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Differing short-term impacts of agricultural tarping on soil-dwelling and surface-active arthropods

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1 **Title: Differing short-term impacts of agricultural tarping on soil-dwelling and surface-**
2 **active arthropods**

3

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18 **Keywords**

19 Agricultural management, biodiversity patterns, community recovery, soil arthropod, plastics

20

21 **Abstract**

22 Agricultural tarping, the practice of placing impermeable plastic tarps over crop beds before
23 planting to suppress weeds, is rising in popularity. However, the use of tarps has uncertain
24 effects on soil arthropod communities. We studied the impact of silage (black plastic) tarps and
25 clear plastic tarps on surface-active and soil-dwelling arthropods by tracking immediate impacts
26 and arthropod recovery for 5 weeks after tarps were removed. We also assessed how well
27 environmental and experimental variables explained arthropod diversity and composition.
28 During tarp application, we found that both silage and clear plastic tarps had significant negative
29 impacts on surface-active arthropod diversity, while only clear plastic tarps impacted soil-
30 dwelling arthropods. Surface-active arthropod diversity recovered by 1-3 weeks after tarping, but
31 at 5 weeks after tarping soil-dwelling arthropod diversity was significantly lower in silage tarp
32 and clear plastic plots than control plots. Tarps also led to compositional changes in the
33 arthropod communities, though these changes were only significant during tarp cover. The
34 variables that best explained arthropod diversity and community composition were treatment
35 (i.e., silage tarp, clear plastic tarp, or control) during tarping and farm site after tarps were
36 removed. Other variables, such as soil moisture and weed coverage, were not strong model
37 predictors. These results imply that tarps may have temporary impacts on surface-active
38 arthropods but potentially longer-lasting impacts on soil-dwelling arthropods. Continuing to
39 monitor impacts on tarps on soil arthropods will better inform the sustainability of this practice.

40 **1. Introduction**

41 Tarping is an agricultural practice that has grown in popularity in the last decade, especially in
42 the northeastern USA, sparking management guides (Lounsbury et al., 2022) and regional
43 research (Lounsbury et al., 2018; Birthisel et al., 2019; Rylander et al., 2020). Tarps have been
44 used globally for decades to kill pathogens and pests (Stapleton and DeVay, 1986; Al-Asa'd and
45 Abu-Gharbieh, 1990; Coelho et al., 1999) and, more recently, have been implemented on small-
46 scale farms as a low-input method to suppress weeds or terminate cover crops while also
47 reducing tillage needs (Rylander et al., 2020). Growers place tarps, often silage (black plastic)
48 tarps or clear plastic tarps, over the soil for several weeks before planting crops. Silage tarps kill
49 weeds via occultation (shading), whereas clear plastic tarps kill weeds via solarization (extreme
50 heating) (Rubin and Benjamin, 1984; Johnson and Fennimore, 2005). Both silage tarps and clear
51 plastic tarps create a barrier over the soil and can increase soil temperatures by around 6°C or
52 15°C (respectively) (Birthisel and Gallandt, 2019), which may affect various aspects of the soil
53 ecosystem.

54 Relatively little is known about the impacts of tarping on soil arthropods, a diverse group
55 in agriculture including spiders, beetles, ants, mites, and collembolans. In general, the use of
56 plastic films in agriculture has been identified as a threat to soil biodiversity because they seal
57 the soil surface and can alter physical and chemical soil properties (Birthisel et al., 2019; Tibbett
58 et al., 2020). However, literature on tarps' effects is limited: existing studies mostly focus on the
59 effects of clear tarps (despite heavy use of silage tarps) and analyze only certain arthropod
60 groups (Seman-Varner, 2005; Gill and McSorley, 2010). In these studies (which both took place
61 in the hot and humid climate of the state of Florida, USA), the use of clear tarps reduced
62 collembolan and mite populations (Seman-Varner, 2005), but did not affect other groups, such as

63 spiders, ants, grasshoppers, crickets, and Elateridae beetles (Gill and McSorley, 2010).
64 Investigating these dynamics in a different climate, comprehensively analyzing the entire
65 arthropod community, and testing the impacts of both silage tarps and clear plastic tarps are
66 therefore needed to expand knowledge around tarps. As well, arthropods' roles in providing key
67 ecosystem services for agriculture, such as controlling pests by serving as natural enemies
68 (Kromp, 1999; Lang, 2003; Schmidt et al., 2003; van Lenteren et al., 2018) and improving soil
69 health by contributing to decomposition and soil structure (Lavelle et al., 2006; Brussaard et al.,
70 2007; Briones, 2018), further emphasize the importance of understanding tarps' impacts on this
71 community.

72 To produce results that can best inform conservation, research should be designed to
73 capture complexity within the soil arthropod community (Magurran, 2021), for example taking
74 into account that the effects of tarping can differ among taxonomic or ecological groups due to
75 trait differences and environmental factors (Franken et al., 2018; Yekwayo et al., 2018).

76 Literature on the impacts of plastic mulch (thin plastic sheeting used for weed suppression
77 during, rather than before, crop growth) shows that the direction of impacts varies among
78 arthropod groups (Tuovinen et al., 2006; Addison et al., 2013; Schirmel et al., 2018). This
79 suggests that the effects of tarps could also vary, perhaps relating to arthropods' tolerances to
80 heat or preferences for light versus dark environments (Dindal, 1990; Briones et al., 2009;
81 Bokhorst et al., 2012). The impacts of tarps may also differ between surface-active and soil-
82 dwelling arthropods, which vary in their sizes, diets, mobility, and exposure to disturbances
83 (Dindal, 1990). Analyzing both groups can give a more comprehensive understanding of tarps'
84 effects and uncover potentially unique responses of ground- and soil-dwelling communities
85 (Briones et al., 2009; Liu et al., 2018).

86 It is also important to consider that tarping may have long-term and indirect effects – for
87 example, Birthisel et al. (2019) found that the impact of tarps on soil microorganisms intensified
88 in the weeks after tarp removal. Such lingering effects could relate to tarps’ impacts on the
89 ecosystem, such as on soil temperature, soil moisture, and weed coverage (Birthisel et al., 2019),
90 which change habitat suitability to soil biological communities (Altieri et al., 1985; Schirmel et
91 al., 2018). Arthropod diversity and composition could also vary due to elements of experimental
92 design, including sampling at different sites or times (Campbell et al., 2011; Kirse et al., 2021).
93 Identifying predictors of arthropod diversity and composition during and after tarping will help
94 explain tarps’ impacts and uncover sources of complexity within our system.

95 In this study, we tested the impact of agricultural tarping on surface-active and soil-
96 dwelling arthropod communities. We specifically asked: 1) how do tarps impact the diversity and
97 community composition of soil arthropods?, 2) how do soil arthropod communities respond after
98 tarps are removed?, and 3) how do the experimental and environmental factors in our system
99 explain variability of the soil arthropod community? We hypothesized that both tarp types would
100 decrease soil arthropod diversity, with more negative effects under clear plastic tarps due to the
101 higher temperatures found there. We also hypothesized that tarps’ effects would differ depending
102 on taxonomic groups, creating unique arthropod composition under tarps. Finally, we
103 hypothesized that treatment would best predict soil arthropod diversity during tarp application,
104 but that environmental factors, like soil moisture and weed coverage, would become more
105 predictive after tarps were removed. Monitoring the effects of tarps on soil arthropods is an
106 important step to assessing the sustainability of this practice.

107

108 **2. Methods**

109 2.1. Site descriptions

110 We conducted this experiment in western Chittenden County in Vermont, which is situated in the
111 northeastern United States and has a temperate climate. Our study took place at three farms:
112 Intervale Community Farm (ICF; 44.49820 N, 73.20567 W, 30 m above sea level), Diggers'
113 Mirth Farm (Diggers'; 44.49888 N, 73.20991 W, 30 m above sea level), and Catamount Farm
114 (Catamount; 44.43237 N, 73.20083 W, 70 m above sea level). ICF and Diggers' are relatively
115 close to one another (separated by 300 m), while Catamount is 8 km south of ICF and Diggers'.
116 All three farms are located adjacent to semi-natural areas: ICF and Diggers' are situated near a
117 network of recreational forested areas within the Intervale floodplains west of the Winooski
118 River, and Catamount is located within a residential area but is surrounded by large strips of
119 forest and grassland/shrubland. All three farms mainly grow annual vegetables (Catamount also
120 grows perennial fruits) and have used tarps for weed suppression.

121 The three farms had key environmental and management differences. To understand
122 baseline soil differences, we collected soil using an auger (2 cm diameter, 15 cm depth) and
123 composite sampling (18 soil samples from each farm, taken from the center of each 4.5 m by 1.5
124 m plot; see Section 2.2). Soils were analyzed by the UVM Agricultural and Environmental
125 Testing Lab (<https://www.uvm.edu/extension/agricultural-and-environmental-testing-lab>) and
126 tested for soil texture, pH, available phosphorus, available nitrate, soil organic carbon (SOC),
127 and effective cation exchange capacity (Table 1). One key difference among sites was that
128 Catamount's soils were sand, Diggers' soils were loam, and ICF's soils were sandy loam. Each
129 farm also had a different composition of major weeds, with Catamount dominated by *Portulaca*
130 *oleracea* L. (common purslane) and *Digitaria* grasses (crabgrass), ICF containing high cover of
131 *Chenopodium album* L. and *Chenopodium glaucum* L. (white and oak-leaved goosefoot), and

132 Diggers' containing mostly *Amaranthus retroflexus* L. (redroot pigweed). These ecological
133 differences allow us to understand how tarps function in contrasting environments. We also used
134 irrigation and fertilizer schemes typical to each farm (rather than controlling management among
135 farms) because these practices are honed to each farm's environment, reflect the operational
136 capacities of each farm, and represent realistic scenarios. Catamount used drip irrigation due to
137 its sandy, well-drained soil, while ICF and Diggers' used a combination of sprinklers and hand
138 watering, and each farm received different water amounts depending on the appropriate levels
139 for their soil type.

140 Weather conditions during our experiment were hotter and drier than historical means for
141 the month of June (when tarps were on the field), with a mean precipitation of 0.2 cm per day
142 and a mean temperature of 22°C. In July, after tarps were removed, weather conditions were
143 wetter and cooler than historical means, with a mean precipitation of 0.5 cm per day and a mean
144 temperature of 21°C (NOAA, 2021).

145

146 Table 1: Baseline soil characteristics for the three farm sites. Soil series were determined from
 147 the Web Soil Survey (NRCS USDA, 2022).

	Catamount	Diggers'	ICF
Soil texture	Sand (87% sand, 9% silt, 4% clay)	Loam (44% sand, 48.5% silt, 7.5% clay)	Sandy loam (70.5% sand, 27% silt, 2.5% clay)
Soil series (with Great Group)	Mix of Scarboro (Humaquept), Adams (Haplothod), and Windsor (Udipsamments)	Winooski (Dystrudept)	Hadley (Udifluvent)
World Reference Base (WRB) soil classification	Fluvisol	Cambisol	Cambisol
pH	6.8	7	7
Available phosphorus (ppm)	19.4	104.9	29.5
Available nitrate (mg N / Kg)	17.6	17.2	21.6
Soil organic carbon (%)	2.3	3.1	1.5
Effective cation exchange capacity (meq/100g)	6.5	12.1	7.5

148

149 **2.2. Treatments and experimental design**

150 We tested two types of plastic tarps, silage tarps and clear plastic tarps, along with an uncovered
151 hoed control. We chose to study silage tarps and clear plastic tarps because they are commonly
152 used in New England (Birthisel et al., 2019; Lounsbury et al., 2022) and have key material and
153 functional differences (e.g., clear plastic tarps create hotter soil conditions than silage tarps and
154 are translucent, while silage tarps are opaque). In control plots, we suppressed weeds by hoeing
155 the plots weekly for the duration that the tarps were on the fields. This method is a common
156 organic practice for weed control and was identified as a likely alternative practice to tarping.
157 We chose this control rather than an unmanaged control (e.g., not applying disturbance) because
158 it is not realistic that a farmer would grow crops without any weed suppression practice.
159 Therefore, results comparing soil arthropod communities in tarped areas versus undisturbed areas
160 would not yield useful information to farmers weighing certain management practices.

161 The three treatments were replicated six times on each farm, with the experimental units
162 organized as a completely randomized design (18 plots on each farm; 54 plots total across the
163 three farms). Plots were 4.5 m by 1.5 m (1.5 m is the width of a crop bed), with a buffer space of
164 0.5 m between plots. The study site at each farm consisted of 2-3 adjacent crop rows.

165 Our silage tarps were 0.13 mm polyethylene plastic with a black up-facing side and white
166 down-facing side (Klerks Hyplast Inc., Chester, South Carolina, USA). Our clear plastic tarps
167 were 0.15 mm polyethylene plastic (Poly-Ag Corp, San Diego, California, USA) and were
168 donated from a local farm where they had been used on a hoop house. Repurposing clear plastic
169 from hoop houses represents realistic farm practices (Birthisel et al., 2019).

170 Before treatments were applied, the study site at each farm was tilled and prepared with
171 fertilizer. Additionally, we irrigated the study sites 1-2 days before applying the tarps to

172 stimulate weed germination and increase effectiveness of tarps for killing weeds (Lounsbury et
173 al., 2022). We installed the tarps in late May (late spring) and secured the tarps by burying
174 roughly 15 cm of the tarp edges under around 3 cm of soil. These buried areas were not
175 considered part of the treatment plot. The tarps were installed for 25 days (Table A1).

176 After tarp removal, we did not remove weeds in any plots (tarped or control) for the
177 remainder of the experiment. This reflects the reality of many small organic farms, which do not
178 have the time, labor, or expenses for extensive hand weeding (Fennimore, 2014). We
179 additionally direct seeded two rows of lettuce (*Lactuca sativa*; Encore Lettuce Mix, Product ID:
180 2366G from Johnny’s Selected Seeds; Winslow, ME, USA) in each plot using a Jang seeder
181 (Jang Automation Co., LTD, South Korea) a week after tarp removal. This mimics the use of
182 tarps to prepare beds for crops. The lettuce was not harvested until after the experiment ended
183 (more information on crop yields is available in Kinnebrew et al., 2022a).

184

185 **2.3. Arthropod sampling and identification**

186 We sampled soil arthropods 5 times throughout the field season (Table A1). First, we sampled
187 before tarps were applied to capture baseline diversity patterns in our field areas (mid-May). We
188 then sampled 3 weeks into tarp placement and 1, 3, and 5 weeks after tarp removal (mid-June to
189 mid-July; 5 total sampling periods). We chose sampling dates where weather was clear to
190 capture arthropods when they are most active and to avoid sample losses from rain.

191 We sampled soil arthropods using two methods: pitfall traps and the Berlese Funnel
192 method. We term arthropods caught by pitfall traps “surface-active arthropods” and arthropods
193 captured with the Berlese funnel method “soil-dwelling arthropods.” Pitfall traps consisted of
194 plastic collection cups (95 mm diameter lid, 120 mm deep) placed in the soil with their lids level

195 to the soil surface (Southwood and Henderson, 2009). This method captures active litter- and
196 surface-dwelling arthropods that fall and become trapped in the cups as they move across the soil
197 surface.

198 We installed one pitfall trap in the center of each plot (at least 0.75 m from the tarp edge
199 and 2 m from the nearest adjacent pitfall trap). Each pitfall trap consisted of two stacked cups.
200 The upper cup was used to collect arthropods, while the purpose of the lower cup was to hold the
201 soil in place and avoid repeatedly digging new holes (and disturbing the soil) at each sampling
202 period. When the pitfall traps were not in use (between sampling periods), we removed the upper
203 cups and covered the lower cups with lids and a thin layer of soil. During collection times, we
204 used a killing agent of 50% propylene glycol and 50% water, and left pitfall traps out for 3 days
205 (we collected all aggregated arthropods on the third day). We chose 3 day intervals for the pitfall
206 traps due to high catch rate and because high frequency of rain storms (which are destructive to
207 samples) made it logistically difficult to plan longer trapping intervals. The 3 days were
208 consecutive for almost all samplings. An exception was the last sampling (5 weeks after tarp
209 removal), when we collected arthropods after 2 days, waited 1 day while a heavy storm passed,
210 and then set out new traps for 1 day. We pooled the data from these 3 days. To sample
211 arthropods under the tarps, we cut a 0.5 m hole in the tarp, set up the pitfall trap (and collected
212 soil for the Berlese funnels), and then sealed the tarp using either black duct tape for the silage
213 tarps or clear tape for the clear plastic tarps. Our collection of arthropods from pitfall traps in this
214 sampling coincided with the end of the tarp treatment.

215 The Berlese funnel method reflects soil-dwelling arthropod presence. For this method, we
216 took 3 soil cores (5 cm diameter, 10 cm deep) from each plot and composited them. We sampled
217 in a stratified random pattern, taking 1 core from each of 3 regions in each plot. Subsequently,

218 we extracted arthropods by placing collected soil in a funnel apparatus and exposing it to a 60-
219 Watt light bulb for 72 hours (Woolley, 1965; Southwood and Henderson, 2009). Arthropods
220 were collected and preserved in 95% ethanol.

221 Because pitfall traps are biased towards catching larger and more mobile organisms while
222 Berlese funnels are biased towards capturing smaller and less mobile organisms (Sabu et al.,
223 2011), we only analyze macrofauna from pitfall traps and mesofauna from Berlese funnels.
224 Macrofauna are generally defined as organisms larger than 2 mm (including small insects), while
225 mesofauna are between 0.2 – 2 mm (Lavelle et al., 1997; Gongalsky, 2021). Orders we include
226 from pitfall traps include: Araneae, Opilliones, Coleoptera, Hemiptera, Lepidoptera
227 (caterpillars), Orthoptera, Psocoptera, Thysanoptera, Diplopoda, Dermaptera, Isopoda, and
228 Lithobiomorpha. Orders we include from the Berlese funnels include: Symphypleona,
229 Astigmata, small insect larva, Protura, Entomobryomorpha, Oribatida, Mesostigmata,
230 Poduromorpha, Prostigmata, and Symphyla. We do not include ants (Hymenoptera) from either
231 sampling method because ants' central foraging behavior drives their organization on the
232 landscape and can make these sampling methods inadequate for determining their abundance
233 (Higgins and Lindgren, 2012). Abundance results for ants are included in the Appendix (Figure
234 A1).

235 In the lab, we used a stereo microscope to classify all soil arthropods to morphospecies.
236 While identifying organisms to species is valuable (Ward and Stanley, 2004), using
237 morphospecies can be an efficient and effective method to monitor arthropod communities, often
238 yielding similar numbers to species (Hackman et al., 2017).

239 We taxonomically identified all soil arthropods to the order level, while some were
240 identified to family, genus, or species level as resources, time, and expertise allowed (for

241 example, all beetles were identified to family). We identified arthropods using keys within
242 Dindal (1990) (for all arthropods), and Evans (2014) and Bousquet (2010) (for beetles). We also
243 utilized taxonomic resources and community identification on online sites bugguide.net (Iowa
244 State University, 2021) and iNaturalist.org (iNaturalist, 2021).

245

246 **2.4. Environmental variables**

247 We collected data on soil temperature, soil moisture, and weed coverage to further understand
248 their indirect effects on soil arthropod community composition. We chose these variables
249 because they represent major documented effects of tarps (Birthisel and Gallandt, 2019) and are
250 known to affect soil arthropod communities (Philpott et al., 2014; Gkisakis et al., 2016).

251 Soil temperature was automatically monitored every 30 minutes while tarps were on the
252 fields using iButtons (Thermochron, Baulkham Hills, NSW, Australia). Two iButtons were
253 buried in each plot — one at the surface (below 1 cm of soil) and one 10 cm below the surface.
254 We removed the iButtons at the time of tarp removal. We measured soil moisture using a
255 FieldScout TDR 350 Economy Soil Moisture Meter (Spectrum Technology, Inc., Aurora, IL,
256 USA) with 12.2 cm probe tips. We took point samples of soil moisture 3 weeks into the tarp
257 experiment, when arthropods were sampled, and subsequently every time arthropods were
258 sampled (1, 3, and 5 weeks after tarp removal; Table A1).

259 We surveyed weeds weekly after tarps were removed (Table A1). For each treatment
260 plot, we placed a 1 x 2 m sampling frame in the center of each plot. We visually estimated
261 percent cover for each present weed species using the following classes: less than 1%, 1–5%, 5–
262 15%, 15–25%, 25–50%, 50–75%, and 75–100% (Peet et al., 1998). For analysis, we converted
263 cover ranges to the midpoint of the range. We then calculated weed richness (total weed species

264 per plot) and total weed coverage by summing the cover for each species. Total weed cover
265 could exceed 100% when multiple layers of vegetation were present. Weeds were identified
266 using Uva et al. (1997).

267

268 **2.5. Statistical analyses**

269 We performed all statistical analyses in R version 4.0.4 (R Core Team, 2021) and used $P < 0.05$
270 to indicate significance. We first tested how abundances of dominant taxonomic groups differed
271 among treatments during tarp application. We determined the 7 most abundant orders (>95% of
272 all captured individuals) for surface-active and soil-dwelling arthropods each during tarp
273 application and tested how their abundance differed among treatments. To analyze data at a finer
274 resolution, we additionally tested how the abundance of the 7 most abundant Coleoptera families
275 (>95% of captured Coleoptera individuals) differed between treatments during tarp application.
276 We tested for significance using either linear mixed models or generalized linear mixed models
277 fit with the Poisson distribution, depending on residual structures, with farm as a random effect
278 with random intercepts (lmer and glmer in the lme4 package; Bates et al., 2015). We made
279 pairwise comparisons using estimated marginal means tests (emmeans in the emmeans package;
280 Lenth et al., 2018).

281 To understand how tarps impacted arthropod diversity, we calculated richness, Shannon's
282 diversity (vegan package; Oksanen et al., 2007), and total abundance separately for surface-
283 active and soil-dwelling arthropod morphospecies and at each sampling time (pre-tarps, during
284 tarping, and 1, 3, and 5 weeks after tarping). We used repeated measures linear mixed effects
285 models (lmer function from the lme4 package) for richness and Shannon's diversity data, and
286 repeated measures generalized linear mixed effects models fit with a negative binomial

287 distribution (glmer.nb function in the lme4 package) for abundance data. All models included
288 farm site as a random effect (with random intercepts). Model types were chosen based on which
289 yielded normality of residuals, which was tested using quantile-quantile plots and histograms.
290 We made multiple comparisons among treatments at each sampling period using estimated
291 marginal means tests with the emmeans package.

292 We then tested how morphospecies composition related to treatment using Principal
293 Coordinates Analysis (PCoA) of Bray-Curtis dissimilarities. We ran separate PCoAs for each
294 sampling period to see the impact and recovery of tarps on arthropod composition. We chose
295 PCoA because it performs well with species abundance data (McArdle and Anderson, 2001). We
296 ran the PCoA using the vegdist (vegan package) and wcmdscale functions, and then tested how
297 well treatment explained the composition data with a permutational multivariate analysis of
298 distance matrices (adonis function with “bray” distance method in vegan package). Pairwise
299 differences between treatments were also computed with the pairwise.perm.manova function in
300 the RVAideMemoire package (Hervé, 2018).

301 Finally, we created models to test how environmental and experimental variables,
302 including treatment, farm, soil moisture, soil temperature, weed coverage, and sampling time (1,
303 3, or 5 weeks after tarps were removed), related to the soil arthropod communities. “Treatment”
304 specifically refers to whether plots were treated with silage tarps, clear plastic, or the control, and
305 “farm” indicates at which of the three farms (Catamount, ICF, or Diggers’) the plots were
306 located. Our primary model structures were based off *a priori* assumptions of the ecological
307 system. In models representing conditions during tarp cover, we included an interaction term
308 between treatment and farm to understand whether tarps’ effects on arthropods differed by farm
309 site; for models representing the sampling periods after tarps were removed, we included an

310 interaction between treatment and sampling time to see if tarps' effects changed through time.
311 We tested model structures with AIC to find the most parsimonious models and avoid overfitting
312 (confirming at this step to only include one interaction term per model). We then assessed
313 collinearity among explanatory variables using variance inflation factors (VIF; vif function in the
314 car package; Fox et al., 2012), and removed variables with $GVIF^{(1/2*DF)}$ scores of over 5 (Fox and
315 Monette, 1992). The only variables we eliminated were soil surface temperature and soil (10 cm
316 below the surface) temperature, due to high collinearity with treatment (Figure A2). Final
317 variables of interest when tarps were on the fields included treatment, farm, soil moisture, and
318 the interaction between treatment and farm. Variables of interest when tarps were removed
319 included treatment, farm, soil moisture, weed coverage, sampling time, and the interaction
320 between treatment and sampling time.

321 With these variables, we used multiple linear regression models to predict soil arthropod
322 richness and redundancy analysis (RDA, a multivariate regression analysis and constrained
323 ordination technique; vegan package) to predict arthropod composition (we chose to build
324 models for richness rather than Shannon's diversity because the results were very similar and
325 richness is a more intuitive metric). We created separate models for when tarps were on and off
326 the fields because these times reflect different dynamics and numbers of samplings, thus having
327 different explanatory variables. We additionally created separate models for surface-active and
328 soil-dwelling arthropods, due to these groups being caught with different methods and
329 representing different ecological groups.

330 Within our final models, we tested for significance of the multiple linear regression
331 models using Type 3 ANOVAs from the car package. We chose Type 3 ANOVAs to test
332 variables regardless of their order in the model and to take into consideration interaction terms

333 (Shaw and Mitchell-Olds, 1993). We additionally assessed the relative importance of variables in
334 the linear regression models with the “lmg” method within the relaimpo package (Grömping,
335 2007). The “lmg” method computes the average R^2 over all possible model structures (orders of
336 variables), and we hereafter call this statistic “variable importance” (Lindeman, 1980). For the
337 RDA models, we tested significance using permutational ANOVAs (anova.cca function from the
338 vegan package), using the “margin” option to obtain Type 3 effects.

339

340 **3. Results**

341 **3.1. Summary of arthropod data**

342 We collected 8,027 surface-active (macrofauna) arthropods in total in the pitfall traps (2,710
343 from Catamount, 2,591 from Diggers’, and 2,726 from ICF), comprising 102 morphospecies and
344 12 orders. At the order level, the most abundant group were beetles with 5,559 total individuals.
345 Beetles were also the order with the most morphospecies. Of the 75 identified beetle
346 morphospecies, all were identified to family, 43 were identified at least to genus (57.9% of the
347 beetle specimens), and 15 were identified to species (14.7% of the beetle specimens).

348 In the Berlese funnel samples, we collected 823 soil-dwelling (mesofauna) arthropods,
349 comprising 20 morphospecies and 10 total orders. Collembolans, mostly from the order
350 Entomobryomorpha, comprised 48% of the Berlese samples, while mites, mostly Prostigmata,
351 made up 39% of the samples.

352

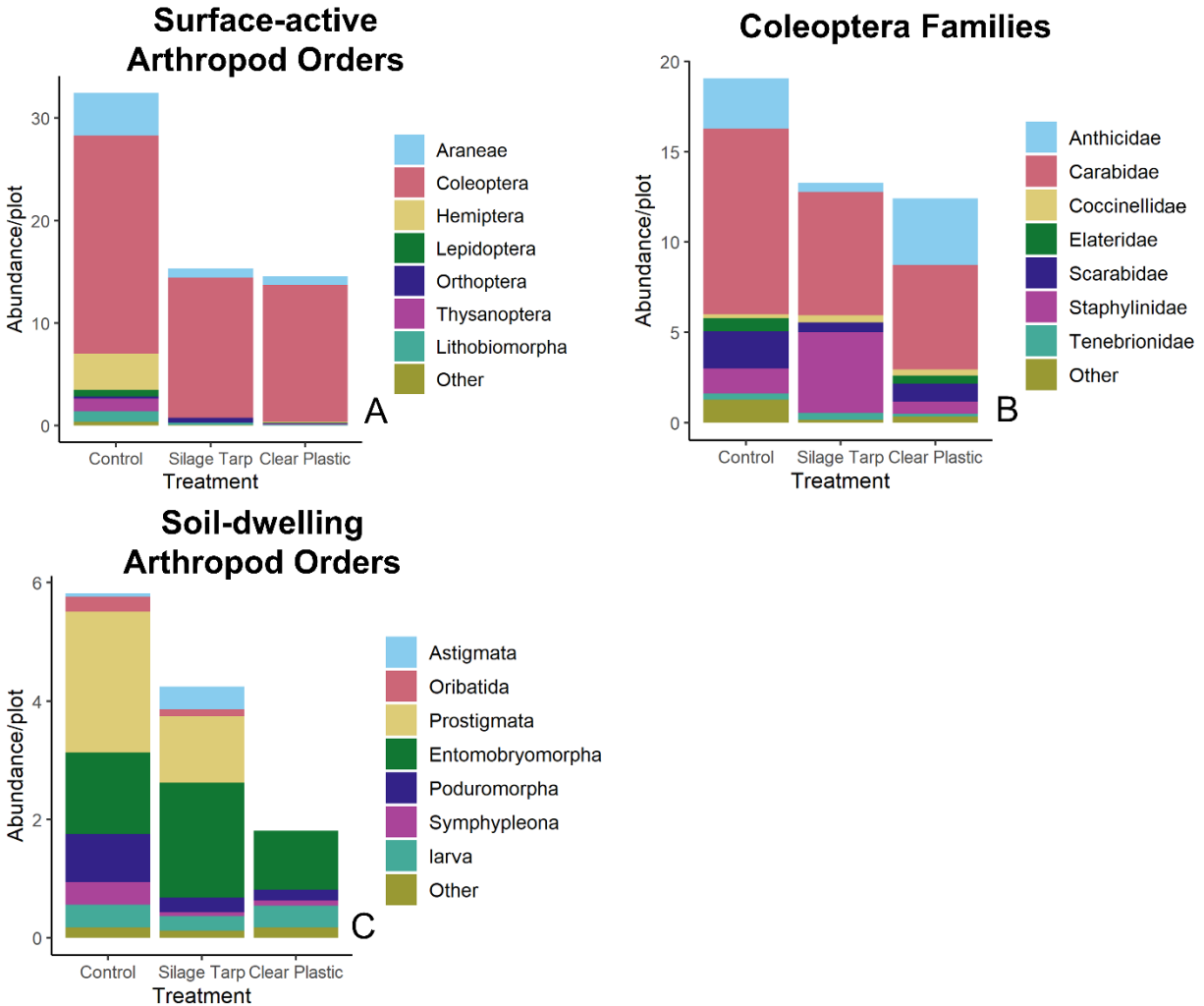
353 **3.2. Taxa abundance during tarping**

354 Tarp application significantly affected many surface-active arthropod taxa. At the order level,
355 most orders — including Araneae, Coleoptera, Hemiptera, Thysanoptera, and Lithobiomorpha

356 — had significantly lower abundances in the tarped plots than control plots (Figure 1A, Table
357 A2). Changes amongst Coleoptera families were less consistent. Compared to the control,
358 Carabidae and Scarabidae were both less abundant in the silage tarp ($P = 0.001$ & $P < 0.001$,
359 respectively) and clear plastic tarp plots ($P < 0.001$ & $P = 0.031$), while Anthicidae were less
360 abundant in the silage tarp plots ($P < 0.001$) but not clear plastic tarp plots ($P = 0.296$).
361 Conversely, Staphylinidae individuals were more than twice as abundant in silage tarp plots than
362 control or clear plastic tarp plots (both $P < 0.001$). Other families, like Coccinellidae, Elateridae,
363 and Tenebrionidae, had lower abundances and did not significantly differ among treatments
364 (Figure 1B, Table A2).

365 For soil-dwelling arthropods, clear plastic tarps had a negative impact on several orders.
366 We found lower abundances in clear plastic than in control plots for Prostigmata ($P = 0.004$) and
367 Poduromorpha springtails ($P = 0.035$), but there were no other significant differences between
368 treatments for these groups. Entomobryomorpha springtails were most abundant under silage
369 tarps and least abundant under clear plastic tarps (silage tarp - clear plastic $P = 0.008$). There
370 were no differences among treatments for Symphypleona springtails or insect larva (Figure 1C;
371 Table A2).

372



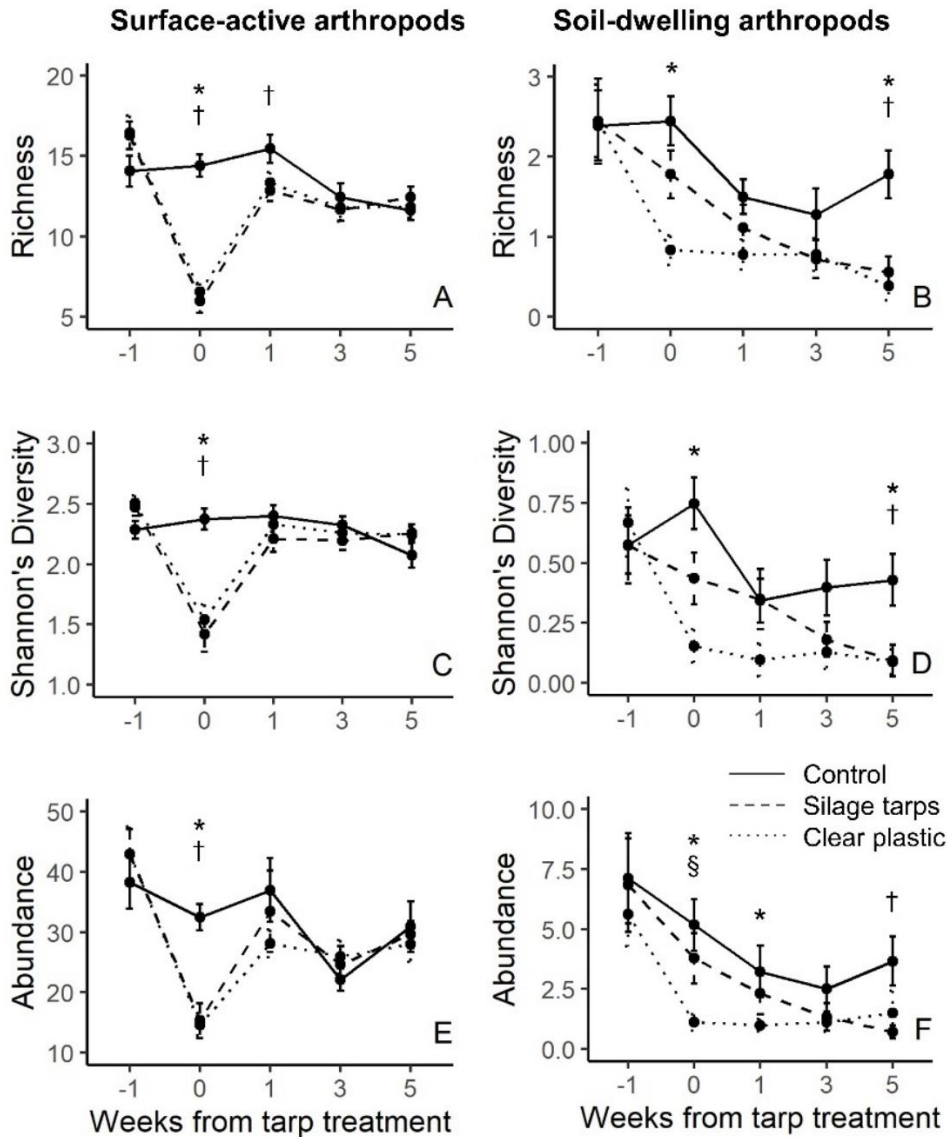
373 Figure 1: Abundances of individuals within surface-active arthropod orders (A), Coleoptera
374 families (B), and soil-dwelling arthropod orders (C) among the treatments when tarps were on
375 the field. “Abundance/plot” relates to the average number of individuals captured per pitfall trap
376 (A, B) or Berlese funnel sample (B).

377

378 **3.3. Diversity and abundance**

379 For surface-active arthropods, during the tarping treatment both silage tarps and clear plastic
380 tarps had significantly lower richness, Shannon's diversity and total abundance than the control
381 plots (all $P < 0.001$; Figure 2A,C,E; all statistics in Table A3). One week after we removed the
382 tarps, silage tarps had lower richness than control plots ($P = 0.044$), but there were no other
383 significant differences. At 3 and 5 weeks after tarp removal, there were no significant differences
384 for richness, Shannon's diversity, or abundance among the treatments.

385 During the tarping treatment, richness and Shannon's diversity of soil-dwelling
386 arthropods were significantly lower in the clear plastic tarp plots than in the control plots (both P
387 < 0.001 ; Figure 2B,D; Table A3). Abundance of soil-dwelling arthropods during the tarp
388 treatment was significantly lower in the clear plastic tarps compared to both control ($P = 0.002$)
389 and silage tarp plots ($P = 0.034$; Figure 2F). Soil-dwelling arthropod abundance remained
390 significantly lower in the clear plastic tarp plots than the control 1 week after tarp removal ($P =$
391 0.033). While there were no differences in richness or Shannon's diversity values 1 and 3 weeks
392 after tarp removal, 5 weeks after tarp removal silage tarp and clear plastic tarp plots had
393 significantly lower richness ($P = 0.011$ & $P = 0.050$) and Shannon's diversity values ($P = 0.003$
394 & $P = 0.045$) than control plots. Additionally, abundance was significantly lower in silage tarp
395 plots than control plots 5 weeks after tarp removal ($P = 0.004$; Figure 2F; Table A3).

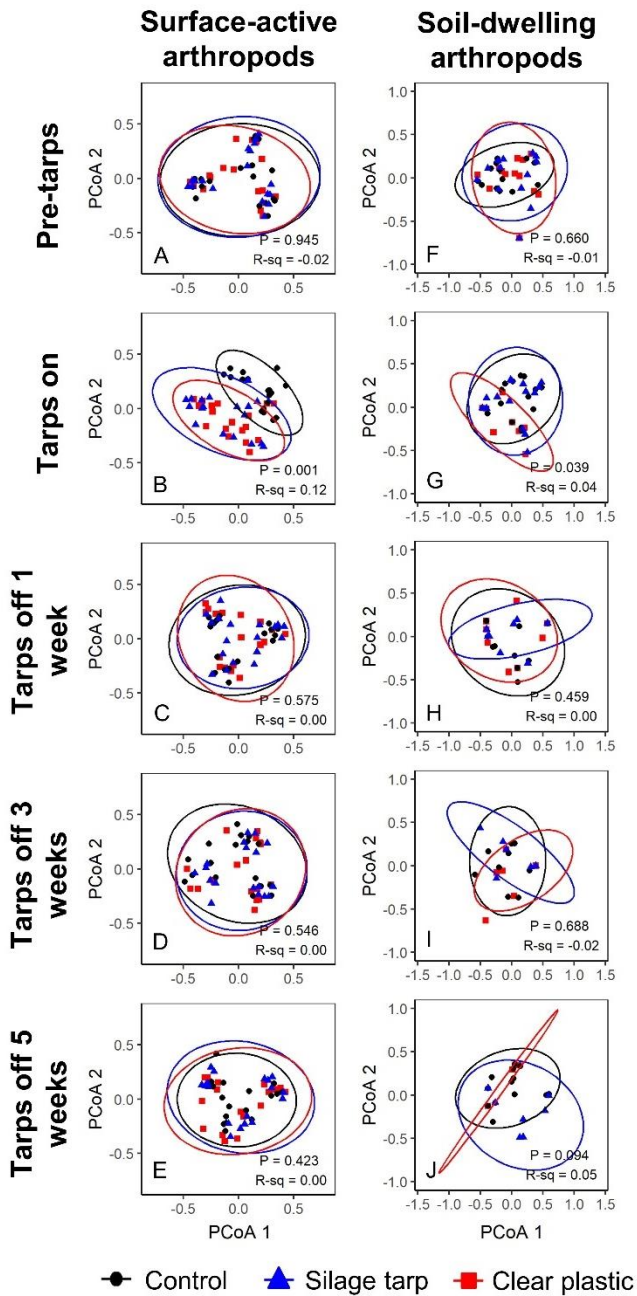


397 Figure 2: Impact of tarps on richness, Shannon's diversity, and abundance for surface-active and
 398 soil-dwelling arthropod morphospecies. Values indicate change from baseline (a pre-tarp
 399 sampling). An asterisk (*) indicates significant differences between control and silage tarp plots,
 400 † indicates significant difference between control and clear plastic tarp plots, and § indicates
 401 significance difference between silage tarp and clear plastic plots. "-1" represents the week
 402 before tarps were applied, "0" represents sampling the last week of the tarp treatment, and "1",
 403 "3", and "5" indicate weeks since tarp removal.

404 **3.4. Composition analyses**

405 In the principal coordinates analyses (PCoA), treatment significantly explained surface-active
406 communities during the tarp treatment ($P < 0.001$; $R^2 = 0.12$), with significant pairwise
407 differences among all treatment pairs (control – silage tarp $P < 0.001$; control – clear plastic $P <$
408 0.001 ; silage tarp – clear plastic $P = 0.039$; Figure 3B). Treatment was also a significant
409 predictor of soil-dwelling arthropod communities during the tarp treatment ($P = 0.039$; $R^2 =$
410 0.04), but there were no differences between treatment pairs (control – silage tarp $P = 0.064$;
411 control – clear plastic $P = 0.064$; silage tarp – clear plastic $P = 0.381$; Figure 3G). After tarps
412 were removed, treatment no longer significantly explained arthropod composition in the PCoAs
413 for either surface-active or soil-dwelling arthropods, and there were additionally no significant
414 differences before tarps were applied (Figure 3).

415



417 Figure 3: Principal coordinates analysis (PCoA) showing the separation of surface-active (A-E)
 418 and soil-dwelling (F-J) arthropod morphospecies composition in each treatment. Trends are
 419 shown for each sampling period. R^2 and P values in each plot correspond to the fit and
 420 significance of treatment.

421

422 **3.5. Importance of experimental and environmental variables**

423 When tarps were on the fields, environmental and experimental variables explained over 75% of
424 the variation in surface-active arthropod richness and around 43% of the variation in surface-
425 active arthropod composition (Table 2). Treatment had the greatest variable importance score,
426 and both treatment and the interaction between treatment and farm significantly explained both
427 surface-active arthropod richness and composition (Figure A3, Table A4 and A5). Farm
428 additionally significantly explained surface-active composition.

429 For soil-dwelling arthropods, our models explained around 44% of the variation in
430 richness and 28% of the variation in composition during tarp cover (Table 2). Treatment again
431 had the greatest variable importance score and significantly explained soil-dwelling arthropod
432 richness, though no other variables were significant.

433 When tarps were removed, our models explained around 20% and 30% of the variability
434 in surface-active arthropod richness and composition, respectively (Table 2). Farm site had the
435 highest variable importance score and was significant in both models. While no other variables
436 significantly explained surface-active arthropod richness, composition was significantly
437 explained by sampling period, soil moisture, and treatment.

438 Our model for soil-dwelling arthropod richness and composition showed similar results
439 (model R^2 of 23% and 24%, respectively). While treatment again had the largest variable
440 importance score and was significant, farm site was additionally a significant predictor for both
441 models. No other variables were significant predictors of soil-dwelling arthropods after tarp
442 removal, and the interaction between treatment and sampling was not significant in any models
443 (Figure A3). Summaries of how environmental variables differed among treatments can be
444 found in the Appendix (Table A6).

445 Table 2: Results from models predicting arthropod morphospecies richness (multiple linear
 446 regression) and composition (RDA) with environmental and experimental variables. We include
 447 F statistics and P-values from all models and variable importance (standardized regression
 448 coefficients, totaling 1) for the richness models. Models were run separately for surface-active
 449 and soil-dwelling arthropods and for two sampling periods: when tarps were on the fields (“tarps
 450 on”), and 1, 3, and 5 weeks after tarp removal (“tarps off”).
 451

	Surface-active arthropods					Soil-dwelling arthropods				
	Richness			Composition		Richness			Composition	
	Variable importance	F	P	F	P	Variable importance	F	P	F	P
Tarps on										
Treatment	0.84	7.40	0.002	3.21	0.001	0.63	7.31	0.002	1.48	0.131
Farm	0.05	3.06	0.057	4.03	0.001	0.03	0.14	0.874	1.55	0.082
Soil moisture	0.02	2.76	0.104	1.54	0.098	0.10	1.06	0.308	0.53	0.809
Treatment:Farm	0.09	3.00	0.029	2.21	0.001	0.24	1.97	0.116	1.18	0.248
<i>Total model fit (R²)</i>	76.6%			42.6%		43.7%			27.9%	
Tarps off										
Treatment	0.10	1.34	0.265	1.39	0.044	0.40	6.01	0.003	1.20	0.233
Farm	0.35	6.21	0.003	3.23	0.001	0.31	6.56	0.002	4.59	0.002
Sampling	0.28	2.05	0.132	13.9	0.001	0.08	0.45	0.454	1.57	0.143
Soil moisture	0.06	0.70	0.405	3.11	0.001	0.01	0.48	0.489	0.62	0.641
Weed coverage	0.09	0.57	0.452	0.74	0.738	0.11	0.01	0.918	2.26	0.076
Treatment: Sampling	0.12	0.80	0.525	0.99	0.479	0.09	0.91	0.462	1.11	0.336
<i>Total model fit (R²)</i>	20.3%			31.3%		22.9%			23.6%	

452

453 **4. Discussion**

454 Tarps had immediate detrimental impacts on the diversity of both surface-active and soil-
455 dwelling arthropods and changed arthropod community composition, supporting other studies in
456 tarping and plastic mulch research (Seman-Varner, 2005; Tuovinen et al., 2006; Addison et al.,
457 2013). While surface-active arthropod diversity recovered within 1-3 weeks after tarps were
458 removed, soil-dwelling arthropods showed a less clear recovery — five weeks after tarps were
459 removed, soil-dwelling arthropod richness was significantly lower in plots that had been tarped
460 than control plots. These results suggest that tarps’ impacts may be temporary for surface-active
461 arthropods but could be longer lasting for soil-dwelling arthropods.

462

463 **4.1. Impacts during tarp application**

464 While tarps affect the soil ecosystem in a variety of ways, including soil sealing and being
465 impermeable, a large factor likely determining arthropod responses to tarping is heat tolerance.
466 As ectotherms with no (or little) internal control over their body temperature, arthropods are
467 susceptible to external temperature fluctuations. Their ability to withstand heat is highly
468 interspecific and poorly understood, likely relating to body size, exoskeleton color and thickness,
469 life stage, and trophic level, among other factors (Franken et al., 2018; González-Tokman et al.,
470 2020).

471 For larger and more mobile arthropods, tarping may trigger a “stay” or “go” response.
472 For instance, low abundance of many surface-active arthropod orders under tarps likely
473 represents migration out of the tarped area. However, other arthropods may be more resilient or
474 unbothered by tarps’ effects, such as detected for Coleoptera (Figure 1B). Coleoptera have a
475 relatively thick cuticle, which may make them more heat tolerant and resistant of desiccation

476 compared to other invertebrates (Wikars and Schimmel, 2001). High mobility of many
477 Coleoptera taxa may also allow them to pass through tarps quickly without succumbing to heat
478 effects. Mobile arthropods may also take cover under tarps during cooler temperatures or as a
479 shelter from predators.

480 In contrast, soil-dwelling arthropods have generally low mobility, with some taxa moving
481 as little as only a few centimeters per day (Ojala and Huhta, 2001). Therefore, with low dispersal
482 ability, soil-dwelling arthropod composition under tarps may reflect a “live” or “die” response.
483 We found that soil-dwelling arthropods were affected by clear plastic tarps but not silage tarps
484 during tarp application, suggesting that higher soil temperatures under clear plastic tarps (8°C
485 warmer than silage tarps) were inhospitable (Table A6, Figure A2). The impact of silage tarps on
486 surface-active arthropods but not soil-dwelling arthropods during tarp application may support
487 the theory that small-bodied organisms have higher heat tolerances than larger organisms (Smith
488 et al., 2009; Sheridan and Bickford, 2011), or may simply reflect the decreasing temperature
489 effects of tarps with soil depth (Oz et al., 2017).

490

491 **4.2. Recovery of arthropods after tarp removal**

492 In response to disturbances, like tarping, biological communities can have very different
493 trajectories, with some engaging in recovery (Moretti et al., 2006; Pryke and Samways, 2012) –
494 as seen for surface-active arthropods – while others experience diversity declines (Birthisel et al.,
495 2019) – as seen for soil-dwelling arthropods. Recovery (or recolonization) of arthropods after
496 tarps reflects either the dispersal of organisms back into the disturbed space or the regeneration
497 of populations and communities (Bengtsson, 2002). Because our farm sites were adjacent to
498 forested areas, there was likely high dispersal of larger and more mobile arthropods back into our

499 experimental plots after tarping. Conversely, the dispersal of less mobile soil-dwelling
500 arthropods may have been limited during our relatively short study, though it is possible that
501 these arthropods engaged in vertical dispersal (Moradi et al., 2020). To better understand the role
502 of dispersal, a study specifically looking at arthropods' movement patterns would be useful
503 (Perry et al., 2021).

504 To support population regeneration, certain requirements need to be met, including
505 having sufficient numbers of mates (for sexually reproducing organisms), food availability, lack
506 of competition and predation, and abiotic suitability (Menge and Sutherland, 1987). For tarps,
507 the spatial scale of effects may be felt differently for surface-active and soil-dwelling
508 communities. For larger arthropods, tarps may create heterogeneity within their habitat, but not
509 have a large enough impact to prohibit them from finding resources or mates. Conversely,
510 smaller and less mobile arthropods, especially those living belowground, depend more on local
511 conditions, especially because some are restricted to movement within existing soil pore
512 networks (Vreeken-Buijs et al., 1998), and thus the impact of tarps may encompass their entire
513 range. Tarps may also have fundamental impacts on the food resources, community and
514 population dynamics, and habitat conditions for soil-dwelling organisms, creating complex and
515 even cascading effects (Bengtsson, 2002). All these effects may explain why we see a fast
516 recovery of surface-active arthropods and a less clear recovery of soil-dwelling arthropods.

517

518 **4.3. Importance of environmental and experimental variables**

519 The importance of experimental and environmental variables differed when tarps were on the
520 field and after they were removed. During tarping, treatment was extremely predictive of soil
521 arthropods but after tarp removal, while it remained significant in some models, its relative

522 importance declined (Table 2). Conversely, farm was not as predictive as treatment in most
523 models during tarping but became one of the most predictive variables after tarps were removed.
524 The strength of farm at predicting arthropod richness and composition is not surprising, as our
525 three farm sites had considerable differences, including for soil texture, soil nutrient profiles, and
526 weed composition (Table 1), and such biophysical differences can lead to unique arthropod
527 communities (Schaffers et al., 2008; Philpott et al., 2014; Ghiglieno et al., 2021). The result that
528 farm site was relatively not as important during tarping demonstrates the strong impact of tarping
529 at driving trends. However, we did interestingly find a significant interaction between treatment
530 and farm during tarping, showing that tarps' effects may differ depending on the site.
531 Specifically, we found that tarps had less effect on surface-active arthropod richness at
532 Catamount Farm (Figure A3; farm-separated data summaries listed in Tables A4 and A5). An
533 explanation for this is that Catamount had ambiently higher soil temperatures than the other
534 farms (by 1-2°C) due to sandy soil, and thus arthropods there may have been more thermally
535 adapted (Brans et al., 2017).

536 Other factors within our system, including soil moisture, weed coverage, and the
537 sampling period, less consistently explained arthropod dynamics. Both weed coverage and soil
538 moisture can be important drivers of soil arthropod communities (Norris and Kogan, 2000; Gear
539 and Schmitz, 2005; Tsiafouli et al., 2005), and we had particularly expected weed coverage to
540 relate to soil arthropod communities, as a possible food source and habitat. Lack of relationships
541 for weed coverage and soil moisture may be due to relationships with other variables, for
542 example between soil moisture and farm and weed coverage and sampling (though collinearity
543 was not detected for these variable pairs). Similarly, we likely did not detect a strong effect of
544 sampling because of the quick recovery of arthropods following tarping and because, while

545 arthropod richness patterns can change inter-annually (Liu et al., 2016; Kirse et al., 2021),
546 sampling for five weeks after tarping may not have been enough time to see significant changes.

547 Finally, while soil temperature was removed from the models due to high collinearity
548 with treatment (Table A6), we found a significant negative relationship between soil temperature
549 and arthropod richness (Figure A2), though it is difficult to decouple the effects here attributable
550 only to soil temperature. Tested models for surface-active arthropod richness including soil
551 temperature but not treatment performed relatively poor (yielding $R^2 = 0.40$, compared to the
552 current $R^2 = 0.77$), demonstrating that tarps' effects stretch beyond temperature effects and
553 losses in soil arthropod richness are driven by other factors as well (such as light-availability
554 differences or soil sealing).

555

556 **4.4. Experimental limitations**

557 A major limitation in our study is that our tarped treatment plots are much smaller than tarps
558 used in practice — our tarps were 2 by 4.5 m, while tarps in practice are often 10 by 15 m or
559 larger. Furthermore, while we sealed the edges of tarps with soil, some farmers use sandbags to
560 hold tarp edges down (especially for silage tarps), which may allow for more airflow. Larger tarp
561 sizes may make it more difficult for mobile organisms to migrate out of the tarped area,
562 potentially causing more negative effects. Conversely, increased airflow and ultimately lower
563 temperatures under tarps may lead to less negative temperature-related diversity declines. While
564 we decided to use small tarp pieces to maximize the replication of treatments on our limited land
565 area, scaling up this experiment might provide more insights.

566 The long-term impacts of tarps also remain unclear. Our study was limited to 5 weeks
567 after tarp removal, but effects may continue long after this period — in studies on fire, soil

568 arthropod communities took decades to recover (Pressler et al., 2019). As well, while we saw
569 recovery of surface-active arthropods during our experimental time frame, recovery patterns are
570 complex and not always linear; thus, sampling for longer periods of time may reveal different
571 dynamics than we observed. Additionally, many farmers use tarps twice or even three times a
572 summer for quick growing crops. The continuous use of tarps could expound effects by not
573 allowing communities to fully recover. Another possible outcome of frequent use of tarps is
574 adaptation. In fact, studies on urbanization suggest that arthropods can adapt to warm
575 temperatures (Diamond et al., 2017; Yilmaz et al., 2021). It is important to consider these
576 potential long-term changes to species, community structure, and biological function.

577

578 **4.5. Conclusions and implications for agricultural management**

579 Tarps are an exciting new practice which may help farmers transition away from intensive
580 practices like tillage and herbicide use, but we are only starting to understand the impacts of
581 tarps on biodiversity. This study has unveiled important information on the short-term effects of
582 tarps on arthropods but, going forward, more research will be necessary to contextualize our
583 results. For example, it will be valuable to compare the effects of tarps and other weed
584 management techniques, or to conduct tarp research over multiple years and in different
585 geographic regions, seasons, and soil types. Additionally, while we did not specifically look at
586 tarps' impacts on pests, this has been one major application of tarps (Stapleton and DeVay,
587 1986) and remains a topic of interest among farmers (Kinnebrew et al., 2022a). We encourage
588 future research on tarps' effects on pests, though suggest consideration of concurrent impacts on
589 beneficial arthropods, including on natural enemies of pests. In conclusion, we hope this study

590 helps inform agricultural management that can be effective for both crop production and
591 biodiversity conservation (Díaz et al., 2015).

592

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