiperda* larvae were added one week later using the same method. Forty *S. frugiperda* larvae
143 per plant are well within the natural range of infestation, as egg batches typically consist of 100
144 or more individuals. On the 22nd of July, when the *D. virgifera* larvae were in the second larval
145 stadium, 4-6 plants with clear caterpillar damage were selected and harvested from each plot.
146 On the 24th of July, when the first *D. virgifera* infestation began to reach the pupal stage and
147 the first maize plants were tasseling, another 500 WCR eggs were added to 8 plants per plot,
148 and the plants were marked for later recovery. These plants had previously been attacked by
149 early emerging *D. virgifera* larvae, followed by either *S. frugiperda* (“infested”) or no leaf-
150 herbivory (“controls”). A non-diapausing strain was used for the second infestation. This strain
151 is similar in many aspects to the diapausing *D. virgifera*, but develops somewhat faster on the
152 plants. This enabled a second, successful establishment of the root herbivore larvae on the
153 plants before they were too old (Hibbard *et al.*, 2008). We also hypothesized that in a natural
154 situation in Mexico, late arriving *D. virgifera* larvae would likely be second-generation
155 individuals that did not enter diapause. Two groups of plants were used for this second
156 application: One half that had already reached the tasseling stage and another half that were
157 still in the whorl stage due to late sowing or replanting. On the 7th of August, when the larvae of
158 the first infestation had pupated and the second *D. virgifera* infestation had reached the second
159 instar, the infested plants were harvested. To gain insight into the number of *D. virgifera* larvae
160 that were able to successfully develop to adult beetles, the remaining plants (around 50/plot)
161 were left in the tents until the end of the adult emergence period of the first infestation of *D.*

162 *virgifera*. The field experiment was terminated on the 20th of September, when a heavy storm
163 destroyed the tents.

164

165 *Recovery of D. virgifera larvae, root damage rating and adult emergence*

166 Plant root systems (4-8 per plot, see above) were harvested from the field by digging the
167 roots out together with the surrounding soil. The root balls were then transferred to
168 commercial onion bags and suspended in a greenhouse as described by Hibbard *et al.* (2004).
169 Under each bag, a plastic pan filled with water was installed. The high temperature in the
170 greenhouse (40-50° C) dried the soil balls and prompted the *D. virgifera* larvae to move down
171 and fall into the water below. Larvae were counted and recovered twice a day over a period of
172 10 days and preserved in ethanol. Roots were then washed and rated for damage using the 0 to
173 3 node-injury scale (Oleson *et al.*, 2005). Starting on the 7th of August, emergence of adult *D.*
174 *virgifera* beetles in the tents was monitored every week until the 16th of September. The
175 emerging insects were collected, sexed and preserved in ethanol.

176

177 *Laboratory plants and insects*

178 To confirm the results obtained in the field in a better controlled environment, we
179 carried out additional experiments in the laboratory. Cultivated maize and teosinte plants were
180 grown in bottom-pierced, aluminium-wrapped plastic pots (diameter, 4cm; depth, 11cm) in a
181 phytotron (23±2°C, 60% r.h., 16:8 hr L/D, and 50,000 lm/m²). Before planting, the seeds were
182 rinsed with water to remove any storage residuals. They were then sown in sand (lower 8 cm)
183 and topped with commercial potting soil (upper 3 cm, Ricoter Aussaaterde, Aarberg,

184 Switzerland). Cultivated maize plants (*Zea mays mays*, var. Delprim) had two fully expanded
185 primary leaves and were 9-10 days old. Teosinte seeds (*Zea mays mexicana*) had been collected
186 from two wild populations near Texcoco (Mexico) in 1998. As the teosinte plants grew slower
187 than the cultivated hybrid Delprim, they were left in the phytotron for 20 days, until they had 2-
188 3 fully developed leaves. All plants were watered with 10ml of tap water every day.
189 Experiments were carried out under light benches in a climatized laboratory ($25\pm 2^{\circ}\text{C}$, $40\pm 10\%$
190 r.h., 16:8 hr L/D, and 8000 lm/m^2). *S. frugiperda* eggs were obtained from an in-house colony
191 reared on artificial diet. *D. virgifera* eggs (non-diapausing strain) were obtained from the USDA-
192 ARS-NCARL Brookings (US) and kept on freshly germinated maize seedlings until use.

193

194 *D. virgifera* performance experiments

195 Laboratory experiments were carried out to specifically test whether physiological
196 changes in the plants are important for the differential effects of sequence of arrival for the
197 impact of *S. frugiperda* on *D. virgifera*. One experiment was performed using cultivated maize,
198 and a second one with teosinte. The following procedure was used for both trials: Before the
199 beginning of the experiments, the pots of 10 day old plants were covered at the bottom with
200 aluminium foil to prevent root herbivores from escaping through the two drainage holes in the
201 bottom of each pot, and transparent 1,5l PET bottles with their bottoms removed (30cm height,
202 conal shape, top-diameter: 8cm) were placed upside down over the AG part of the plants to
203 confine leaf-herbivores. The PET tubes were held in place with parafilm. The plants were then
204 divided into three groups (n=12-15). All groups were infested with 4 pre-weighed early second
205 instar *D. virgifera* larvae by putting them on the soil with a fine brush. One set of plants had

206 been infested with 12 L2 *S. frugiperda* larvae 48h prior to root herbivore infestation, while the
207 second set was infested with the leaf herbivore 48h after *D. virgifera* had started feeding. In
208 both cases, the *S. frugiperda* larvae were removed from the plants after 48h of feeding. The
209 third group did not receive any leaf-herbivore treatment. We had intended to add an additional
210 leaf-herbivore treatment to the teosinte experiment, but a lack of suitable *S. frugiperda* larvae
211 prevented this and we therefore had a teosinte control group that consisted of a total of 24
212 independent replicates. After five days of feeding, the *D. virgifera* larvae were recovered from
213 the soil and weighed to determine their weight increase. Leaves of the different plants were
214 harvested and their fresh weight (FW) was determined.

215

216 *Data analysis*

217 For the field experiment, the parameters recorded were averaged for the different
218 plots, resulting in eight independent replicate values per treatment. Two-way Analyses of
219 Variance (ANOVAs) were carried out on the number of recovered root herbivore larvae and
220 emerging adults with the factors treatment and environment. The environment was either
221 “flooded” (8 plots) or “non-flooded” (8 plots) depending on the soil-water condition within the
222 field tents, and the two treatments were “control” (8 plots) and “*S. frugiperda* infested” (8
223 plots). Interaction terms were included in the models. To assess the effect of big and small
224 plants, plant size was included as a nested factor in a general linear model (GLM). Larval growth
225 and leaf fresh-weight in the lab-experiment were assessed using one-way ANOVAs. In all cases,
226 normality and homogeneity of variance was assessed using the Kolmogorov-Smirnov and
227 Levene’s test respectively. Because the number of emerged *D. virgifera* adults in the field

228 experiment did not conform to normality and the variance was unequal for this dataset, the
229 analysis was carried out on rank-transformed data. *D. virgifera* weight gain on maize and
230 teosinte were analyzed on $\log_{10}+2$ transformed data to ensure normality of distribution.
231 Significant effects were subjected to pair-wise comparisons using Holm-Sidak post hoc tests.
232 Association between variables was tested using Pearson Product Moment Correlations and
233 Sum-of-Squares linear regression. Statistical analyses were performed with SigmaStat v3.5 and
234 MiniTab v15.
235

236 **Results**

237 *Recovery of D. virgifera larvae*

238 The tents prevented natural infestation of the two major leaf-pests of corn, *Ostrinia*
239 *nubilalis* and *S. frugiperda*, as no infestation of the control plots by these species was observed.
240 Individual cattail (*Simyra spp.*) and yellow woollybear (*Spilomena virginica*) caterpillars on the
241 other hand were occasionally encountered on the leaves of control plants. Control plants
242 showing clear damage by these herbivores were not used for root-herbivore recovery. From the
243 first infestation of *D. virgifera*, a total of 216 larvae were recovered from the roots. There was
244 no natural infestation by *D. virgifera* in this particular field. The number of recovered root-
245 herbivore larvae from the first infestation was not affected by the presence of *S. frugiperda*
246 (ANOVA: $p=0.536$). Root masses from plots that had suffered from elevated soil moisture
247 carried significantly lower numbers of larvae than the roots from plots with normal water status
248 (ANOVA: $p<0.001$; Holm-Sidak post-hoc test: $p=0.001$; Fig. 1a). From the second infestation, a
249 total of 129 larvae were retrieved. The first infestation larvae had reached the pupal stage by
250 the time the second generation was sampled. It is therefore unlikely that individuals from this
251 group ended up in the collection pans and indeed, no third instar larvae or pupae were
252 recovered. The environmental block factor (high moisture levels early on) did not show a
253 significant effect on this infestation of *D. virgifera* (ANOVA: $p=0.607$). On the other hand, the
254 presence of *S. frugiperda* significantly reduced the number of surviving root herbivore larvae of
255 the second infestation (ANOVA: $p=0.027$; Holm-Sidak post-hoc test: $p=0.0275$; Fig. 1b). In the
256 plots that were not infested with *S. frugiperda*, an average of 1.5 larvae/plant was retrieved,

257 whereas in the presence of leaf-herbivores, larval recovery was reduced by 79% to 0.3
258 larvae/plant.

259

260 *Influence of plant growth stage and AG damage*

261 It was observed that the smaller plants suffered significantly more from *S. frugiperda*
262 feeding damage than the plants that were already tasseling: In mid-season (during the period
263 when the root herbivores were recovered) the small plants (growth stage V8) were largely
264 defoliated with only the midrib of the youngest leaves remaining, while the bigger plants
265 (growth stage VT, tasseling) showed only traces of herbivory and minimal notable loss of
266 biomass. Only later in the season (at the beginning of the adult-emergence period) did the VT
267 plants also suffer from major defoliation. This difference was most probably due to the fact that
268 tasseling plants had tougher leaves (Williams *et al.*, 1998) and no whorl tissue that serves as an
269 important protective structure for *S. frugiperda*. To test whether this difference in defoliation
270 had an effect on *D. virgifera* resistance, we added plant size (big vs. small) as an additional
271 parameter into the model. The nested ANOVA (with plant size as a nested parameter) showed
272 no significant effect of elevated soil moisture (ANOVA: $p=0.555$) or plant size ($p=0.668$), but the
273 effect of *S. frugiperda* was highly significant for the second infestation (ANOVA: $p=0.008$; Fig.
274 1c).

275

276 *Root damage rating*

277 The clear difference in the numbers of larvae recovered from the differentially shoot-
278 infested plants was not reflected in the observed root damage. One explanation for this is that

279 overall, the level of *D. virgifera* infestation was relatively low (Hibbard *et al.* 2010), and damage
280 scores were between 0-1 for most root systems, which corresponds to less than one node of
281 pruning. Damage to the first batch of rated plants (attacked by the first infestation of *D.*
282 *virgifera*) was not affected by *S. frugiperda* feeding (ANOVA: $p=0.815$), but was reduced in
283 plants growing in soil with high early humidity levels (ANOVA: $p=0.022$; Fig. 2a). The second set
284 of plants (sequentially attacked by both infestations of *D. virgifera*) showed the same pattern,
285 with no significant effect of *S. frugiperda* (ANOVA: $p=0.505$) and a negative effect of flooding
286 (ANOVA: $p=0.012$; Fig. 2b).

287

288 *D. virgifera* adult emergence

289 In total, 338 adult *D. virgifera* beetles were collected from the field tents over 6 weeks.
290 The beetles were from the first infestation only, as the larvae of the second infestation did not
291 have enough time to reach the adult stage before the termination of the experiment. The
292 number of adults was affected by the elevated soil moisture factor (ANOVA: $p=0.042$), as well
293 as by *S. frugiperda* feeding ($p<0.001$): Significantly fewer adults emerged from the plots that
294 had experienced flooding, and the same was true for plots in which *S. frugiperda* had fed on the
295 leaves (Figs. 2c and d). When tested separately, the negative effect of *S. frugiperda* feeding was
296 significant for both male (ANOVA: $p<0.001$) and female (ANOVA: $p=0.002$) emergence (data not
297 shown).

298

299 *D. virgifera* weight gain

300 Similarly to the field experiment, larval development of *D. virgifera* was negatively
301 affected by *S. frugiperda* feeding in the laboratory. In both cultivated maize and the wild
302 ancestor teosinte, *D. virgifera* larvae on plants that had previously been infested by *S.*
303 *frugiperda* gained less weight over 5 days compared to larvae on plants that were free of *S.*
304 *frugiperda* (Figs. 3a and 4a). Interestingly, *D. virgifera* larvae that had established on the roots
305 before *S. frugiperda* showed similar weight gain as larvae on uninfested maize plants (Fig. 3a)
306 and were affected only slightly on teosinte (Fig. 4a). Leaf-biomass was reduced significantly
307 (~50%) by *S. frugiperda* feeding on the relatively small maize plants used in the laboratory assay
308 (ANOVA: $p < 0.001$). The teosinte plants also suffered from a significant reduction of leaf fresh
309 weight (ANOVA: $p < 0.001$), although this was less pronounced. Leaf biomass was reduced more
310 for the plants that had been infested first with *S. frugiperda* compared to the ones where *S.*
311 *frugiperda* attacked the plants after *D. virgifera* (Holm-Sidak post-hoc test: $p < 0.05$; Figs. 3b and
312 4b). As it is known that leaf-to-root effects can directly depend on the extent of defoliation
313 (Kaplan *et al.*, 2008), we tested if there was a relationship between leaf-biomass removal and
314 *D. virgifera* weight gain. In accordance with our observations in the field, no significant
315 correlation was found between these two factors, neither in maize ($R^2 = 0.032$; Fig. 3c) nor
316 teosinte ($R^2 = 0.003$; Fig. 4c).

317

318 Discussion

319 To the best of our knowledge, the presented study shows for the first time that the
320 sequence of arrival is an important factor shaping plant-mediated interactions between
321 herbivores. In the field experiment, the number of *D. virgifera* larvae recovered from the roots
322 was not changed by *S. frugiperda* feeding on the leaves if *D. virgifera* established on the plants
323 first (Fig. 1a). However, the root-feeding larvae that arrived after *S. frugiperda* were negatively
324 affected by leaf herbivory (Fig. 1b). The same effect was observed in the laboratory, where
325 larval growth was only impaired when the leaf-feeder had attacked the plant first (Figs. 3a and
326 4a). In nature, root herbivores may therefore escape this negative effect by arriving early on the
327 plant. Interestingly, early studies on AG-BG interactions reported enhanced herbivore growth
328 rates rather than induced resistance (Masters *et al.*, 1993). This has been attributed to an
329 increase in primary metabolite concentrations in the systemic tissues (van Dam and Heil, this
330 issue; Kaplan *et al.*, 2008). While phloem feeding aphids and plant parasitic nematodes may
331 indeed benefit from such changes, our study adds to the growing evidence the chewing
332 herbivores are suffering from induced defenses after primary attack (van Dam and Heil, this
333 issue). We are currently investigating if the increase in resistance reported in this study is
334 indeed due to an increase in defensive metabolite concentrations in the roots, or if changes in
335 primary metabolism are involved as well (see below).

336 The laboratory experiments allow a comparison between cultivated and wild maize
337 plants to herbivory. The general pattern regarding the sequence-specificity of leaf-herbivore
338 induced root resistance was similar for teosinte and maize (Figs. 3 and 4), suggesting that the
339 physiological responses have not been altered during the cultivation process. Yet, some small

340 differences between the two systems were observed. First, teosinte suffered less leaf-herbivory
341 by *S. frugiperda* in terms of biomass loss than cultivated maize (Figs. 3b and 4b). It remains to
342 be determined if the wild plant is naturally more resistant to leaf-herbivory than the cultivar, or
343 if the slightly advanced developmental state of the teosinte plants compared to maize (Figs. 3b
344 and 4b) was responsible for this difference. Second, the effect on root herbivore growth was
345 less pronounced in teosinte than in maize (Figs. 3a and 4a). This may be due to the fact that the
346 plants were less induced by the leaf herbivores. Moreover, the somewhat higher standard
347 deviations indicate higher genetic variability in the field-collected teosinte compared to the
348 genetically uniform background of the cultivar. Future experiments could aim at comparing
349 leaf-herbivore induced root resistance in a variety of wild teosinte populations to get insight
350 into possible evolutionary drivers behind the phenomenon.

351 Interestingly, *D. virgifera* infestation has been shown to increase leaf-resistance against
352 *Spodoptera littoralis* in the laboratory (Erb *et al.*, 2009a) and against lepidopteran herbivores in
353 the field (M. Erb, in press). This phenomenon may partially explain why the removal of leaf-
354 biomass was reduced in the laboratory when *S. frugiperda* had to feed on *D. virgifera* infested
355 maize or teosinte plants (Figs. 3b and 4b). Although root herbivore-induced leaf resistance
356 (RISR) is unlikely to be adaptive for the plant (M. Erb, in press), it may help the root herbivore to
357 protect itself against negative effects of AG herbivores. RISR may have contributed to the
358 reduction of negative shoot-to-root effects in the laboratory, but the field experiment was not
359 confounded by this factor because in all treatments, *S. frugiperda* fed on plants that had been
360 infested in the roots before, regardless of the arrival of the second generation. Yet, for the field
361 experiment, it would theoretically be possible that the feeding by the first infestation changed

362 the physiology of the roots differentially depending on the presence of the leaf-herbivore,
363 which then could have influenced the performance of the second infestation. Alternatively,
364 differences in the behavior of the diapausing and non-diapausing strains may have contributed
365 to the observed results (Prischmann *et al.*, 2008). However, the laboratory experiments
366 demonstrate that leaf-herbivore induced root resistance functions independently of such
367 effects, as only one root herbivore generation was present per plant, and the same *D. virgifera*
368 strain was used for all treatments. Taken together, due to their complementary nature, the
369 field and laboratory experiments conclusively show that the sequence of arrival is important for
370 the outcome of plant-mediated insect-plant-insect interactions.

371 AG attack by *S. frugiperda* profoundly influences the physiology and host suitability of
372 maize roots for root-feeding insects. It is unlikely that the lack of assimilate supply from the
373 leaves is responsible for this phenomenon, as i) both heavily defoliated and less-damaged
374 plants supported lower numbers of *D. virgifera* larvae (Fig. 1c), and ii) there was no correlation
375 between the available leaf-biomass and root herbivore growth (Figs. 3c and 4c). On the
376 contrary, leaf-defoliation by grasshoppers has been shown to increase root assimilate flows in
377 maize (Holland *et al.*, 1996). Another possible explanation for the observed reduction in root
378 herbivore performance could be that leaf-herbivory leads to a short-term reduction of root-
379 growth (Hummel *et al.*, 2009) and a long term-decrease of root-biomass (Bardgett *et al.*, 1998).
380 During the course of the field experiment, however, both larval densities and adult emergence
381 numbers were low (Figs. 1 and 2) and the root systems showed only little damage (Fig. 2),
382 implying that root biomass was not a limiting factor. Equally, ample root-biomass was available

383 in the laboratory assays at the end of the experiment. Therefore, the differences in *D. virgifera*
384 performance likely stemmed from changes in secondary metabolism.

385 It has been proposed that highly resistant maize lines produce the defensive protein MIR1-CP in
386 the roots upon leaf-attack by *S. frugiperda* (Lopez *et al.*, 2007). Plants synthesize a variety of
387 secondary metabolites BG to support leaf-defences (Erb *et al.*, 2009c) that may also negatively
388 affect *D. virgifera*. Further research will have to be conducted to characterize the alterations in
389 root physiology that increase BG resistance. It will be interesting to see if these defences are
390 induced differentially in the roots depending on the sequence of arrival. Another focus should
391 be on possible shoot-root signals mediating the interaction. It has been proposed that
392 phytohormone cross-talk may be responsible for a series of plant-mediated interactions
393 between herbivores: The plant's salicylic acid (SA) response for example down regulates
394 jasmonic acid (JA) dependent defense genes (Spoel *et al.*, 2007), which may explain the
395 interference of whiteflies with induced resistance (Zarate *et al.*, 2007) and bacterial
396 colonization belowground (Yang *et al.*, this issue). However, our hormonal profiles suggest
397 that none of the classical stress-response signals (JA, SA and abscisic acid) change in
398 concentration in the roots upon herbivory by *Spodoptera littoralis* (Erb *et al.*, 2009a). This
399 indicates that hormonal crosstalk is not responsible for the reported interaction, and that a
400 hitherto unknown insect-induced compound mediates the increase in systemic resistance BG,
401 which is not surprising, given the complexity of plant hormonal networks (Erb and Glauser,
402 2010).

403 It has also been suggested that early arriving herbivores may “canalize the plant
404 response”, making it less reactive to subsequent changes (Viswanathan *et al.*, 2007).

405 Conversely, other studies show that a prior stress may “accentuate” the response to a
406 secondary attacker (Erb *et al.*, 2009b, Ton *et al.*, 2007). In our field experiment, canalization is
407 an unlikely scenario, as the late arriving *D. virgifera* larvae would have benefited equally from
408 the fact that the early arriving root-feeders would have blocked the leaf-herbivore induced
409 changes. For the same reason, an accentuated response is an equally unlikely, as all the
410 “second generation” *D. virgifera* larvae arrived on plants that had previously been induced in
411 the roots by the early arrivers. This raises the question about the nature of the sequence
412 dependent factor. We hypothesize that an increase in feeding-deterrent and/or repellent
413 secondary metabolites is responsible for the observed effects. Such compounds would interfere
414 with the host-location and host-acceptance behavior of herbivores that arrive on the plant, but
415 not necessarily with the feeding behavior of larvae that have already colonized and burrowed
416 into the roots. In the laboratory set-up, the fact that the *D. virgifera* larvae did grow less over 5
417 days on plants that had been pre-infested in the leaves may therefore have been the
418 consequence of the fact that they did not accept the roots as hosts and therefore did not
419 readily initiate feeding. *D. virgifera*, as a highly specialized herbivore, has been shown to be
420 very responsive to specific root metabolites (Spencer *et al.*, 2009, Bernklau and Bjostad, 2008),
421 and future experiments will aim at characterizing the behavior and feeding pattern of root
422 herbivores in the presence of leaf-attackers.

423 In conclusion, we demonstrate that the sequence of arrival of different insect herbivore
424 species on a plant can be an important determinant shaping the outcome of plant-mediated
425 interactions between them. Further studies involving other systems will be needed to evaluate
426 if this is a general pattern in plant-insect interactions. Our results suggest that in order to

427 understand the interplay between herbivores sharing a host plant, their sequence of arrival has
428 to be addressed. Experimentally imposed insect-treatments in particular may lead to erroneous
429 interpretations if they do not take into account the natural order of insect-succession during
430 the growing season.

431

432

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442

443 **References**

- 444 Bardgett, R. D., Wardle, D. A. & Yeates, G. W. (1998) Linking above-ground and below-ground
 445 interactions: How plant responses to foliar herbivory influence soil organisms. *Soil Biology &*
 446 *Biochemistry*, **30**, 1867-1878.
- 447 Bernklau, E. J. & Bjostad, L. B. (2008) Identification of feeding stimulants in corn roots for western corn
 448 rootworm (Coleoptera : Chrysomelidae) larvae. *Journal of Economic Entomology*, **101**, 341-351.
- 449 Blosssey, B. & Hunt-Joshi, T. R. (2003) Belowground herbivory by insects: Influence on plants and
 450 aboveground herbivores. *Annual Review of Entomology*, **48**, 521-547.
- 451 Branson, T. F. (1976) The selection of a non-diapausing strain of *Diabrotica virgifera* (Coleoptera:
 452 Chrysomelidae). *Entomologia Experimentalis et Applicata*, **19**, 148-154.
- 453 Branson, T. F., and J. L. Krysan. (1981) Feeding and oviposition behavior and life cycle strategies of
 454 *Diabrotica*: an evolutionary view with implications for pest management. *Environmental*
 455 *Entomology*, **10**, 826-831.
- 456 Dangl, J. L. & Jones, J. D. G. (2001) Plant pathogens and integrated defence responses to infection.
 457 *Nature*, **411**, 826-833.
- 458 Erb, M., Koellner, T.G., Degenhardt, J., Zwahlen, C., Hibbard, B.E.& Turlings, T.C.J (in press). The role of
 459 abscisic acid and water stress in root herbivore-induced leaf-resistance. *New Phytologist*.
- 460 Erb, M. & Glauser, G. (2010). Family business: Multiple members of major phytohormone classes
 461 orchestrate plant stress responses. *Chemistry- A European Journal*. Published Online.
- 462 Erb, M., Flors, V., Karlen, D., de Lange, E., Planchamp, C., D'Alessandro, M., Turlings, T. C. J. & Ton, J.
 463 (2009a). Signal signature of aboveground-induced resistance upon belowground herbivory in
 464 maize. *Plant Journal*, **59**, 292-302.
- 465 Erb, M., Gordon-Weeks, R., Camaño, G., Turlings, T. C. J. & Ton, J. (2009b) Belowground ABA boosts
 466 aboveground production of DIMBOA and primes induction of chlorogenic acid in maize. *Plant*
 467 *Signalling & Behavior*, **4**, 639-642.
- 468 Erb, M., Lenk, C., Degenhardt, J. & Turlings, T. C. J. (2009c) The underestimated role of roots in defence
 469 against leaf attackers. *Trends in Plant Science*, **14**, 653-659.
- 470 Erb, M., Ton, J., Degenhardt, J. & Turlings, T. C. J. (2008) Interactions between Arthropod-Induced
 471 Aboveground and Belowground Defences in Plants. *Plant Physiology*, **146**, 867-874.
- 472 Foster, R. E. & Cherry, R. H. (1987) Survival of Fall Armyworm, *Spodoptera frugiperda*, (Lepidoptera,
 473 Noctuidae) exposed to cold temperatures. *Florida Entomologist*, **70**, 419-422.
- 474 Heil, M. & Ton, J. (2008) Long-distance signalling in plant defence. *Trend in plant science*.
- 475 Heil, M. & van Dam, N (this issue). Multitrophic interactions belowground and aboveground: en route to
 476 the next level.
- 477 Hibbard, B. E., Higdon, M. L., Duran, D. P., Schweikert, Y. M. & Ellersieck, M. R. (2004) Role of egg density
 478 on establishment and plant-to-plant movement by western corn rootworm larvae (Coleoptera :
 479 Chrysomelidae). *Journal of Economic Entomology*, **97**, 871-882.
- 480 Hibbard, B. E., Meihls, L. N., Ellersieck, M. R. & Onstad, D. W. (2010) Density-Dependent and Density-
 481 Independent Mortality of the Western Corn Rootworm: Impact on Dose Calculations of
 482 Rootworm-Resistant Bt Corn. *Journal of Economic Entomology*, **103**, 77-84.
- 483 Hibbard, B. E., Schwikert, Y. M., Higdon, M. L. & Ellersieck, M. R.(2008). Maize phenology affects
 484 establishment, damage and development of the western corn rootworm (Coleoptera:
 485 Chrysomelidae). *Environmental Entomology*, **37**, 1558-1564.
- 486 Holland, J. N., Cheng, W. X. & Crossley, D. A. (1996) Herbivore-induced changes in plant carbon
 487 allocation: Assessment of below-ground C fluxes using carbon-14. *Oecologia*, **107**, 87-94.
- 488 Howe, G. A. & Jander, G. (2008) Plant immunity to insect herbivores. *Annual Review of Plant Biology*, **59**,
 489 41-66.

490 Hummel, G. M., Schurr, U., Baldwin, I. T. & Walter, A. (2009) Herbivore-induced jasmonic acid bursts in
491 leaves of *Nicotiana attenuata* mediate short-term reductions in root growth. *Plant Cell and*
492 *Environment*, **32**, 134-143.

493 Kaplan, I. & Denno, R. F. (2007) Interspecific interactions in phytophagous insects revisited: a
494 quantitative assessment of competition theory. *Ecology Letters*, **10**, 977-994.

495 Kaplan, I., Halitschke, R., Kessler, A., Rehill, B. J., Sardanelli, S. & Denno, R. F. (2008) Physiological
496 integration of roots and shoots in plant defence strategies links above- and belowground
497 herbivory. *Ecology Letters*, **11**, 841-851.

498 Lopez, L., Camas, A., Shivaji, R., Ankala, A., Williams, P. & Luthe, D. (2007) Mir1-CP, a novel defence
499 cysteine protease accumulates in maize vascular tissues in response to herbivory. *Planta*, **226**,
500 517-527.

501 Masters, G. J., Brown, V. K. & Gange, A. C. (1993) Plant mediated interactions between aboveground
502 and belowground insect herbivores. *Oikos*, **66**, 148-151.

503 Masters, G. J. (1995) The effect of herbivore density on host-plant mediated interactions between 2
504 insects. *Ecological Research*, **10**, 125-133.

505 Moran, N. A. & Whitham, T. G. (1990) Interspecific Competition between Root-Feeding and Leaf-Galling
506 Aphids Mediated by Host-Plant Resistance. *Ecology*, **71**, 1050-1058.

507 O'Day, M. (1998) *Corn insect pests: A diagnostic guide*. University of Missouri-Columbia, Missouri.

508 Oleson, J. D., Park, Y. L., Nowatzki, T. M. & Tollefson, J. J. (2005) Node-injury scale to evaluate root injury
509 by corn rootworms (Coleoptera : Chrysomelidae). *Journal of Economic Entomology*, **98**, 1-8.

510 Orians, C. (2005) Herbivores, vascular pathways, and systemic induction: Facts and artifacts. *Journal of*
511 *Chemical Ecology*, **31**, 2231-2242.

512 Poelman, E. H., Broekgaarden, C., Van Loon, J. J. A. & Dicke, M. (2008) Early season herbivore
513 differentially affects plant defence responses to subsequently colonizing herbivores and their
514 abundance in the field. *Molecular Ecology*, **17**, 3352-3365.

515 Poelman, E. H., van Loon, J. J. A., van Dam N. M., Vet L. E. M., Dicke, M. (2010) Herbivore-induced plant
516 responses in *Brassica oleracea* prevail over effects of constitutive resistance and result in
517 enhanced herbivore attack. *Ecological Entomology*, **35**, 240-247.

518 Prischmann, D. A., Dashiell, K. E. & Hibbard, B. E. (2008) Assessing larval rootworm behavior after
519 contacting maize roots; impact of germplasm, rootworm species and diapause status. *Journal of*
520 *Applied Entomology*, **133**, 21-32.

521 Rasmann, S., Erwin, A. C., Halitschke, R. & Agrawal, A. A. (this issue). Direct and indirect root defences of
522 milkweed (*Asclepias syriaca*): trophic cascades, trade-offs and novel methods for studying
523 subterranean herbivory.

524 Rasmann, S. & Turlings, T. C. J. (2007) Simultaneous feeding by aboveground and belowground
525 herbivores attenuates plant-mediated attraction of their respective natural enemies. *Ecology*
526 *Letters*, **10**, 926-936.

527 Schwachtje, J. & Baldwin, I. T. (2008) Why does herbivore attack reconfigure primary metabolism? *Plant*
528 *Physiology*, **146**, 845-851.

529 Short, D. E. & Luedtke, R. J. (1970) Larval migration of western corn rootworm. *Journal of Economic*
530 *Entomology*, **63**, 325-&.

531 Soler, R., Bezemer, T. M., Cortesero, A. M., Van der Putten, W. H., Vet, L. E. M. & Harvey, J. A. (2007)
532 Impact of foliar herbivory on the development of a root-feeding insect and its parasitoid.
533 *Oecologia*, **152**, 257-264.

534 Soler, R., Bezemer, T. M., Van der Putten, W. H., Vet, L. E. M. & Harvey, J. A. (2005) Root herbivore
535 effects on above-ground herbivore, parasitoid and hyperparasitoid performance via changes in
536 plant quality. *Journal of Animal Ecology*, **74**, 1121-1130.

537 Spencer, J. L., Hibbard, B. E., Moeser, J. & Onstad, D. W. (2009) Behaviour and ecology of the western
538 corn rootworm (*Diabrotica virgifera virgifera* LeConte). *Agricultural and Forest Entomology*, **11**,
539 9-27.

540 Spoel, S.H., Johnson, J.S. & Dong, X. (2007). Regulation of tradeoffs between plant defenses against
541 pathogens with different lifestyles. *PNAS*, **104**, 18842-18847.

542 Steinger, T. & Müller-Schärer, H. (1992) Physiological and growth-responses of *Centaurea maculosa*
543 (*Asteraceae*) to root herbivory under varying levels of interspecific plant competition and soil-
544 nitrogen availability. *Oecologia*, **91**, 141-149.

545 Sticher, L., MauchMani, B. & Metraux, J. P. (1997) Systemic acquired resistance. *Annual Review of*
546 *Phytopathology*, **35**, 235-270.

547 Suttle, P. J., Musick, G. J. & Fairchil.MI (1967) Study of larval migration of Western Corn Rootworm.
548 *Journal of Economic Entomology*, **60**, 1226-&.

549 Ton, J., D'Alessandro, M., Jourdie, V., Jakab, G., Karlen, D., Held, M., Mauch-Mani, B. & Turlings, T. C. J.
550 (2007) Priming by airborne signals boosts direct and indirect resistance in maize. *Plant Journal*,
551 **49**, 16-26.

552 van Dam, N. M., Raaijmakers, C. E. & van der Putten, W. H. (2005) Root herbivory reduces growth and
553 survival of the shoot feeding specialist *Pieris rapae* on *Brassica nigra*. *Entomologia*
554 *Experimentalis Et Applicata*, **115**, 161-170.

555 van Loon, L. C., Bakker, P. & Pieterse, C. M. J. (1998) Systemic resistance induced by rhizosphere
556 bacteria. *Annual Review of Phytopathology*, **36**, 453-483.

557 van Zandt, P. A., Agrawal, A. A. (2004) Community-wide impacts of herbivore-induced plant responses in
558 milkweed (*Asclepias syriaca*). *Ecology*, **85**, 2616-2629.

559 Viswanathan, D. V., Lifchits, O. A. & Thaler, J. S. (2007) Consequences of sequential attack for resistance
560 to herbivores when plants have specific induced responses. *Oikos*, **116**, 1389-1399.

561 Viswanathan, D. V., Narwani, A. J. T. & Thaler, J. S. (2005) Specificity in induced plant responses shapes
562 patterns of herbivore occurrence on *Solanum dulcamara*. *Ecology*, **86**, 886-896.

563 Voelckel, C. & Baldwin, I. T. (2004) Herbivore-induced plant vaccination. Part II. Array-studies reveal the
564 transience of herbivore-specific transcriptional imprints and a distinct imprint from stress
565 combinations. *Plant Journal*, **38**, 650-663.

566 Williams, W. P., Davis, F. M., Buckley, P. M., Hedin, P. A., Baker, G. T. & Luthe, D. S. (1998) Factors
567 associated with resistance to Fall Armyworm (*Lepidoptera* : *Noctuidae*) and southwestern corn
568 borer (*Lepidoptera*: *Crambidae*) in corn at different vegetative stages. *Journal of Economic*
569 *Entomology*, **91**, 1471-1480.

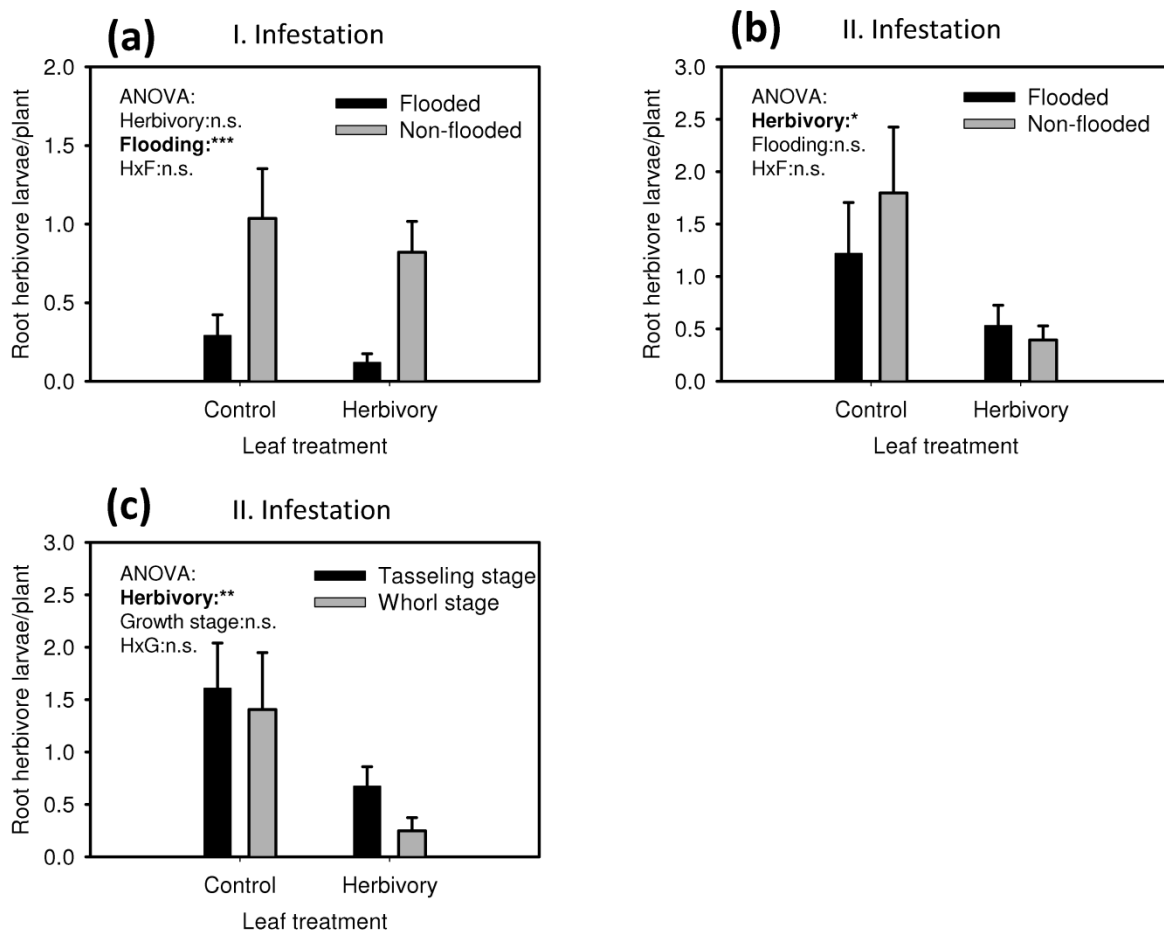
570 Yang, J. W., Yi, H.-S., Lee, B., Lee, S. & Ghim, S.-Y. (this issue). Whitefly infestation elicits defense
571 responses against bacterial pathogens on the leaf and root and belowground dynamic change of
572 microflora in pepper.

573 Zarate, S. I., Kempema, L. A. & Walling, L. L. (2007). Silverleaf whitefly induces salicylic acid defenses and
574 suppressed effectual jasmonic acid defenses. *Plant Physiology*, **143**, 866-875.

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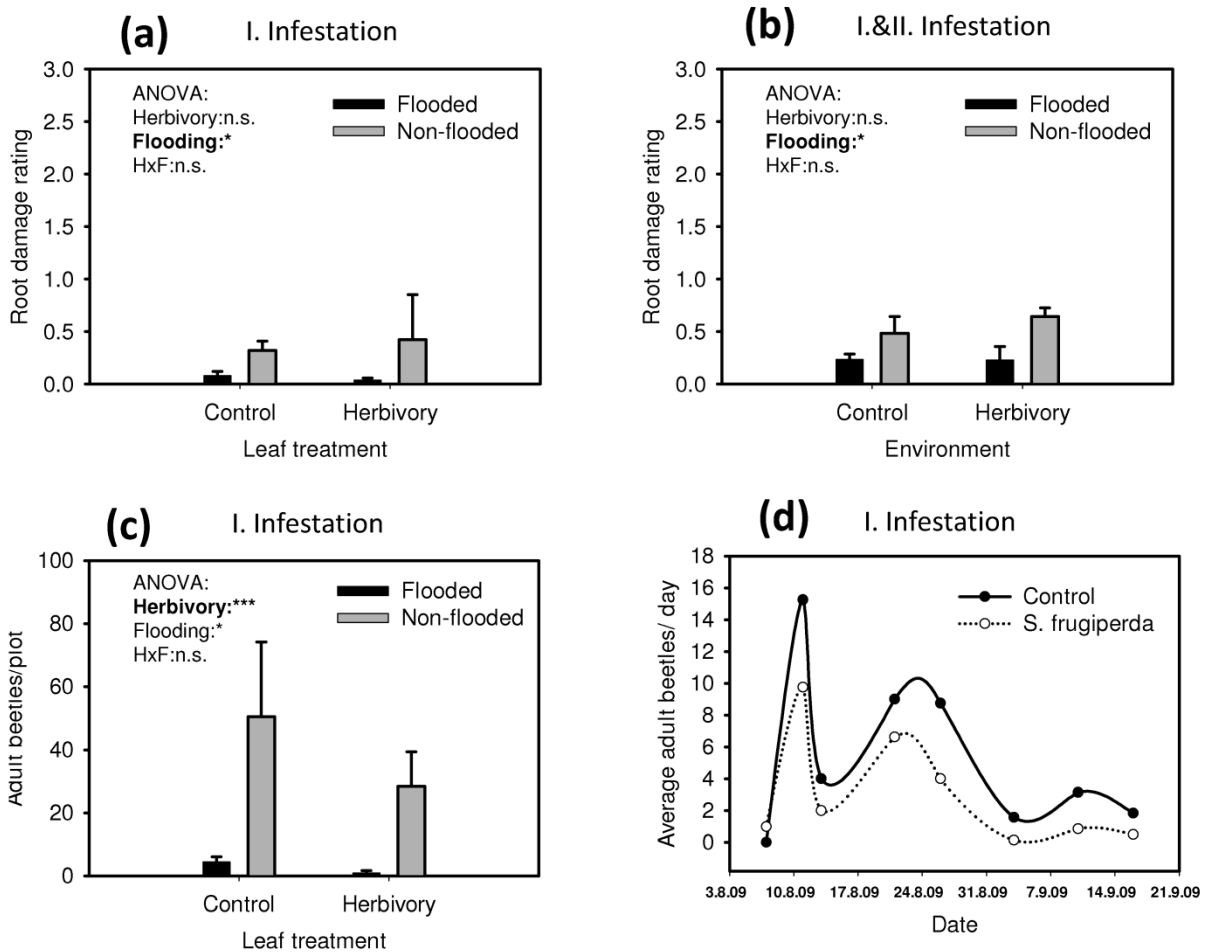
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Fig. 1



577
 578 **Fig. 1:** Influence of leaf herbivory by *S. frugiperda* on recovery rates of root feeding *D. virgifera* larvae. **(a):** Average
 579 number (+SE) of first infestation *D. virgifera* larvae/ plant are shown. *D. virgifera* larvae established on the plants
 580 before onset of *S. frugiperda* herbivory. **(b):** Average number (+SE) of second infestation *D. virgifera* larvae/ plant.
 581 *D. virgifera* larvae established on the plants after onset of *S. frugiperda* herbivory. Numbers recovered from
 582 control plants (left) and *S. frugiperda* infested plants (right) are shown. Plots that suffered from flooding (black
 583 bars) are separated from undisturbed plots (grey bars). Results from two-way ANOVAs are included. Effects of
 584 Herbivory (*S. frugiperda* and control), flooding (flooded and non-flooded), and their interaction (HxF) are depicted.
 585 **(c):** Average number (+SE) of second infestation *D. virgifera* larvae/ plant. Numbers recovered from control plants
 586 (left) and *S. frugiperda* infested plants (right) are shown. Tasseling maize plants (black bars) are separated from
 587 plants in the late whorl stage (grey bars). Effects of Herbivory (*S. frugiperda* and control), growth stage (whorl and
 588 tasseling stage), and their interaction (HxG) are depicted. Stars denote significant factor effects (* $p < 0.05$;
 589 ** $p < 0.01$; *** $p < 0.001$). N=8.
 590

Fig. 2

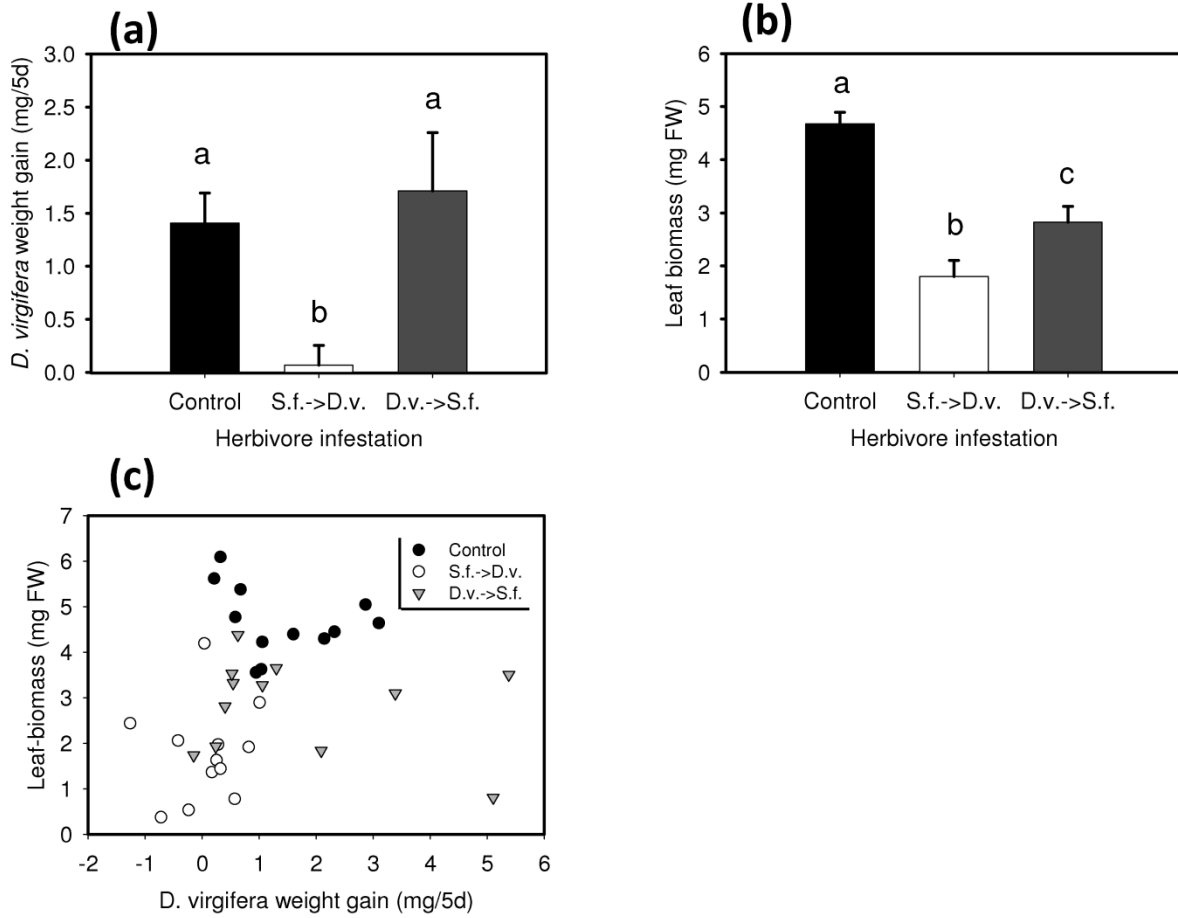


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Fig. 2: Effect of leaf herbivory by *S. frugiperda* on *D. virgifera* root damage and adult emergence. **(a):** Average root rating (+SE) of plants after infestation with the first infestation of *D. virgifera* larvae. **(b):** Average root rating (+SE) of plants after infestation with the first and the second infestation of *D. virgifera* larvae. **(c):** Average number (+SE) of emerging *D. virgifera* adults per plot. Numbers recovered from control plants (left) and *S. frugiperda* infested plants (right) are shown. Plots that suffered from flooding (black bars) are separated from undisturbed plots (grey bars). Results from two-way ANOVAs are included. Effects of Herbivory (*S. frugiperda* and control), flooding (flooded and non-flooded), and their interaction (HxF) are depicted. Stars denote significant factor effects (* $p < 0.05$; ** $p < 0.01$; *** $p < 0.001$). **(d):** Time course of emerging adult beetles over the collection period. Average adult beetles per day from control plants (closed circles) and *S. frugiperda* infested plants (open circles) are shown. N=8.

Fig. 3

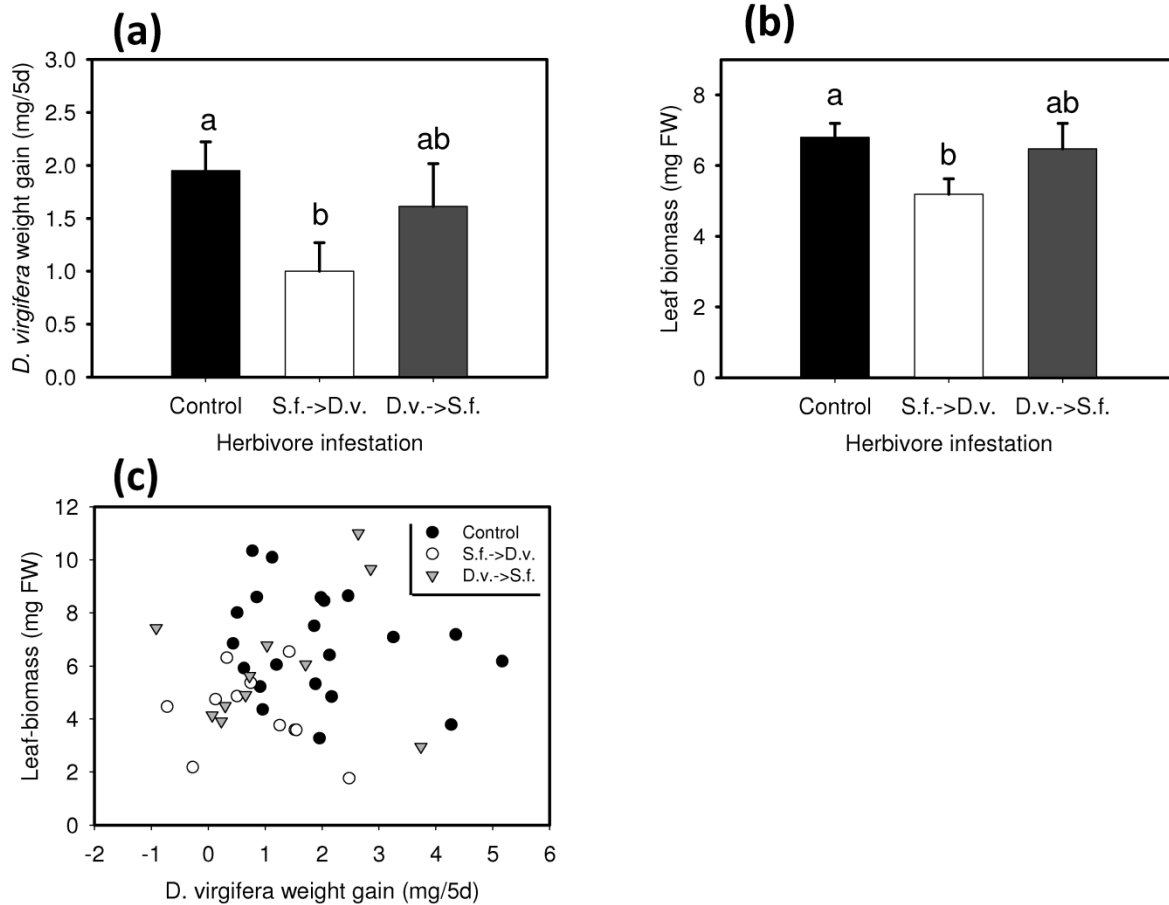
Cultivated maize



603
 604 **Fig. 3:** Influence of leaf herbivory by *S. frugiperda* on *D. virgifera* growth on cultivated maize. **(a):** Average weight
 605 gain (+SE) of *D. virgifera* larvae feeding on leaf-herbivore free plants (control, black bars), previously *S. frugiperda*
 606 infested plants (before onset of root herbivory, S.f.->D.v., open bars) and late *S. frugiperda* infested plants (after
 607 onset of root herbivory, D.v.->S.f., grey bars) are shown. **(b):** Average leaf-biomass of *D. virgifera* and *S. frugiperda*
 608 infested plants. Different letters indicate significant differences between treatments (p<0.05). **(c):** Correlation
 609 between leaf-biomass and *D. virgifera* weight gain on leaf herbivore free plants (filled circles), previously *S.*
 610 *frugiperda* infested plants (empty circles.) and simultaneously *S. frugiperda* infested plants (gray triangles). N=12-
 611 15.
 612

Fig. 4

Wild teosinte



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614

615 **Fig. 4:** Influence of leaf herbivory by *S. frugiperda* on *D. virgifera* growth on teosinte. **(a):** Average weight gain (+SE)
616 of *D. virgifera* larvae feeding on leaf-herbivore free plants (control, black bars), previously *S. frugiperda* infested
617 plants (before onset of root herbivory, S.f.->D.v., open bars) and late *S. frugiperda* infested plants (after onset of
618 root herbivory, D.v.->S.f., grey bars) are shown. **(b):** Average leaf-biomass of *D. virgifera* and *S. frugiperda* infested
619 plants. Different letters indicate significant differences between treatments ($p < 0.05$). **(c):** Correlation between leaf-
620 biomass and *D. virgifera* weight gain on leaf herbivore free plants (filled circles), previously *S. frugiperda* infested
621 plants (empty circles.) and simultaneously *S. frugiperda* infested plants (gray triangles). N=12.
622

Fig. 1

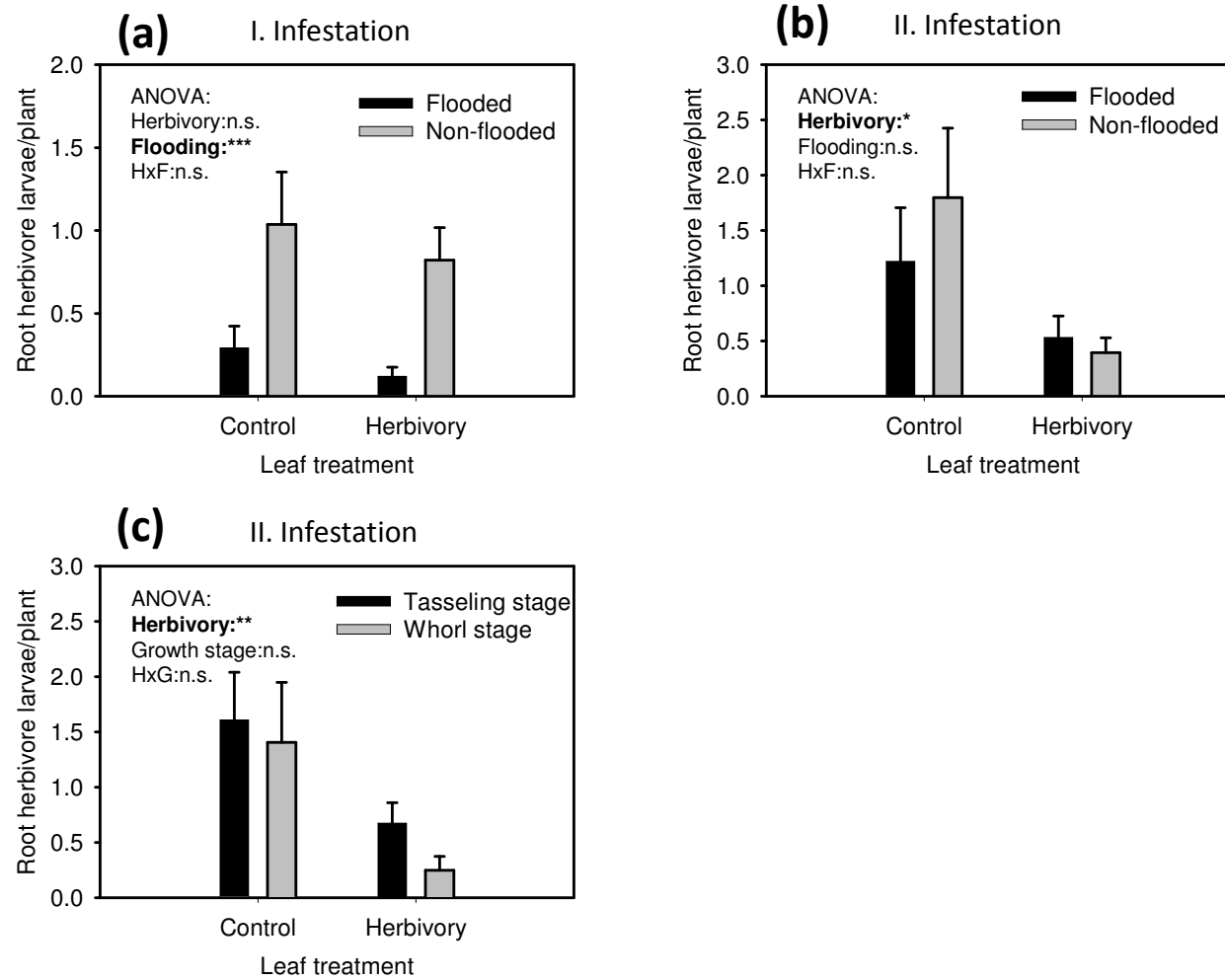


Fig. 2

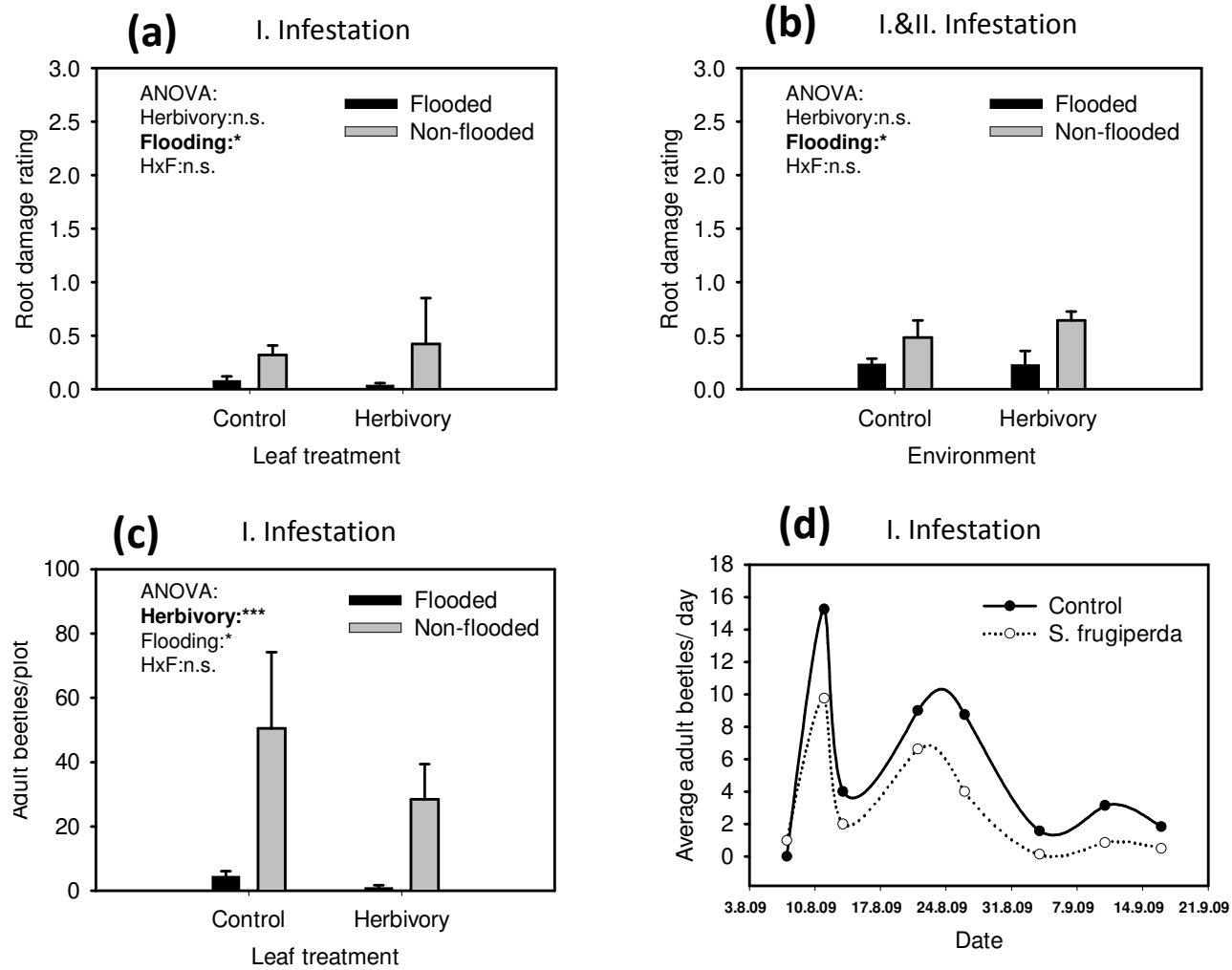


Fig. 3

Cultivated maize

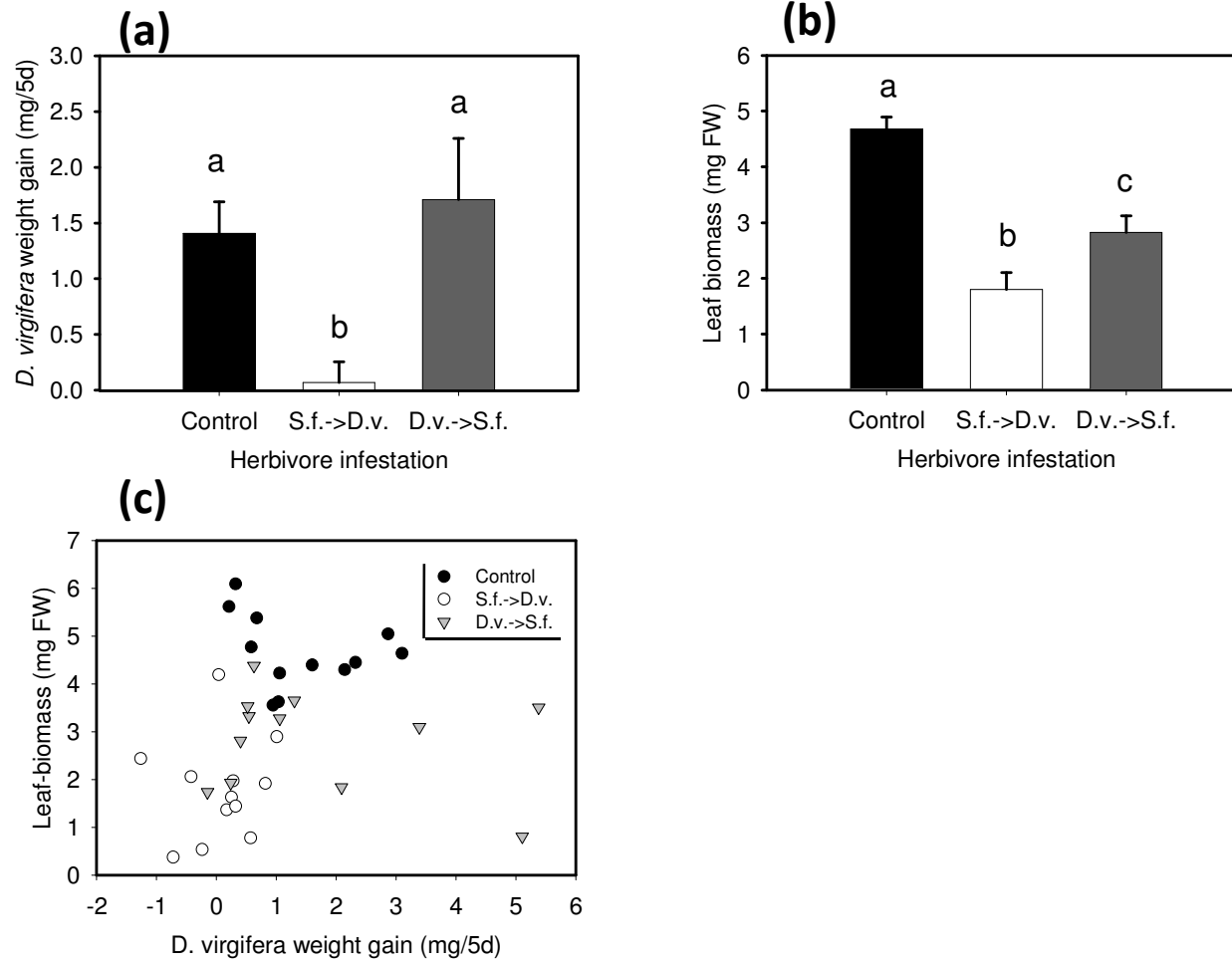


Fig. 4

Wild teosinte

