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2  
3 **Title: Increasing crop heterogeneity enhances multitrophic diversity across agricultural**  
4 **regions**

5  
6 **Short title: Crop heterogeneity and multitrophic diversity**

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81 complementation

82 **ABSTRACT**

83 Agricultural landscape homogenization has detrimental effects on biodiversity and key  
84 ecosystem services. Increasing agricultural landscape heterogeneity by increasing semi-  
85 natural cover can help to mitigate biodiversity loss. However, the amount of semi-natural  
86 cover is generally low and difficult to increase in many intensively-managed agricultural  
87 landscapes. We hypothesized that increasing the heterogeneity of the crop mosaic itself  
88 (hereafter “crop heterogeneity”) can also have positive effects on biodiversity. In eight  
89 contrasting regions of Europe and North America, we selected 435 landscapes along  
90 independent gradients of crop diversity and mean field size. Within each landscape, we  
91 selected three sampling sites in one, two or three crop types. We sampled seven taxa (plants,  
92 bees, butterflies, hoverflies, carabids, spiders, birds) and calculated a synthetic index of  
93 multitrophic diversity at the landscape level. Increasing crop heterogeneity was more  
94 beneficial for multitrophic diversity than increasing semi-natural cover. For instance, the  
95 effect of decreasing mean field size from 5 to 2.8 ha was as strong as the effect of increasing  
96 semi-natural cover from 0.5 to 11 %. Decreasing mean field size benefited multitrophic  
97 diversity even in the absence of semi-natural vegetation between fields. Increasing the number  
98 of crop types sampled had a positive effect on landscape-level multitrophic diversity.  
99 However, the effect of increasing crop diversity in the landscape surrounding fields sampled  
100 depended on the amount of semi-natural cover. Our study provides the first large-scale,  
101 multitrophic, cross-regional evidence that increasing crop heterogeneity can be an effective  
102 way to increase biodiversity in agricultural landscapes without taking land out of agricultural  
103 production.

104

105 **SIGNIFICANCE STATEMENT**

106 Agricultural landscape homogenization is a major ongoing threat to biodiversity and the  
107 delivery of key ecosystem services for human well-being. It is well known that increasing the  
108 amount of semi-natural cover in agricultural landscapes has a positive effect on biodiversity.  
109 However, little is known about the role of the crop mosaic itself. Crop heterogeneity in the  
110 landscape had a much stronger effect on multitrophic diversity than the amount of semi-  
111 natural cover in the landscape, across 435 agricultural landscapes located in eight European  
112 and North American regions. Increasing crop heterogeneity can be an effective way to  
113 mitigate the impacts of farming on biodiversity without taking land out of production.

## 114 INTRODUCTION

115 Agriculture dominates the world's terrestrial area (1, 2). Agricultural landscape  
116 homogenization through the decrease of semi-natural cover, crop specialization and field  
117 enlargement (3–6) represents a continuing worldwide threat to biodiversity and the delivery of  
118 key ecosystem services to people (7, 8). There is ample evidence that enhancing landscape  
119 heterogeneity by reversing the decline in semi-natural cover can benefit biodiversity in  
120 agricultural landscapes (9–12). However, the amount of semi-natural cover keeps decreasing  
121 in many agricultural landscapes, and the efficiency of policies focusing solely on maintaining  
122 or increasing semi-natural cover has been questioned (13).

123 While half of the biodiversity in agricultural landscapes occurs exclusively in semi-  
124 natural cover (14), the crop mosaic offers a wide range of resources to the other half,  
125 including to species occurring exclusively in crop fields and providing key ecosystem  
126 services, such as crop pollination or biological pest control (15–17). It is therefore of  
127 increasing interest to evaluate whether enhancing landscape heterogeneity by increasing the  
128 heterogeneity of the crop mosaic itself (hereafter “crop heterogeneity”) can also benefit  
129 biodiversity (Fig. 1). There is growing pressure on agricultural land for food and energy  
130 production as well as for urbanization. Therefore, measures to benefit biodiversity consisting  
131 of a re-arrangement of the production area, as opposed to measures focusing solely on its  
132 reduction, could provide valuable new sustainable policy options.

133 Crop heterogeneity can be decomposed into compositional heterogeneity, i.e. the  
134 composition of the crop mosaic (e.g. crop diversity), and configurational heterogeneity, i.e.  
135 the shape and spatial arrangement of fields (e.g. mean field size, 18; see further explanation in  
136 *Methods*). These two components of crop heterogeneity may influence farmland biodiversity  
137 in several ways (see detailed alternative hypotheses in SI 1). First, increasing crop diversity  
138 may benefit biodiversity if many species are specialists of distinct crop types (i.e. habitat

139 specialization; **Hyp 1a** in **SI 1**; 19). In that case, sampling increasing numbers of crop types  
140 should lead to observing increasing levels of species diversity. Second, increasing crop  
141 diversity may also benefit biodiversity through a landscape-level effect if many species  
142 require multiple resources provided by different crop types (i.e. landscape complementation;  
143 **Hyp 1b** in **SI 1**; 20). In that case, sampling a given number of crop types surrounded by  
144 increasing levels of crop diversity available in the landscape should lead to observing  
145 increasing levels of species diversity. Third, decreasing mean field size may benefit  
146 biodiversity through a landscape-level effect if small fields provide easier access to adjacent  
147 crop fields for many species (i.e. landscape complementation; **Hyp 2a** in **SI 1**; 20, 21). In that  
148 case, sampling a given number of fields surrounded by fields with decreasing mean sizes  
149 should lead to observing increasing levels of species diversity.

150         Biodiversity responses to crop heterogeneity may be non-linear and non-additive. For  
151 instance, increasing the diversity of crops available in the landscape may benefit biodiversity  
152 in a given field only if fields are small enough for adjacent fields to be reached easily.  
153 Additionally, the effects of increasing crop heterogeneity on biodiversity may depend on the  
154 amount of semi-natural cover in the landscape. For instance, the ‘intermediate landscape-  
155 complexity’ hypothesis (22) predicts that the positive biodiversity-crop heterogeneity  
156 relationship is stronger in landscapes with intermediate amounts of semi-natural habitats (e.g.  
157 5-20%) than in landscapes with little (e.g. <5%) or much semi-natural habitat (e.g. >20%; 10).  
158 Sampling over a wide range of landscapes may therefore be necessary to understand the  
159 general effect of crop heterogeneity on farmland biodiversity.

160         The biodiversity-crop heterogeneity relationship may vary among taxa (e.g. 23, 24).  
161 For instance, it may be more positive for species and taxa that have lower habitat area  
162 requirements (e.g. small species; 25) or higher habitat specialization levels (e.g. 26). Although  
163 in-depth understanding of the effects of crop heterogeneity on each species or taxon is

164 valuable, it is also critical to develop environmental policies that are effective across a wide  
165 range of species (27, 28). To achieve this, we here use a cross-regional sampling scheme in  
166 Europe and North America and a synthetic index integrating information on multiple trophic  
167 groups in order to identify landscape patterns that simultaneously increase the diversity of  
168 most taxa (29).

169 We selected 435 landscapes along orthogonal gradients of mean size and diversity of  
170 crop types available in the landscape in eight contrasting agricultural regions in France, the  
171 United Kingdom, Germany, Spain and Canada (Fig. S2.1 in SI 2). In each landscape, we  
172 selected three sampling sites in one, two or three crop types. We sampled seven taxa  
173 representing a wide range of ecological traits, functions and trophic levels (plants, bees,  
174 butterflies, hoverflies, carabids, spiders and birds) in each field. We then computed a synthetic  
175 index of multitrophic diversity (*Methods*). We tested the relative effects of mean field size,  
176 the number of crop types sampled, the diversity of crop types available in the landscape, and  
177 the amount of semi-natural cover in the landscape on multitrophic diversity and on the species  
178 richness of taxonomic groups. We also evaluated whether the effects of mean field size and  
179 the diversity of crop types available in the landscape were non-linear, non-additive, and  
180 influenced by semi-natural cover (see detailed hypotheses in SI 1).

181

## 182 **RESULTS AND DISCUSSION**

183 Our study provides the first large-scale evidence that crop heterogeneity is a major driver of  
184 multitrophic diversity in agricultural landscapes. The number of crop types sampled in the  
185 landscape, the mean size and diversity of crop types available in the landscape were  
186 consistently included in all models (Fig. 2A). Together, they accounted for 61% of the  
187 explained variance in multitrophic diversity, while semi-natural cover accounted for 24%  
188 (Fig. 2B). Interactions between semi-natural cover and mean size/crop diversity of fields

189 available in the landscape also accounted for an important part of the explained variance  
190 (15%), indicating that the effects of crop heterogeneity is modulated by the amount of semi-  
191 natural cover in the landscape (Fig. 3). The effects of crop heterogeneity on multitrophic  
192 diversity were consistent across the eight European and North American regions (Fig. 4). The  
193 effects of crop heterogeneity on the species richness of taxonomic groups were similar to their  
194 effects on multitrophic diversity and similar across the seven taxa (Fig. 5 and Fig. S5.2 in SI  
195 5). They hold true when considering either landscape-level or field-level multitrophic  
196 diversity, including when focusing only on cereal fields, the most dominant crop type across  
197 our eight regions (Table S5.11 in SI 5). Their effects were also unchanged when potential  
198 confounding factors such as the identity of crop types sampled, land-use intensity within  
199 fields sampled (i.e. an index combining data on ploughing, fertilizer, herbicide and  
200 insecticide), the composition of the crop mosaic, grassland cover or hedgerow length  
201 available in the landscape were taken into account in our analyses (see additional analyses in  
202 SI 5).

203

#### 204 **Consistent positive effects of decreasing mean field size on multitrophic diversity**

205 Decreasing mean field size was the main driver of multitrophic diversity variations, mean  
206 field size and mean field size<sup>2</sup> together accounting for 47.4% of the explained variance in  
207 multitrophic diversity (Fig. 2B). The effect of decreasing mean field size from 5 to 2.75 ha  
208 was as strong as the effect of increasing semi-natural cover from 0.5 to 11 % of the landscape  
209 (Fig. 3B). Such a positive effect of decreasing mean field size on multitrophic diversity is  
210 consistent with the hypothesis that smaller fields provide easier access to multiple cover  
211 patches, in particular for species that require resources occurring in different cover types  
212 (landscape complementation; 20, 21). The positive effect of decreasing mean field size was



213 particularly clear and strong when mean field size fell below 6 ha (93% of landscapes  
214 studied).

215 Although the strength of this effect varied significantly among regions, decreasing  
216 mean field size had a consistent positive effect across all regions studied (Fig. 4 and section  
217 5.3 in SI 5). It was also consistently positive across all group of taxa considered separately,  
218 from primary producers to predators (Fig. 5 and section 5.4 in SI 5). Previous studies have  
219 already reported positive effects of decreasing mean field size on the diversity of several taxa  
220 considered separately (30–34). Our study, based on multiple regions and multiple trophic  
221 groups, shows that the benefits of decreasing mean field size can be generalized to  
222 multitrophic diversity across a wide range of agricultural regions.

223 Previous studies suggested that the positive effect of decreasing mean field size on  
224 multitrophic diversity may be primarily due to the presence of semi-natural vegetation  
225 between fields (30–34). To test this hypothesis, we selected a subset of landscapes for which  
226 mean field size and the length of semi-natural vegetation between fields were uncorrelated  
227 (see details in section 5.5.3 in SI 5). The analysis, based on 274 landscapes, showed that the  
228 positive effect of increasing mean field size on multitrophic diversity cannot be explained  
229 solely by the increase in the length of semi-natural vegetation between fields. Increasing the  
230 amount of semi-natural vegetation between fields had a positive effect on multitrophic  
231 diversity but including this effect in our model did not change the effect of mean field size on  
232 multitrophic diversity (Table S5.8 in SI 5). This result suggests that smaller fields benefit  
233 multitrophic diversity even in the absence of semi-natural vegetation between fields.

234 Finally, the presence of the interaction term between mean field size and semi-natural  
235 cover in our model (Fig. 2A) suggests that the effect of mean field size on multitrophic  
236 diversity tends to be modulated by the amount of semi-natural cover available in the  
237 landscape (Fig. 3B). To further explore this interaction, we used a moving window modeling

238 approach (35; see details in [section 5.7](#) in [SI 5](#)). This analysis confirmed that decreasing mean  
239 field size had a consistent positive effect on multitrophic diversity along the gradient of semi-  
240 natural cover. Moreover, it suggested that this effect is stronger when semi-natural cover is  
241 below 8%, i.e. when semi-natural cover is too scarce to provide access to the multiple  
242 resources required by most species occurring in agricultural landscapes ([Fig. S5.5.B](#) in [SI 5](#)).

243

#### 244 **Complex effects of increasing crop diversity on multitrophic diversity**

245 The number of crop types sampled in each landscape and the diversity of crop types available  
246 in the landscape surrounding sampled fields were consistently included in all models ([Fig.](#)  
247 [2A](#)). This result suggests that both field-level (i.e. habitat specialization) and landscape-level  
248 processes (i.e. landscape complementation and/or spill-over) can contribute to the effect of  
249 crop diversity on multitrophic diversity (see further explanations in [SI 1](#) and [section 4.4.](#) in [SI](#)  
250 [4](#)).

251         Increasing the number of crop types sampled had a significant positive effect  
252 accounting for 13% of the explained variance in landscape-level multitrophic diversity ([Fig.](#)  
253 [2B](#)). This result confirms that increasing crop diversity results in a larger number of distinct  
254 habitats, and therefore higher biodiversity levels by increasing the number of specialist  
255 species in the landscape ([Hyp 1a](#) in [SI 1](#), 26).

256         The main effect of increasing the diversity of crop types available in the landscape was  
257 non-significant but the effect was significantly mediated by semi-natural cover. These effects  
258 were consistent across all regions ([Fig. 4](#)). Together, the diversity of crop types available in  
259 the landscape and its interaction with semi-natural cover accounted for 10% of the explained  
260 variance in multitrophic diversity ([Fig. 2B](#)). The landscape-level effect of increasing crop  
261 diversity on multitrophic diversity ranged from negative in landscapes with low semi-natural  
262 cover to positive in landscapes with high semi-natural cover ([Fig. 3A](#)). This result is

263 consistent with the variability of effects observed across previous studies (30, 32, 34, 36, 37).  
264 To further explore this interaction, we used the same moving window modeling approach  
265 described above (see [section 5.7](#) in [SI 5](#) for details). This analysis confirmed that the  
266 landscape-level effect of increasing crop diversity on multitrophic diversity was positive in  
267 landscapes with more than 11% semi-natural cover (i.e. 50% of landscapes included in our  
268 study), non-significant in landscapes with 4 to 11% semi-natural cover (i.e. 34% of  
269 landscapes), and negative in landscapes with less than 4% semi-natural cover (i.e. 16% of  
270 landscapes; [Fig. S5.5.A](#) in [SI 5](#)).

271         The positive landscape-level effect of increasing crop diversity on multitrophic  
272 diversity observed in landscapes with more than 11% semi-natural cover supports the  
273 ‘landscape complementation’ hypothesis ([Hyp 1b](#) in [SI 1](#)). This finding is consistent with the  
274 fact that a diverse crop matrix provides a temporal continuity of food sources (38) while semi-  
275 natural patches provide stable resources, for example, for nesting or shelter (e.g. 37). Such  
276 complementation among multiple cover types has been described for several species (e.g. 38–  
277 40). Our study, based on multiple regions and multiple trophic groups, shows that the positive  
278 landscape-level effect of increasing crop diversity can be generalized to multitrophic diversity  
279 across many agricultural landscapes (50% of landscapes included in our study).

280         The negative landscape-level effect of increasing crop diversity on multitrophic  
281 diversity in landscapes with less than 4% semi-natural cover supports the ‘minimum total  
282 habitat area requirement’ hypothesis ([Hyp 1c](#) in [SI 1](#)). This finding is consistent with the fact  
283 that landscape simplification tends to filter out species with large body sizes (43), which also  
284 have high minimum total habitat area requirements (44), and may therefore require high  
285 amount of a single crop type. However, the whole range of taxa included in the present study,  
286 associated with a wide range of ecological traits, and therefore a wide range of minimum total  
287 habitat area requirements, showed a consistent response to crop diversity and the interaction

288 of crop diversity and semi-natural cover (Fig. 5). The ‘minimum total habitat area  
289 requirement’ hypothesis therefore seems unlikely to solely explain our results. Other  
290 hypotheses developed in the literature include the role of crop identity and management  
291 practices (e.g. 41). We considered the possibility that, at low levels of semi-natural cover,  
292 landscapes with higher crop diversity may have more intensive management practices, thus  
293 reducing multitrophic diversity (as suggested in 34). For example, in Armorique and PVDS,  
294 the increase in crop diversity was associated with a decrease in the cover of clover, a crop  
295 type associated with extensive management practices, and an increase in the cover of  
296 potatoes, a crop type associated with very intensive management practices (45). Reasons for  
297 the negative landscape-level effect of increasing crop diversity on multitrophic diversity in  
298 landscapes with low semi-natural cover deserve further attention. Future research is needed to  
299 identify conditions under which increasing crop diversity leads to a consistent net positive  
300 effect on multitrophic diversity, i.e. a positive effect of field-level (i.e. habitat specialization)  
301 plus landscape-level (i.e. landscape complementation) processes.

302

### 303 **Implications for agricultural policies**

304 Our study has important implications for large-scale policy schemes implemented across a  
305 wide range of contexts such as the European Common Agricultural Policy and its recent  
306 greening (27), the Canadian Agriculture Policy Frameworks (46), or the United States Farm  
307 Bill (47).

308 First, our results suggest that increasing crop heterogeneity may have a similar or  
309 greater benefit for multitrophic diversity to increasing semi-natural cover (Fig. 2B) or even  
310 decreasing field-level land use intensity (21; Table S5.12 in SI 5). Given current challenges to  
311 increase semi-natural cover and limit chemical use in agricultural landscapes (48), policies  
312 aiming at increasing crop heterogeneity may represent an effective and complementary way to

313 improve biodiversity conservation in agricultural landscapes. Policy measures favoring crop  
314 heterogeneity may be more easily implemented than policies to increase semi-natural cover or  
315 reduce chemical use (49). Associated with adequate economic incentives, they may also be  
316 more favorably perceived by farmers and thus lead to higher uptake than measures requiring  
317 farmers to take land out of production (48). Such measures may also contribute to the  
318 development of new frameworks that reward farmers for sustainable land stewardship (50).

319 We observed a consistent effect of crop heterogeneity on species diversity across  
320 seven taxa representing a wide range of ecological traits, functions and trophic levels (plants,  
321 bees, butterflies, hoverflies, carabids, spiders and birds; Fig. 5). We observed landscapes  
322 where six or even all seven taxa reached the threshold of 60% of the maximum species  
323 richness observed within a given region (Fig. 4). Our study therefore suggests that policies to  
324 increase crop heterogeneity would be an effective way to increase the diversity of all  
325 components of biodiversity simultaneously and restore multitrophic biodiversity in  
326 agricultural landscapes.

327 Finally, our results can contribute to the development of policies adapted to different  
328 landscape contexts. For instance, our results suggest that policy measures aimed at decreasing  
329 field sizes to below 6 ha may be particularly effective to promote multitrophic diversity in  
330 agricultural landscapes, especially in landscapes where semi-natural cover is below 8%. Our  
331 results also caution against a ‘blind’ increase of crop diversity. Measures aimed at increasing  
332 crop diversity may be effective to promote multitrophic diversity in landscapes where semi-  
333 natural cover exceeds 11%. However, they may have little effect or may even have negative  
334 effects in intensive agricultural landscapes with little semi-natural cover. Our study therefore  
335 highlights that measures promoting an increase in crop diversity are more likely to be  
336 effective in promoting multitrophic diversity across all agricultural landscapes if combined  
337 with measures promoting the restoration or maintenance of semi-natural cover.

338

## 339 **CONCLUSION**

340 Our study demonstrates the importance of crop heterogeneity for multitrophic diversity in  
341 agricultural landscapes: the effect of maintaining/increasing crop heterogeneity is likely to be  
342 as important as the effect of maintaining/increasing semi-natural cover. This finding suggests  
343 that field enlargement and crop specialization, especially the former, have been  
344 underestimated drivers of past and ongoing biodiversity declines. More importantly, our study  
345 shows that increasing crop heterogeneity represents a major potential lever to increase  
346 synergies between food production and biodiversity conservation.

347

## 348 **METHODS**

### 349 **1. Region, landscape and sampling site selection**

350 We selected eight agricultural regions (Armorique, Camargue, Coteaux de Gascogne  
351 and Plaine et Val de Sèvre in France, East Anglia in the United Kingdom, Goettingen in  
352 Germany, Lleida in Spain and Eastern Ontario in Canada; **Fig. S2.1 in SI 2**) belonging to six  
353 different ecoregions (51) and differing in topography, climate, field shapes, and agricultural  
354 cover types and products (e.g. rice, dairy, tree crops).

355 We used the best spatial data available within each region prior to field work to  
356 identify all 1 km × 1 km rural landscapes, i.e. those dominated by agricultural cover (>60%,  
357 including all crops and grassland managed for agricultural production). We then developed a  
358 protocol to select a combination of landscapes that maximized the gradients of crop  
359 compositional heterogeneity (crop diversity) and crop configurational heterogeneity (mean  
360 field size) while minimizing the correlation between them (52). Crop diversity may  
361 theoretically be constrained by the number and size of fields in landscapes with large fields.  
362 However, in our dataset, mean field size was smaller than 12 ha and was therefore not a

363 limiting factor for crop diversity within the 1 km x 1 km landscapes. We selected between 32  
364 and 93 landscapes within each region, totaling 435 landscapes across all regions.

365 We selected three sampling sites within each landscape, totaling 1305 sampling sites  
366 across all regions. The number of crop types sampled ranged from one to three per landscape.  
367 Where feasible, we located sampling sites in dominant agricultural cover types within each  
368 region (e.g. wheat fields and oilseed rape in Goettingen). When this was not feasible, we  
369 located sampling sites in agricultural cover types that were accessible within a given  
370 landscape (SI 3). The three sampling sites were at least 200 m from each other, at least 50 m  
371 from the border of the landscape, and at least 50 m from patches of non-agricultural cover  
372 types such as forests and urban areas.

373

## 374 **2. Multi-taxa sampling**

375 We selected seven taxa representing a wide range of ecological traits, functions and  
376 trophic levels which, combined into a multidiversity index (see below), represent a proxy for  
377 multitrophic diversity: plants, bees, butterflies, hoverflies, carabids, spiders and birds. All taxa  
378 were sampled using standardized sampling protocols across all regions, allowing us to test the  
379 consistency of effects across the eight regions (Section 3.1 in SI 3).

380 At each sampling site, we selected two parallel 50 m ‘transects’, one located at the  
381 field edge and the other inside the field 25 m away from the first transect (Fig. S3.1 and S3.2  
382 in SI 3). Birds were sampled using point-counts centered on the field-edge transect. Plants  
383 were surveyed along both transects. Butterflies were surveyed visually using timed walks  
384 along both transects. Bees and hoverflies were sampled using colored pan traps on poles  
385 erected at each end and in the center of all transects. Carabids and spiders were sampled using  
386 pitfall traps installed at each end of all transects. Captured arthropods were preserved in  
387 ethanol prior to identification. Multiple survey visits were conducted during the season when

388 relevant (SI 3). Each landscape was sampled during one year and sampling of landscapes was  
389 distributed across two years within each region, between 2011 and 2014 (see further details on  
390 the timing of our sampling in Table S3.1 in SI 3).

391 We identified more than 167,000 individuals from 2795 species (Table S3.2 in SI 3).  
392 For each taxon, we calculated species richness at the landscape level, i.e. across all three  
393 sampling sites and across all visits when multiple survey visits were conducted. The average  
394 species richness per landscape varied greatly among taxa, from 5.4 for butterflies to 44.9 for  
395 plants. Correlations in average species richness between pairs of taxa were weak ( $<0.41$ ), with  
396 an average correlation of 0.07 (Table S3.3 in SI 3).

397

### 398 **3. Multitrophic diversity index**

399 Our objective was to identify landscapes where the diversity of most taxa increases  
400 simultaneously. A first approach used in the literature consists of calculating the average,  
401 standardized diversity across taxa (53). However, this approach has limitations (see section  
402 3.3 in SI 3). Although very high/low values imply that all taxa exhibit high/low diversity,  
403 intermediate values are difficult to interpret as they may correspond to situations where (i)  
404 diversity values are intermediate for all taxa, or (ii) diversity values are high for some taxa  
405 and low for others, i.e. trade-offs among taxa.

406 To overcome this limitation, we used a threshold approach initially developed to  
407 aggregate multiple ecosystem functions (29, 54). For each taxon and each region, we  
408 identified the maximum species richness observed across all landscapes. We actually used the  
409 95<sup>th</sup> percentile as the maximum observed species richness (hereafter ‘SR max’) in order to  
410 minimize the effect of outliers. Next, we identified which landscapes attained a given  
411 threshold (x) of SR max. We then tallied the proportion of taxa that exceeded the given



412 threshold in order to produce a multidiversity index (Tx.landscape) for each landscape, based  
413 on the following formula:

$$414 \quad \text{Multidiversity (Tx.landscape)} = \frac{1}{n} \sum_{i=1}^{i=n} (\text{SR}_i > (x \times \text{SR}_{\text{max.region } j}))$$

415 where n is the number of taxa for which data were available in a given landscape (see  
416 details in [section 3.2 in SI 3](#)),  $\text{SR}_i$  is the number of species for taxon i, x is the minimum  
417 threshold to be reached and  $\text{SR}_{\text{max.region } j}$  is the maximum species richness for taxon i in the  
418 region the landscape considered belonged to. This multidiversity index ranges between 0 and  
419 1.

420 We calculated this multidiversity index for each threshold x between 20 and 90%  
421 (every 10%). For each threshold x, the multidiversity index was smoothed by calculating the  
422 average over the interval [x - 10%, x + 10%] (55; see details in [section 3.3 in SI 3](#)).  
423 Multidiversity indices calculated for different thresholds were strongly correlated. We chose  
424 to use the intermediate threshold T60.landscape because 1) intermediate thresholds have been  
425 shown to provide an effective measure of multitrophic diversity in agricultural landscapes  
426 (53) and 2) T60.landscape shows a distribution ranging from 0, i.e. none of the taxa reach  
427 60% of the regional maximum, to 100, i.e. all taxa reach 60% of the regional maximum (mean  
428 value for T60.landscape = 45.1). Nevertheless, we verified that our results were not sensitive  
429 to the threshold selected ([Fig. S5.2 in SI 5](#)). For simplicity, we hereafter refer to “landscape-  
430 level multitrophic diversity” rather than T60.landscape.

431

#### 432 **4. Crop compositional and configurational heterogeneity**

433 We used a standardized protocol across all regions to produce land cover maps  
434 allowing us to compare consistency of effects across the eight regions ([SI 4](#)). We conducted  
435 extensive ground-truthing surveys during the field seasons to map all fields, linear elements  
436 between adjacent fields, and non-agricultural covers. We built a common land cover

437 classification for the eight regions. Agricultural cover types included all crops, as well as  
438 temporary and permanent grassland managed for production purposes (SI 4). Linear elements  
439 between fields included hedgerows, grassy margins, ditches and tracks. Non-agricultural  
440 cover types included woodland (including woody linear elements), open land (e.g. extensive  
441 grassland, shrubland, grassy linear elements), wetland and built-up areas (including roads).  
442 We then used these standardized, detailed maps to calculate four explanatory variables for  
443 each landscape: crop diversity, mean field size, semi-natural cover and total length of semi-  
444 natural linear elements between fields.

445 We used the Shannon diversity of agricultural cover types (hereafter “crop diversity”;  
446 CD) as a measure of crop compositional heterogeneity. We used mean field size in hectares  
447 (MFS) as a measure of crop configurational heterogeneity. Neither CD nor MFS was  
448 correlated with local land use intensity (an index combining data on ploughing, fertilizer,  
449 herbicide and insecticide, see section 5.6.3 in SI 5) or the overall composition of the crop  
450 mosaic (section 5.5.1 in SI 5) across all regions. CD and MFS were moderately correlated  
451 with the type of crops sampled in some regions and MFS was moderately correlated with the  
452 proportion of grassland in the crop mosaic, but none of these correlations affected our  
453 conclusions (sections 5.5.1 and 5.5.2 in SI 5). We calculated the percentage of semi-natural  
454 cover types, i.e. woodland, open land and wetland (SNC), in each landscape. We also  
455 calculated the total length of linear semi-natural elements between fields, e.g. hedgerows,  
456 grassy margins (SNL; measured in meters). SNL and MFS were highly correlated in some  
457 regions (Table S5.6 in SI 5). As a result, we did not include SNL in the main analyses and  
458 only tested the relative effect of MFS and SNL using a subset of our dataset for which MFS  
459 and SNL were not strongly correlated (section 5.5.3 in SI 5).

460

## 461 **5. Data analysis**

462 We first tested the effect of crop heterogeneity on multitrophic diversity (Model 1).  
463 We fitted a linear mixed model with Restricted Maximum Likelihood using the landscape-  
464 level multidiversity index (T60.landscape) as the response variable. We included the number  
465 of crop types sampled per landscape (CropNb), crop diversity (CD), mean field size (MFS)  
466 and semi-natural cover (SNC) as explanatory variables (see alternative hypotheses on crop  
467 heterogeneity-biodiversity relationships in SI 1). We included both interaction effects and  
468 quadratic effects. Due to a positive skew in the distribution of mean field size, we used log  
469 mean field size in all analyses. To reflect the large-scale spatial and temporal structure of our  
470 dataset, we added sampling year (Year), nested within study region (Region), as a random  
471 effect. To reflect the spatial structure of our dataset within each region, we included the  
472 longitude and latitude of the center of each landscape (Lat, Lon) as covariates. We  
473 standardized all fixed effects to allow for a direct comparison of estimates.

474

475 *Model 1: lmer (T60.landscape ~ CD \* MFS \* SNC + CD<sup>2</sup> + MFS<sup>2</sup> + SNC<sup>2</sup> + CropNb + Lat + Lon + (1|*  
476 *Region/Year))*

477

478 To test whether the effects of crop diversity, mean field size and semi-natural cover on  
479 multitrophic diversity measured at the landscape level (T60.landscape) varied significantly  
480 among regions we added random effects for region on the slopes of crop diversity, mean field  
481 size, semi-natural cover as well as the interaction between crop diversity and semi-natural  
482 cover (model 2). We assumed that the effects of region on the intercept and slopes were  
483 uncorrelated. To test whether Region had a significant effect on the slope of either crop  
484 diversity, mean field size, semi-natural cover as well as the interaction between crop diversity  
485 and semi-natural cover, we used the function exactRLRT from package RLRsim.

486

487 *Model 2: lmer (T60.landscape ~ CD \* MFS \* SNC + CD<sup>2</sup> + MFS<sup>2</sup> + SNC<sup>2</sup> + CropNb + Lat + Lon +*  
488 *(1|Region/Year) + (0+CD|Region)) + (0+MFS|Region) + (0+SNC|Region) + (0+CD:SNC|Region))*

489

490 We then tested the effects of crop heterogeneity on the species richness of taxonomic  
491 groups (Model 3). To do this, we fitted a similar model, using the landscape-level species  
492 richness of taxonomic groups (SR) standardized within each taxon and region as the response  
493 variable. To reflect that species pools vary between taxa, we added Taxon as a random effect.

494

495 *Model 3: lmer(SR ~ CD\*MFS\*SNC + CD<sup>2</sup> + MFS<sup>2</sup> + SNC<sup>2</sup> + CropNb + Lat + Lon + (1|Region/Year) +*  
496 *(1|Taxon))*

497

498 To test whether the effects of crop diversity, mean field size and semi-natural cover on  
499 the species richness of taxonomic groups varied significantly among taxa we added random  
500 effects for Taxon on the slopes of crop diversity, mean field size, semi-natural cover as well  
501 as the interaction between crop diversity and semi-natural cover (model 4). We assumed that  
502 the effects of Taxon on the intercept and slopes were uncorrelated. To test whether Taxon had  
503 a significant effect on the slope of either crop diversity, mean field size, semi-natural cover or  
504 the interaction between crop diversity and semi-natural cover, we used the function  
505 exactRLRT from package RLRsim.

506

507 *Model 4: lmer(SR ~ CD \* MFS \* SNC + CD<sup>2</sup> + MFS<sup>2</sup> + SNC<sup>2</sup> + CropNb + Lat + Lon + (1|Taxon) +*  
508 *(1|Region/Year) + (0+CD|Taxon)) + (0+MFS|Taxon) + (0+SNC|Taxon) + (0+CD:SNC|Taxon))*

509

510 We fitted all models with the R lme4 package using LMER (56), we removed outliers  
511 using function romr.fnc from package LMERConvenienceFunctions (57) and we ran  
512 diagnostic tools to verify that residuals were independently and normally distributed, and  
513 showed no spatial autocorrelation. For each model, a multimodel inference procedure was  
514 applied using the R MuMIn package (58). This method allowed us to perform model selection  
515 by creating a set of models with all possible combinations of the initial variables and sorting

516 them according to the Akaike Information Criterion (AIC) fitted with Maximum Likelihood  
517 (59). We selected all models with  $\Delta AIC < 2$  and used the model averaging approach using  
518 LMER to estimate parameters and associated p-values, using the function model.avg. We ran  
519 all analyses using the software R 3.4.0 (60).

520 We ran additional analyses to check that the composition of the crop mosaic, the  
521 proportion of grassland in the crop mosaic, and the amount of semi-natural vegetation  
522 occurring between fields did not affect our conclusions (section 5.5 in SI 5). We also ran  
523 complementary analyses using field-level multidiversity (T60.field) as the response variable -  
524 instead of the landscape-level multidiversity index (T60.landscape) - to check that our results  
525 hold true at the field level, in particular within a subset of cereal fields, and that the type of  
526 crop sampled or the level of land-use intensity within sampled fields did not affect our  
527 conclusions (section 5.6 in SI 5). Finally, we used a moving window analysis to identify  
528 potential discontinuities in multitrophic diversity response to crop diversity and mean field  
529 size along the gradient of semi-natural cover (section 5.7 in SI 5).

530

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554

#### 555 **Author contribution**

556 CS and NG wrote the first draft of the manuscript; BP, FB, TT, VB, GS, AO, LB, JLM and LF  
557 designed the FarmLand project; CS, ABB, CB, RC, AH, LH, PM, AA, JG, DG, G Bota, FC,  
558 AGT, RG, SH, JR, XOSS, IR, JB, JAB, AR, MAM-G, JM and GS contributed data; CS, NG,  
559 ABB, CB, RC, AH, LH, PM and AA analyzed data; all co-authors provided feedback on the  
560 manuscript.

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562

563

#### 564 **References**

- 565 1. Tilman D, et al. (2001) Forecasting agriculturally driven global environmental change.  
566 *Science* 292(5515):281–284.
- 567 2. Foley JA, et al. (2005) Global consequences of land use. *Science* 309(5734):570–574.
- 568 3. Robinson RA, Sutherland WJ (2002) Post-war changes in arable farming and  
569 biodiversity in Great Britain. *J Appl Ecol* 39(1):157–176.
- 570 4. White EV, Roy DP (2015) A contemporary decennial examination of changing  
571 agricultural field sizes using Landsat time series data. *Geo Geogr Environ* 2(1):33–54.
- 572 5. Barr CJ, Gillespie MK (2000) Estimating hedgerow length and pattern characteristics in  
573 Great Britain using Countryside Survey data. *J Environ Manage* 60(1):23–32.
- 574 6. Aguilar J, et al. (2015) Crop Species Diversity Changes in the United States: 1978–2012.  
575 *PLoS ONE* 10(8). doi:10.1371/journal.pone.0136580.
- 576 7. Newbold T, et al. (2015) Global effects of land use on local terrestrial biodiversity.  
577 *Nature* 520(7545):45–50.
- 578 8. Maxwell SL, Fuller RA, Brooks TM, Watson JEM (2016) Biodiversity: The ravages of  
579 guns, nets and bulldozers. *Nature* 536(7615):143–145.
- 580 9. Benton TG, Vickery JA, Wilson JD (2003) Farmland biodiversity: is habitat  
581 heterogeneity the key? *Trends Ecol Evol* 18(4):182–188.
- 582 10. Batáry P, Fischer J, Báldi A, Crist TO, Tscharrntke T (2011) Does habitat heterogeneity  
583 increase farmland biodiversity? *Front Ecol Environ* 9(3):152–153.
- 584 11. Miyashita T, Chishiki Y, Takagi SR (2012) Landscape heterogeneity at multiple spatial  
585 scales enhances spider species richness in an agricultural landscape. *Popul Ecol*  
586 54(4):573–581.
- 587 12. Perović D, et al. (2015) Configurational landscape heterogeneity shapes functional  
588 community composition of grassland butterflies. *J Appl Ecol* 52(2):505–513.

- 589 13. Batáry P, Dicks LV, Kleijn D, Sutherland WJ (2015) The role of agri-environment  
590 schemes in conservation and environmental management. *Conserv Biol* 29(4):1006–  
591 1016.
- 592 14. Lüscher G, et al. (2016) Farmland biodiversity and agricultural management on 237  
593 farms in 13 European and two African regions. *Ecology* 97(6):1625–1625.
- 594 15. Henderson IG, Vickery JA, Carter N (2004) The use of winter bird crops by farmland  
595 birds in lowland England. *Biol Conserv* 118(1):21–32.
- 596 16. Holzschuh A, Dormann CF, Tschardt T, Steffan-Dewenter I (2013) Mass-flowering  
597 crops enhance wild bee abundance. *Oecologia* 172(2):477–484.
- 598 17. Raymond L, et al. (2014) Immature hoverflies overwinter in cultivated fields and may  
599 significantly control aphid populations in autumn. *Agric Ecosyst Environ* 185:99–105.
- 600 18. Fahrig L, et al. (2011) Functional landscape heterogeneity and animal biodiversity in  
601 agricultural landscapes. *Ecol Lett* 14(2):101–112.
- 602 19. Weibull A-C, Östman Ö, Granqvist Å (2003) Species richness in agroecosystems: the  
603 effect of landscape, habitat and farm management. *Biodivers Conserv* 12(7):1335–1355.
- 604 20. Dunning JB, Danielson BJ, Pulliam HR (1992) Ecological processes that affect  
605 populations in complex landscapes. *Oikos* 65(1):169–175.
- 606 21. Batáry P, et al. (2017) The former Iron Curtain still drives biodiversity-profit trade-offs  
607 in German agriculture. *Nat Ecol Evol* 1(9):1279–1284.
- 608 22. Tschardt T, et al. (2012) Landscape moderation of biodiversity patterns and processes  
609 - eight hypotheses. *Biol Rev* 87(3):661–685.
- 610 23. Burel F, Butet A, Delettre YR, de la Pena NM (2004) Differential response of selected  
611 taxa to landscape context and agricultural intensification. *Landsc Urban Plan* 67(1–  
612 4):195–204.
- 613 24. Dormann CF, et al. (2007) Effects of landscape structure and land-use intensity on  
614 similarity of plant and animal communities. *Glob Ecol Biogeogr* 16(6):774–787.
- 615 25. Ponjoan A, Bota G, Mañosa S (2012) Ranging behaviour of little bustard males, *Tetrax*  
616 *tetrax*, in the lekking grounds. *Behav Processes* 91(1):35–40.
- 617 26. Gaba S, Chauvel B, Dessaint F, Bretagnolle V, Petit S (2010) Weed species richness in  
618 winter wheat increases with landscape heterogeneity. *Agric Ecosyst Environ* 138(3–  
619 4):318–323.
- 620 27. Pe'er G, et al. (2014) EU agricultural reform fails on biodiversity. *Science*  
621 344(6188):1090–1092.
- 622 28. Díaz S, et al. (2015) The IPBES Conceptual Framework — connecting nature and  
623 people. *Curr Opin Environ Sustain* 14:1–16.

- 624 29. Allan E, et al. (2014) Interannual variation in land-use intensity enhances grassland  
625 multidiversity. *Proc Natl Acad Sci U S A* 111(1):308–313.
- 626 30. Fahrig L, et al. (2015) Farmlands with smaller crop fields have higher within-field  
627 biodiversity. *Agric Ecosyst Environ* 200:219–234.
- 628 31. Collins SJ, Fahrig L (2017) Responses of anurans to composition and configuration of  
629 agricultural landscapes. *Agric Ecosyst Environ* 239(Supplement C):399–409.
- 630 32. Monck-Whipp L, Martin AE, Francis CM, Fahrig L (2018) Farmland heterogeneity  
631 benefits bats in agricultural landscapes. *Agric Ecosyst Environ* 253(Supplement C):131–  
632 139.
- 633 33. Šálek M, et al. (2018) Bringing diversity back to agriculture: Smaller fields and non-crop  
634 elements enhance biodiversity in intensively managed arable farmlands. *Ecol Indic*  
635 90:65–73.
- 636 34. Hass AL, et al. (2018) Landscape configurational heterogeneity by small-scale  
637 agriculture, not crop diversity, maintains pollinators and plant reproduction in western  
638 Europe. *Proc R Soc B* 285(1872):20172242.
- 639 35. Berdugo M, et al. Aridity preferences alter the relative importance of abiotic and biotic  
640 drivers on plant species abundance in global drylands. *J Ecol* 0(0). doi:10.1111/1365-  
641 2745.13006.
- 642 36. Josefsson J, Berg Å, Hiron M, Pärt T, Eggers S (2017) Sensitivity of the farmland bird  
643 community to crop diversification in Sweden: does the CAP fit? *J Appl Ecol* 54(2):518–  
644 526.
- 645 37. Olimpi EM, Philpott SM (2018) Agroecological farming practices promote bats. *Agric*  
646 *Ecosyst Environ* 265:282–291.
- 647 38. Schellhorn NA, Gagic V, Bommarco R (2015) Time will tell: resource continuity  
648 bolsters ecosystem services. *Trends Ecol Evol* 30(9):524–530.
- 649 39. Sirami C, Brotons L, Martin J (2011) Woodlarks *Lullula arborea* and landscape  
650 heterogeneity created by land abandonment. *Bird Study* 58(1):99–106.
- 651 40. Pope SE, Fahrig L, Merriam NG (2000) Landscape complementation and  
652 metapopulation effects on leopard frog populations. *Ecology* 81(9):2498–2508.
- 653 41. Mueller T, Selva N, Pugacewicz E, Prins E (2009) Scale-sensitive landscape  
654 complementation determines habitat suitability for a territorial generalist. *Ecography*  
655 32(2):345–353.
- 656 42. Marrec R, et al. (2015) Crop succession and habitat preferences drive the distribution  
657 and abundance of carabid beetles in an agricultural landscape. *Agric Ecosyst Environ*  
658 199:282–289.
- 659 43. Gámez-Virués S, et al. (2015) Landscape simplification filters species traits and drives  
660 biotic homogenization. *Nat Commun* 6:8568.



- 661 44. Baguette M, Stevens V (2013) Predicting minimum area requirements of butterflies  
662 using life-history traits. *J Insect Conserv* 17(4):645–652.
- 663 45. Agreste (2013) Les indicateurs de fréquence de traitement (IFT) en 2011. *Doss* 18.
- 664 46. Deaton BJ, Boxall P (2017) Canadian Agricultural Policy in the Twenty-First Century:  
665 Looking Back and Going Forward. *Can J Agric Econ Can Daposagroeconomie*  
666 65(4):519–522.
- 667 47. Reimer A (2015) Ecological modernization in U.S. agri-environmental programs: Trends  
668 in the 2014 Farm Bill. *Land Use Policy* 47(Supplement C):209–217.
- 669 48. Pe’er G, et al. (2017) Adding Some Green to the Greening: Improving the EU’s  
670 Ecological Focus Areas for Biodiversity and Farmers. *Conserv Lett* 10(5):517–530.
- 671 49. Rodríguez C, Wiegand K (2009) Evaluating the trade-off between machinery efficiency  
672 and loss of biodiversity-friendly habitats in arable landscapes: The role of field size.  
673 *Agric Ecosyst Environ* 129(4):361–366.
- 674 50. Mathevet R, Bousquet F, Raymond CM (2018) The concept of stewardship in  
675 sustainability science and conservation biology. *Biol Conserv* 217:363–370.
- 676 51. Olson DM, et al. (2001) Terrestrial Ecoregions of the World: A New Map of Life on  
677 Earth A new global map of terrestrial ecoregions provides an innovative tool for  
678 conserving biodiversity. *BioScience* 51(11):933–938.
- 679 52. Pasher J, et al. (2013) Optimizing landscape selection for estimating relative effects of  
680 landscape variables on ecological responses. *Landsc Ecol* 28(3):371–383.
- 681 53. Byrnes JEK, et al. (2014) Investigating the relationship between biodiversity and  
682 ecosystem multifunctionality: challenges and solutions. *Methods Ecol Evol* 5(2):111–  
683 124.
- 684 54. Zavaleta ES, Pasari JR, Hulvey KB, Tilman GD (2010) Sustaining multiple ecosystem  
685 functions in grassland communities requires higher biodiversity. *Proc Natl Acad Sci U S*  
686 *A* 107(4):1443–1446.
- 687 55. Le Bagousse-Pinguet Y, et al. (2019) Phylogenetic, functional, and taxonomic richness  
688 have both positive and negative effects on ecosystem multifunctionality. *Proc Natl Acad*  
689 *Sci*:201815727.
- 690 56. Bates D, Mächler M, Bolker BM, Walker SC (2015) Fitting linear mixed-effects models  
691 using lme4. *J Stat Softw* 67(1):1–48.
- 692 57. Tremblay A, Ransijn J (2015) *LMERConvenienceFunctions: Model Selection and Post-*  
693 *hoc Analysis for (G)LMER Models*.
- 694 58. Barton K (2009) MuMIn : multi-model inference, R package version 0.12.0. *Httppr-Forg-*  
695 *Proj*. Available at: <https://ci.nii.ac.jp/naid/10030574914/> [Accessed August 1, 2018].

- 696 59. Zuur A, Ieno EN, Walker N, Saveliev AA, Smith GM (2009) *Mixed effects models and*  
697 *extensions in ecology with R* (Springer New York) Available at:  
698 <http://www.springer.com/in/book/9780387874579> [Accessed June 24, 2017].
- 699 60. R Core Team (2016) *R: A language and environment for statistical computing* (R  
700 Foundation for Statistical Computing. ISBN 3-900051-07-0, URL [http://www.R-](http://www.R-project.org)  
701 [project.org](http://www.R-project.org)., Vienna, Austria).
- 702 61. Grace JB, Bollen KA (2005) Interpreting the Results from Multiple Regression and  
703 Structural Equation Models. *Bull Ecol Soc Am* 86(4):283–295.
- 704
- 705

706 **Figure legends**

707

708 **Figure 1.** A) Traditional representations of agricultural landscapes have focused on the  
709 amount of semi-natural covers and semi-natural vegetation between fields, often considering  
710 the farmed part of the landscape as a homogeneous matrix. These representations are  
711 associated with the hypothesis that increasing the amount of semi-natural covers and semi-  
712 natural vegetation between fields benefits biodiversity. B) Novel representations of  
713 agricultural landscapes consider the heterogeneity of the crop mosaic. These representations  
714 are associated with new hypotheses: increasing crop heterogeneity by increasing crop  
715 diversity and/or decreasing mean field size, while maintaining semi-natural cover and semi-  
716 natural vegetation between fields constant, benefits biodiversity (large squares represent  
717 landscapes; adapted from 18).

718

719 **Figure 2.** Response of multitrophic diversity to the diversity of crop types available within the  
720 landscape (CD), the number of crops sampled (Crop Nb), mean field size (MFS), semi-natural  
721 cover (SNC), and interaction terms (CD:SNC, MFS:SNC, see further details in *Methods*),  
722 based on data collected in 435 landscapes located in eight agricultural regions. Covariates  
723 (Lon, Lat) were excluded from the figure for simplicity. A) Importance of each variable in the  
724 model averaging approach (model 1), estimated as the proportion of submodels where the  
725 variable was selected (see details in *SI 5*). B) The relative effect of each variable corresponds  
726 to the ratio between its parameter estimate and the sum of all parameter estimates (i.e. the %  
727 of variance explained, as explained in 60). Parameter estimates and confidence intervals,  
728 based on a model averaging approach applied to model 1 (*Methods*). ° p<0.1; \* p<0.05; \*\*  
729 p<0.01; \*\*\* p<0.001. Variables are grouped in three components: orange = crop  
730 heterogeneity (MFS, MFS<sup>2</sup>, CD, CD<sup>2</sup>, MFS:CD, Crop Nb), green = semi-natural cover (SNC,  
731 SNC<sup>2</sup>), blue = interactive effects between crop heterogeneity and semi-natural cover  
732 (CD:SNC, MFS:SNC, CD:MFS:SNC). The % of variance explained by CD is too small to be  
733 visible.

734

735 **Figure 3.** Effect of the diversity of crop types available within the landscape (CD), mean field  
736 size (MFS), semi-natural cover (SNC), and their interaction terms on landscape-level  
737 multitrophic diversity (see further details in *Methods*), based on data collected in 435  
738 landscapes located in eight agricultural regions. A) Interactive effects of crop diversity and  
739 semi-natural cover on multitrophic diversity. B) Interactive effects of mean field size and  
740 semi-natural cover on multitrophic diversity. The direction of the mean field size axis is  
741 reversed to improve readability. The parameter estimates of all other variables were fixed to  
742 their mean values, i.e. zero, as all predictors were scaled. Black dots and surfaces correspond  
743 to values of multitrophic diversity predicted by the model averaging approach applied to  
744 model 1 (*Methods*). The color gradient corresponds to multitrophic diversity values, ranging  
745 from low values (blue) to high values (red). Grey dots show the overall gradients of crop  
746 diversity, mean field size and semi-natural cover across the 435 landscapes located in eight  
747 regions.

748

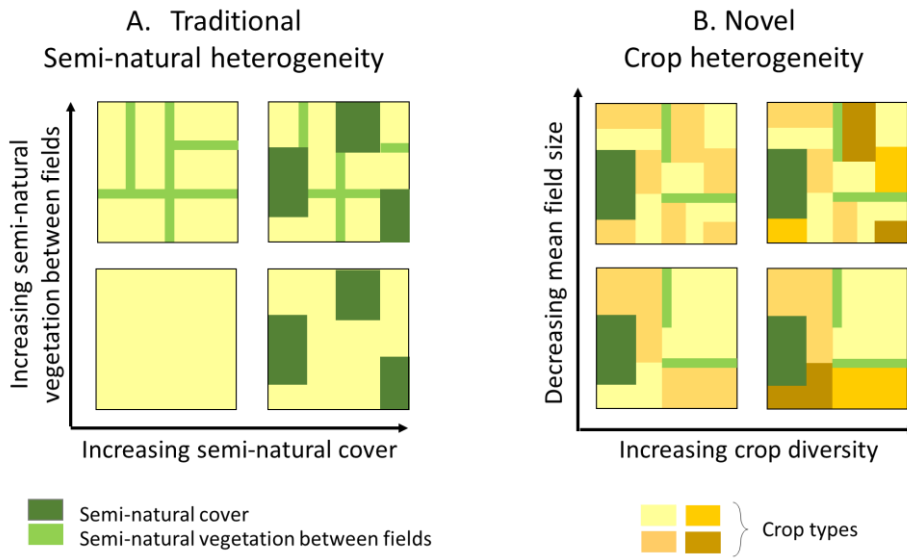
749 **Figure 4.** Effects of the diversity of crop types available in the landscape (CD), mean field  
750 size (MFS), semi-natural cover (SNC) and the interaction between crop diversity and semi-  
751 natural cover (CD:SNC) on multitrophic diversity in different regions (see further details in  
752 *Methods*). Slopes are based on the outputs of model 2 including a random effect of region on  
753 these four slopes (n=435 landscapes). Colors indicate the region.

754

755 **Figure 5.** Effects of the diversity of crop types available in the landscape (CD), mean field  
756 size (log MFS), semi-natural cover (SNC) and the interaction between the diversity of crop  
757 types available in the landscape and semi-natural cover (CD:SNC) on the landscape-level  
758 species richness of taxonomic groups (see further details in *Methods*). Slopes are based on the  
759 outputs of model 10 including a random effect of taxon on these four slopes (n=435  
760 landscapes). Colors indicate the taxon.  
761

762 **Figure 1.**  
763

### Representations of farmland heterogeneity

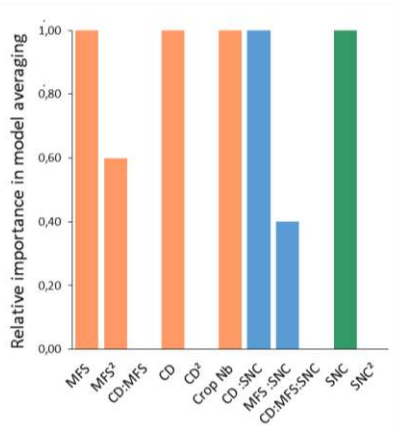


764

765 **Figure 2**

766

767 **A**



768

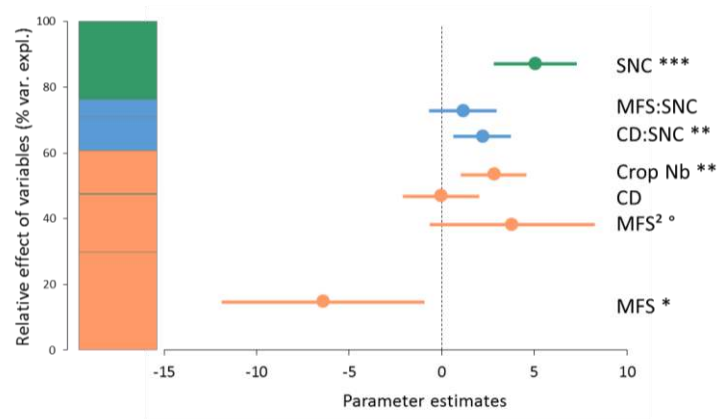
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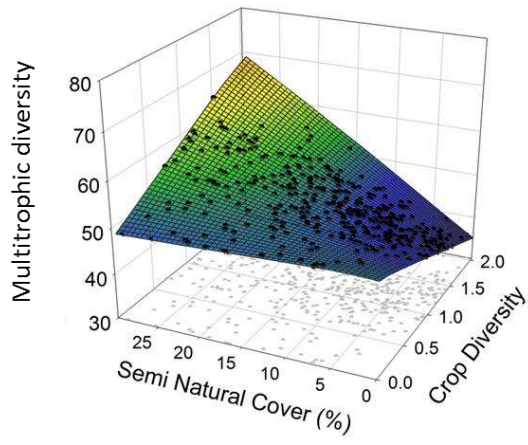
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**B**

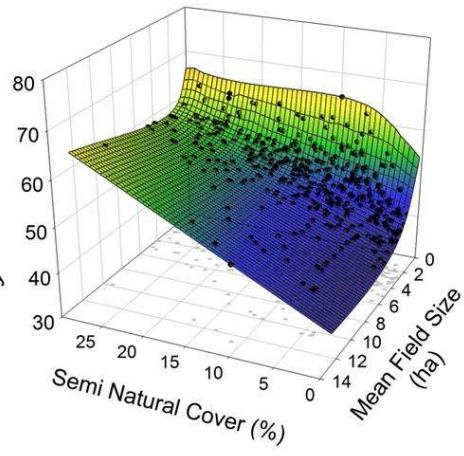


773 **Figure 3.**

774 A



B

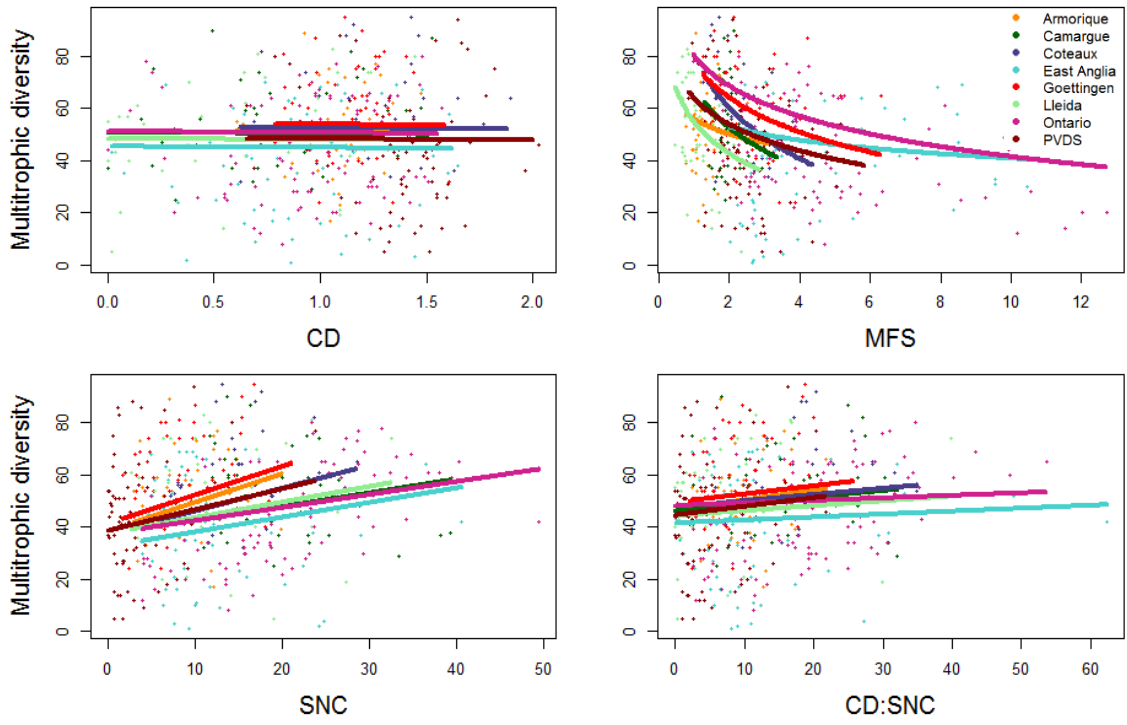


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776

777 **Figure 4.**

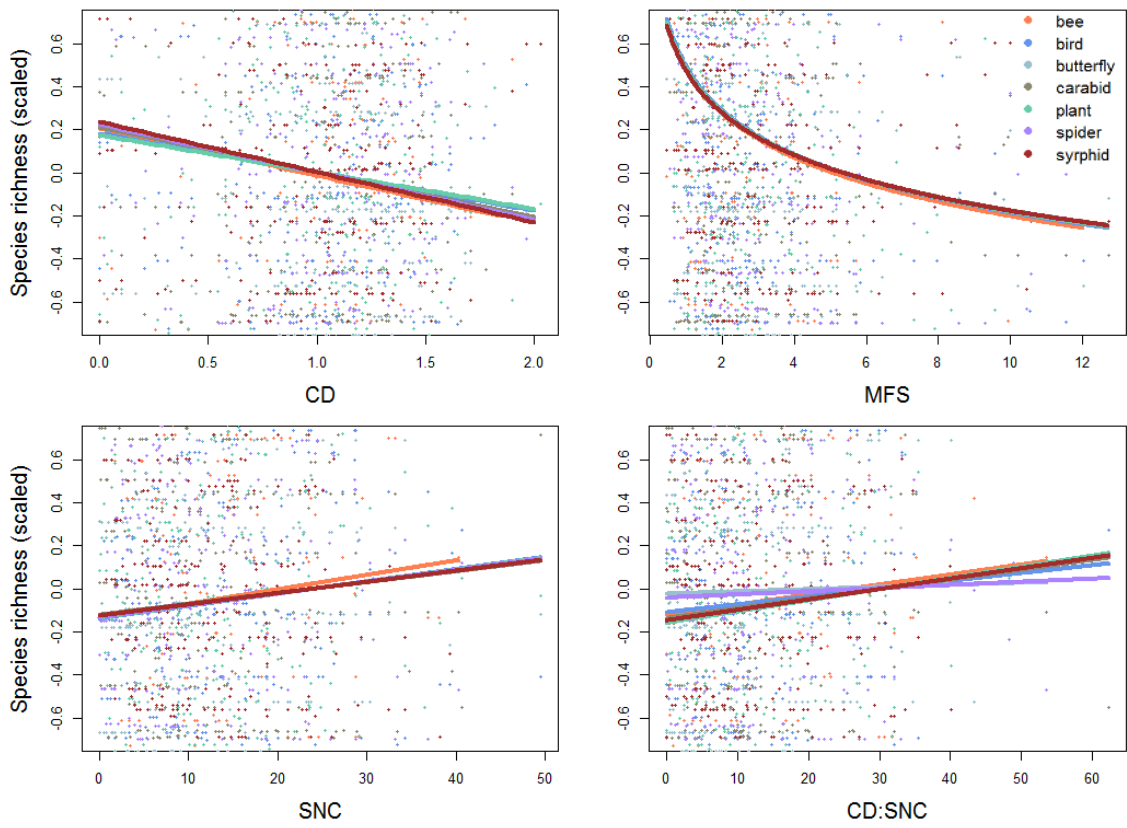
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779  
780



781 **Figure 5.**



782  
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787 **Supporting Information**

788

789 SI 1 – Hypotheses on the effects of crop heterogeneity on biodiversity

790 SI 2 – Region and landscape selection

791 SI 3 – Multitrophic diversity sampling

792 SI 4 – Land cover mapping and landscape metrics

793 SI 5 – Complementary analyses

794

795 **SI 1 – Hypotheses on the effects of crop heterogeneity on biodiversity**

796

797

798

**1.1. Crop compositional heterogeneity (increasing crop diversity)**

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800

**Hyp 1a.** Biodiversity **increases** with increasing crop diversity if different crop types can serve as habitat for different specialized species (single habitat specialization; Andreasen et al. 1991; Weibull et al. 2003). In that case, sampling more crop types will result in observing higher landscape-level biodiversity.

804

805

**Hyp 1b.** Biodiversity **increases** with increasing crop diversity if different crop types provide different resources required for single species (landscape complementation; Dunning et al. 1992), or if specialist species spillover from other crop types in the landscape into the fields sampled (Duelli 1997, Schneider et al. 2016). In that case, for a given number of crop types sampled, landscapes with higher crop diversity will result in observing higher landscape-level biodiversity.

810

811

**Hyp 1c.** Biodiversity **decreases** with crop diversity if most species have high minimum total habitat area requirements, i.e. require large amounts of a single crop type. An increase in the number of crop types available in the landscape results in a decrease in the total area of each crop type available in the landscape, which could hypothetically result in insufficient resources for species associated with individual crop types (Fahrig et al. 2011; Tscharntke et al. 2012).

816

817

**Hyp 1d.** Biodiversity shows a **peaked relationship** with crop diversity available in the landscape (Allouche et al. 2012) if there is an initial increase in biodiversity with increasing crop diversity for reasons explained in Hyp 1a-1b, but at higher levels of crop diversity, each crop type has a lower spatial cover and biodiversity decreases for reasons explained in Hyp 1c.

821

822

823

**1.2. Crop configurational heterogeneity (decreasing mean field size)**

824

825

**Hyp 2a.** Biodiversity **increases** with decreasing mean field size if landscapes with smaller fields provide easier access to multiple fields for species that require resources occurring in different crop types (landscape complementation).

828

829

**Hyp 2b.** Biodiversity **increases** with decreasing mean field size if landscapes with smaller fields also have higher density of crop edges. This could increase biodiversity measured in sampled crop fields by increasing spillover from adjacent fields or from adjacent semi-natural vegetation occurring between fields.

833

834

**Hyp 2c.** Biodiversity **decreases** with decreasing mean field size if most species show negative edge effects and/or if most species have minimum patch size requirements (separate from their total habitat area requirements, see Hyp1c).

837

838

**Hyp 2d.** Biodiversity shows a **peaked relationship** with decreasing mean field size if there is an initial increase in biodiversity for reasons explained in Hyp 2a-2b and then biodiversity decreases when mean field size reaches minimum patch size requirements for most species (Hyp 2c).

840

841

842

843

**1.3. Interactions between crop compositional and configurational heterogeneity**

844

845 **Hyp 3a.** The positive effect of crop diversity on biodiversity is **stronger** when mean field size  
846 decreases (and vice-versa) if most species require multiple land cover types easily accessible  
847 (landscape complementation). This is because increasing crop diversity increases the chance that all  
848 required crop types are available, and decreasing field sizes increases accessibility among the  
849 required crop types.

850  
851 **Hyp 3b.** The positive effect of crop diversity on biodiversity is **weaker** when mean field size is low if  
852 most species require landscape complementation and have minimum patch size requirements.  
853 Similarly, the positive effect of decreasing mean field size on biodiversity is **weaker** when crop  
854 diversity is high if the presence of a distinct crop type in the adjacent field results in a negative edge  
855 effect for most species within the sampled field.

856  
857 **Hyp 3c.** The positive effect of crop diversity on biodiversity is **independent** of mean field size if most  
858 species are highly mobile and can access multiple fields regardless of mean field size. The positive  
859 effect of decreasing mean field size on biodiversity is **independent** of crop diversity if most species in  
860 landscapes with low mean field size primarily benefit from an easier access to semi-natural cover, in  
861 particular to semi-natural linear elements, rather than to multiple fields.

862  
863

#### 864 **1.4. Interactions between crop heterogeneity and semi-natural cover**

865

866 **Hyp 4a.** The positive effect of **crop diversity** on biodiversity is **stronger** when semi-natural cover  
867 (SNC) increases if most species require complementary resources found in semi-natural cover types  
868 and several crop types (e.g. species require SNC + crop A + crop B).

869

870 **Hyp 4b.** The positive effect of decreasing **mean field size** on biodiversity is **stronger** when semi-  
871 natural cover (SNC) increases if most species in landscapes with low mean field size primarily benefit  
872 from an easier access to semi-natural cover, in particular to semi-natural linear elements, rather than  
873 an easier access to multiple fields.

874

875 **Hyp 4c.** The positive effects of crop heterogeneity on biodiversity is stronger in landscapes with  
876 **intermediate amounts of semi-natural cover** than in landscapes with very low or very high semi-  
877 natural cover (Tscharrntke et al. 2012). In landscapes with no or very low semi-natural cover, species  
878 pool may be small and species may be well adapted to intensive agriculture, and biodiversity may  
879 therefore remain unaffected by crop heterogeneity levels. In landscapes with high semi-natural  
880 cover, biodiversity levels may be high everywhere due to widespread spill-over effects, and may  
881 remain unaffected by crop heterogeneity levels.

882

883

#### 884 **References**

885

886 Allouche O, et al. (2012) Area–heterogeneity tradeoff and the diversity of ecological communities. *Proc Natl Acad*  
887 *Sci* 109(43):17495–17500.

888 Andreassen C, et al. (1991) Soil properties affecting the distribution of 37 weed species in Danish fields. *Weed Res*  
889 31(4):181–187.

890 Dunning JB, et al. (1992) Ecological processes that affect populations in complex landscapes. *Oikos* 65(1):169–  
891 175.

892 Fahrig L, et al. (2011) Functional landscape heterogeneity and animal biodiversity in agricultural landscapes. *Ecol*  
893 *Lett* 14(2):101–112.

894 Schneider G, Krauss J, Boetzel FA, Fritze M-A, Steffan-Dewenter I (2016) Spillover from adjacent crop and forest  
895 habitats shapes carabid beetle assemblages in fragmented semi-natural grasslands. *Oecologia*  
896 182(4):1141–1150.

- 897 Tschardtke T, et al. (2012) Landscape moderation of biodiversity patterns and processes - eight hypotheses. *Biol*  
898 *Rev* 87(3):661–685.
- 899 Weibull A-C, et al. (2003) Species richness in agroecosystems: the effect of landscape, habitat and farm  
900 management. *Biodivers Conserv* 12(7):1335–1355.

901 **SI 2 – Region and landscape selection**

902

903

904 **2.1. Region selection**

905

906 We selected eight agricultural regions (Fig. S2.1) that belong to six different ecoregions (Olson et al;  
907 2001)(51) : Eastern Great Lakes lowland forests (Eastern Ontario in Canada), Celtic broadleaf forests  
908 and English lowland beech forests (East Anglia in United Kingdom), Atlantic mixed forests  
909 (Armorique, Plaine et Val de Sèvre in France), Western European broadleaf forests (Goettingen in  
910 Germany, Coteaux de Gascogne in France), Iberian sclerophyllous and semi-deciduous forests (Lleida  
911 in Spain) and Northeastern Spain & Southern France Mediterranean forests (Camargue in France).  
912 Topography varied from flat (e.g. Camargue, Eastern Ontario) to intermediate (e.g. Goettingen,  
913 Lleida), to hilly (e.g. Coteaux de Gascogne). Climate varied from dry (e.g. Lleida) to humid (e.g. East  
914 Anglia). Complexity in crop field shapes varied from rectilinear (e.g. Camargue, Eastern Ontario) to  
915 intermediate complexity (e.g. Coteaux de Gascogne, Armorique) to complex field shapes (e.g. Lleida).  
916 Specific agricultural products were found in some regions, e.g. dairy (Armorique), olives (Lleida) or  
917 rice (Camargue). Diversity of agricultural cover types varied from low (e.g. Camargue, Lleida) to high  
918 (e.g. Coteaux de Gascogne, Plaine et Val de Sèvre). Mean field size varied from 1.2 ha in Lleida and  
919 1.4 ha in Armorique to 4.4 ha in Eastern Ontario and 4.7 ha in East Anglia.

920



921

922 **Figure S2.1.** Locations of the eight study regions in Europe and North America.

923

924

925 **2.2. Landscape selection**

926

927 The purpose of the landscape selection protocol was to select in each region a set of landscapes  
928 in a pseudo-experimental design (also called a "mensurative experiment") which aimed at selecting  
929 agricultural landscapes (between 60 and 100% of agricultural cover) along two independent  
930 gradients of crop compositional and configurational heterogeneity. The general protocol is detailed  
931 in Pasher et al. (2013).

932

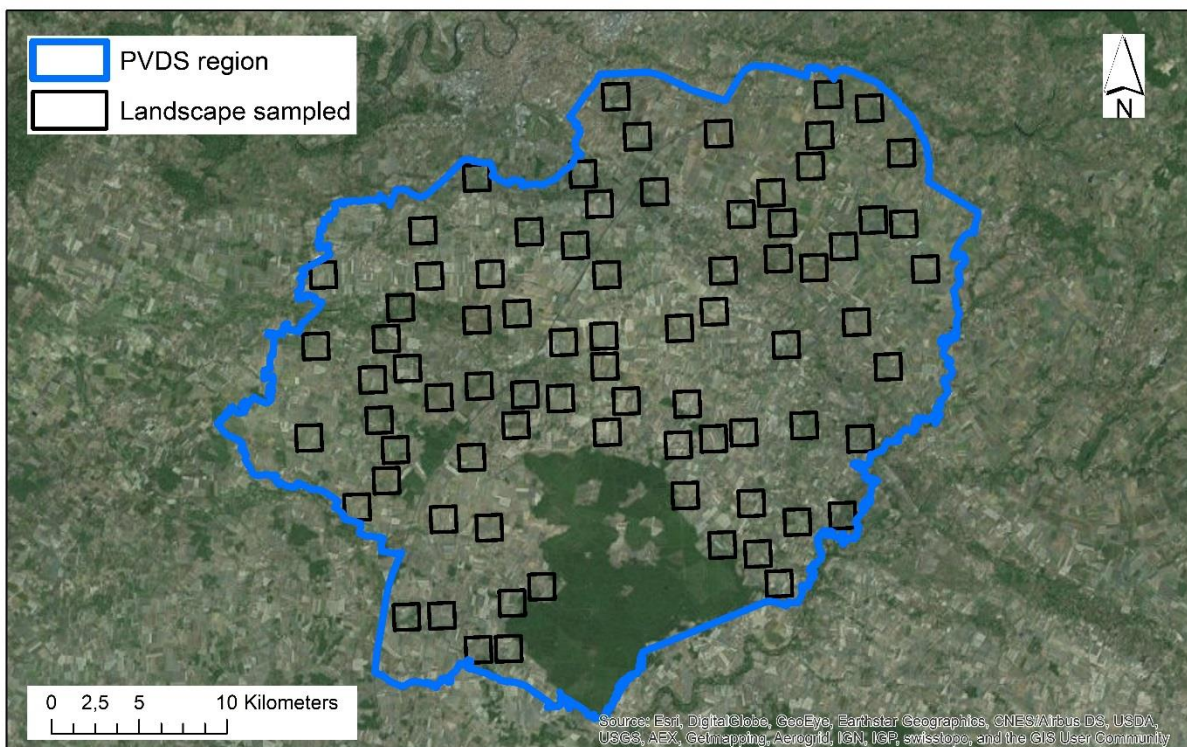
933 We used the highest resolution and most recent remotely sensed data or the best land cover  
934 map available within each region. We delineated all fields (contiguous production cover), even when  
935 adjacent fields contain the same agricultural cover type (as they may belong to different farmers or  
may be managed differently). We attributed each field to one of the following 34 agricultural cover

936 types: cereal, fallow, alfalfa, clover, ryegrass, grassland, rice, corn, sunflower, sorghum, millet, moha,  
937 oilseed rape, mustard, pea, bean, soybean, linseed, orchard, almond, olive, vineyard, mixed  
938 vegetables, sugar beet, asparagus, carrot, onion, parsnip, potato, tomato, melon, strawberry,  
939 raspberry, wild bird cover (i.e. a spring sown crop left unharvested over winter to provide food for  
940 farmland birds). We also delineated patches of non-agricultural cover (woodland, open land, wetland  
941 and built-area).

942 We then calculated crop compositional heterogeneity (Shannon diversity index of the crop  
943 mosaic) and crop configurational heterogeneity (mean size of agricultural fields) as well as  
944 agricultural cover.

945 We selected spatially independent agricultural landscapes (between 60 and 100% of agricultural  
946 cover) within each region (Fig. S2.2), representing the maximum variation for both crop  
947 compositional heterogeneity and crop configurational heterogeneity.

948  
949



950  
951 **Figure S2.2.** Spatial distribution of landscapes sampled in one of the eight regions (PVDS = Plaine et Val  
952 de Sèvre).

953  
954

#### 955 **References**

956  
957 Olson, D.M., Dinerstein, E., Wikramanayake, E.D., Burgess, N.D., Powell, G.V.N., Underwood, E.C., D'amico, J.A.,  
958 Itoua, I., Strand, H.E., Morrison, J.C., Loucks, C.J., Allnutt, T.F., Ricketts, T.H., Kura, Y., Lamoreux, J.F.,  
959 Wettengel, W.W., Hedao, P. & Kassem, K.R. (2001) Terrestrial Ecoregions of the World: A New Map of Life  
960 on Earth A new global map of terrestrial ecoregions provides an innovative tool for conserving  
961 biodiversity. *BioScience*, 51, 933–938.

962 Pasher, J., Mitchell, S., King, D., Fahrig, L., Smith, A. & Lindsay, K. (2013) Optimizing landscape selection for  
963 estimating relative effects of landscape variables on ecological responses. *Landscape Ecology*, 28, 371–  
964 383.

965  
966

967 **SI 3 – Multitrophic diversity sampling**

968  
969

970 **3.1. Sampling site selection**

971 Disentangling the effects of crop diversity and mean field size on multitrophic diversity required  
972 sampling many landscapes. Trade-offs between the number of landscapes sampled and the number  
973 of sampling sites per landscape were unavoidable. Whereas studies assessing the effect of landscape  
974 structure on biodiversity are often based on a single sampling site per landscape, we decided to  
975 sample three sampling sites (i.e. three agricultural fields) within each landscape of 1 x 1 km (Fig.  
976 S3.1). These sites were located at least 200 m apart from each other, at least 50 m from the border  
977 of the 1km x 1km landscape, and at least 50 m from non-agricultural cover such as forests.

978  
979 We sampled either one, two or three distinct crop types per landscape. We located these sampling  
980 sites in dominant crop types within each region. When this was not feasible, we located sampling  
981 sites in crop types available within a given landscape while limiting correlations between crop types  
982 sampled and the two heterogeneity gradients within each region (see further details in SI 5).

983  
984 At each sampling site, we selected two parallel 50 m ‘transects’, one located at the field edge and the  
985 other inside the field 25 m away from the first transect (Fig. S3.2).  
986



987  
988 **Figure S3.1.** Example landscape showing the three selected sampling sites.  
989

990 **3.2. Multitrophic diversity sampling within each sampling site**

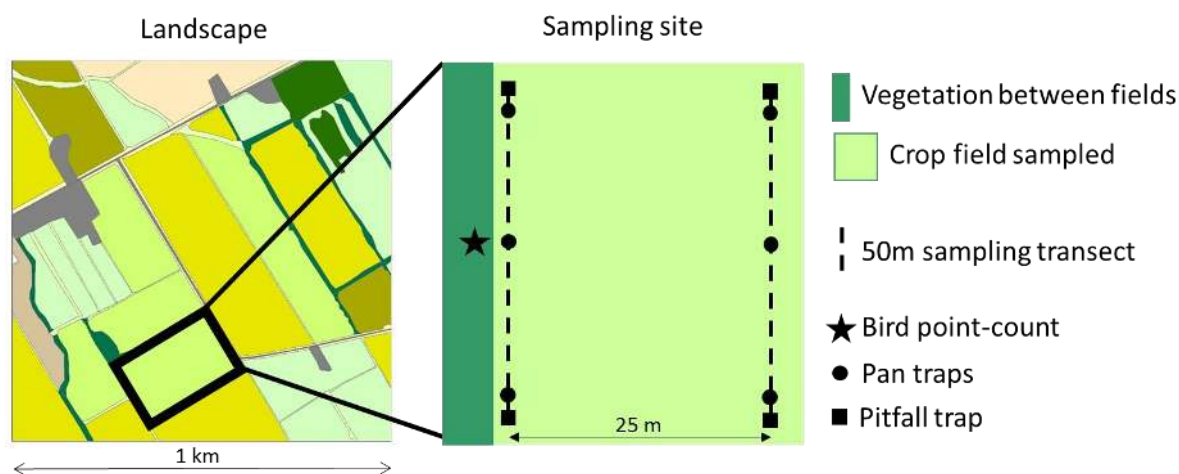
991 Multitrophic diversity sampling occurred between 2011 and 2014 depending on the region and  
992 landscape (Table S3.1).  
993

994 **Table S3.1.** Number of landscapes sampled and main crop types sampled within each region and each year.

Region	2011	2012	2013	2014	Total	Crop types sampled
Armorique			30	10	40	cereal, corn, grassland
Camargue			32	8	40	rice, cereal
Coteaux			20	12	32	cereal, corn, sunflower
East Anglia		30	30		60	cereal, sugar beet, oilseed rape
Goettingen			32	20	52	cereal, oilseed rape, grassland
Lleida			25	15	40	cereal, almond, olive
Eastern Ontario	46	47			93	corn, soybean, grassland



995  
 996 All taxa were sampled using sampling methods commonly used in the literature (point counts, traps,  
 997 visual surveys; Fig. S3.2; Fahrig et al. 2015).  
 998



999  
 1000 **Figure S3.2.** Multitrophic diversity sampling design within each sampling site within each landscape (1  
 1001 km x 1 km).  
 1002

1003 While trade-offs between the number of sites sampled and sampling intensity at each site were  
 1004 necessary, our sampling efforts (see below: number of traps, length of transects, number of visits)  
 1005 were consistent with the literature (e.g. Pollard and Yates 1993, Bibby et al. 2005, Geiger et al. 2010).  
 1006 **Table S3.2** shows the number of species and specimens we sampled for each taxa.

1007  
 1008 **Table S3.2.** Number of species and specimens (occurrences for plants) for each taxa.

	Species	Specimens
All taxa	2795	167028
Bees	343	13326
Birds	208	10911
Butterflies	109	10605
Carabids	256	42547
Hoverflies	146	21491
Plants	1229	30276
Spiders	504	37872

1009  
 1010 **Plants** - Plant surveys were conducted along the field edge and in the field interior transects.  
 1011 Percentage cover was recorded for each species. Each transect was 1 m wide and 50 m long and  
 1012 represented a total surveyed area of 20 m<sup>2</sup>, except in Eastern Ontario where plant survey transects  
 1013 were 2m wide, represented a total surveyed area of 100 m<sup>2</sup> and the field edge transect included both  
 1014 the field and the boundary vegetation. Plant surveys were conducted once, except in Eastern  
 1015 Ontario, Goettingen and East Anglia where surveys were conducted twice.  
 1016

1017 **Bees and hoverflies** – Bees and hoverflies were sampled using colored pan traps, except for  
 1018 hoverflies in Eastern Ontario which were sampled by sweep-netting along the two transects. Plastic  
 1019 bowls painted in UV blue, white or UV yellow were placed in pairs at each end and at the center of  
 1020 each transect. As a result, we used six pan traps per transect, 12 pan traps per sampling site and 36  
 1021 pan traps per landscape. The height of pan traps was adjusted to vegetation height. Cups were filled  
 1022 with water, with three drops of odorless soap added per 1L of water. The traps were left in the field  
 1023 for four days. The insects were then stored in 70 % ethanol and later identified to species level. Bee  
 1024 and hoverfly sampling was carried out twice during the growing season (April-July), the dates being

1025 selected in each region based on regional climatic conditions. Therefore rarefied species richness  
1026 could not be calculated. Due to technical and financial constraints, bees could only be identified to  
1027 species level in seven of the eight regions, and in a total of 183 landscapes. This did not affect our  
1028 results (see section 3.3 of this SI).

1029  
1030 **Carabids and spiders** - Carabids and spiders were sampled using pitfall traps (Bertrand et al. 2016).  
1031 Cups were half-filled with a solution of 10 drops of soap and 10 g of salt per 1L of water and placed in  
1032 the ground. One trap was placed at each end of each transect (two traps per transect and four per  
1033 sampling site in total). The traps were left in the field for four days. Arthropods were then stored in  
1034 70 % ethanol and carabids and adult spiders were later identified to species level. Carabids and  
1035 spiders were sampled at the same time as the bee and hoverfly sampling (above). They were carried  
1036 out only once in East Anglia in 2012 due to bad weather conditions and could not be conducted in  
1037 rice fields in Camargue due to the presence of water.

1038  
1039 **Butterflies** - Butterfly surveys were conducted along the field edge and in the field interior transects  
1040 (Pollard and Yates 1993). Surveys were conducted on calm (Beaufort scale < 3), sunny days, when the  
1041 temperature was > 15°C. The observer recorded all butterfly species observed within an imaginary 5  
1042 m-sided box (2.5 m to each side, 5 m in front and 5 m high) during approximately 10 min per transect  
1043 (Pollard and Yates 1993). Individuals that could not be identified by sight were captured with a  
1044 butterfly net for closer examination (survey time was stopped during capture and identification).  
1045 Surveys were conducted once, except in Eastern Ontario, Goettingen and Lleida where surveys were  
1046 conducted twice.

1047  
1048 **Birds** - Birds were surveyed using 10-minutes point counts (Bibby et al. 2005) located at the center of  
1049 the border transect. All individuals singing or seen within a distance of 100m were recorded. Birds  
1050 flying across were considered as transients and thus not included. Counts were conducted twice,  
1051 except in East Anglia in 2012 due to bad weather conditions, in Ontario and in rice fields in Camargue  
1052 due to the specific phenology of this crop type, where they were conducted once. Surveys were  
1053 conducted during the peak breeding season, between April and June depending on the region, and  
1054 during peak activity hours, from 1 to 4 hours after sunrise and under good weather conditions.

1055  
1056 **Note on detection and rare species** – Our sampling scheme presents the following characteristics : 1)  
1057 the three fields within each landscape often correspond to different crop types and therefore  
1058 correspond to different species pools; 2) we only sampled each landscape during a single year; 3) we  
1059 sampled some taxa across two sessions within the sampling season but these sessions target distinct  
1060 communities (e.g. spring versus summer spider communities); 4) some protocols involve multiple  
1061 sampling within the field (e.g. several pitfall traps along the edge transect and several pitfall traps  
1062 along the center transect) but these traps cannot be considered as replicates due to the high level of  
1063 heterogeneity within fields, both between transects and within a transect. As a result, we do not  
1064 think we have truly replicated data that would allow us computing species richness estimators such  
1065 as the Chao estimator. Nevertheless, because we used standard protocols commonly used in the  
1066 literature, we believe that when pooling the data at the landscape level, our uncorrected data is a  
1067 good proxy of species richness for each taxa studied.

### 1068 1069 **3.3. Multidiversity**

1070 An important challenge when studying the overall effects of crop heterogeneity on multitrophic  
1071 diversity is that different taxa might respond differently (Flynn et al. 2009; Kormann et al. 2015;  
1072 Concepción 2016). Indeed, we observed weak correlations among taxa within our dataset (Table  
1073 S3.3) and significant differences in the response of taxa (Fig. 4 in the main text).

1074  
1075 **Table S3.3.** Mean species richness per landscape  $\pm$  standard deviation for each taxa and correlations among  
1076 taxa (Pearson correlation coefficients). \*  $p < 0.05$ ; \*\*  $p < 0.01$ ; \*\*\*  $p < 0.001$ .

1077

	Mean SR	birds	bees	butterflies	carabids	hoverflies	plants
birds	18.7±6.7						
bees	11.2±4.6	0.11					
butterflies	5.4±2.9	0.03	0.14				
carabids	12.3±6.8	0.01	-0.18*	0.13**			
hoverflies	6.4±3.7	-0.04	0.14	0.09	0.25		
plants	44.9±17.5	0.19	-0.07	0.23	-0.21	0.12	
spiders	20.6±11.5	0.17*	0.41***	-0.20**	0.34***	0.16***	-0.27

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To test the overall effects of crop heterogeneity on multitrophic diversity, we investigated methods developed by Allan et al. (2014) to study ecosystem multifunctionality. Such approach differs from testing how crop heterogeneity impacts each taxa separately by searching for optimal landscape conditions that promote most taxa simultaneously.

A first approach to achieve this is to calculate a multidiversity index based on the averaged approach (Byrnes et al. 2014). This approach consists simply in calculating the average standardized values of multiple taxonomic diversities for each landscape, as follows:

1088

$$\text{Average-based Multidiversity} = \frac{1}{7} \times \sum_{i=1}^{n=7} \text{scale}(SR_i, \text{center}=T, \text{scale}=T)$$

1089

where  $SR_i$  is the number of species for taxa  $i$  in a given landscape.

1090

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1097

Although this averaging approach provides an intuitive method to assess changes in diversity across multiple taxa simultaneously (Allan et al. 2014), the averaged-approach includes some biases. For instance, very high averaged-multidiversity values implies that all groups exhibit high diversity. However, intermediate averaged-multidiversity values are difficult to interpret and it is impossible to differentiate situations where (i) diversity values are intermediate for all taxa simultaneously; or (ii) diversity values are very high for some groups while they are very low for others, i.e. trade-offs among taxa (Byrnes et al. 2014).

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To overcome this limitation, we used a threshold approach (Zavaleta et al. 2010) not biased by potential trade-offs among taxa (Byrnes et al. 2014). The objective of this approach is to assess the ability of agricultural landscapes to simultaneously host at least a given percentage, or threshold ( $x$ ), of the maximum species richness observed for each taxa ( $SR_{max}$ ). Because  $SR_{max}$  is likely to vary between regions, we chose to use the 95th percentile of the maximum observed species richness within each region as  $SR_{max,region}$  for each taxa. We then calculated the multidiversity index based on the following formula:

1105

$$\text{Threshold – based Multidiversity (Tx.landscape)} = \frac{1}{7} \sum_{i=1}^{n=7} (SR_i > (x \times SR_{max,region j}))$$

1106

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where  $SR_i$  is the number of species for taxa  $i$ ,  $x$  is the minimum % to reach and  $SR_{max,region}$  is the maximum species richness for group  $i$  in the region the landscape considered belong to. For a given taxon, if  $SR_i$  is above the threshold, this taxon is associated with the value 1. The sum ranges between 0 and 7, and the multidiversity index ranges between 0 and 1.

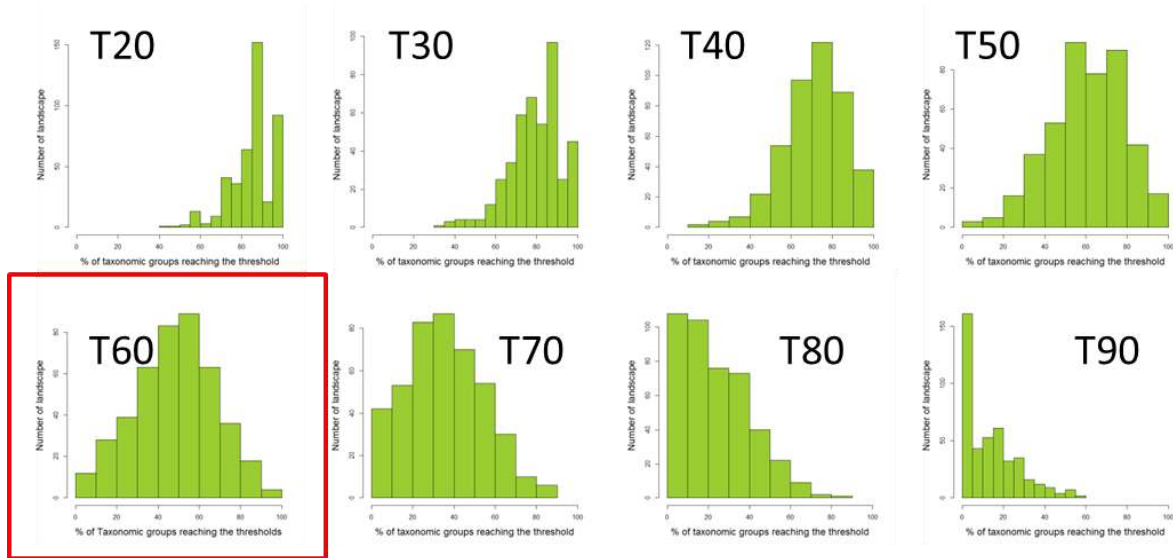
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We calculated this multidiversity index for each threshold  $x$  between 20 and 90% (every 10%). For each threshold  $x$ , the multidiversity index was smoothed by calculating the average over the interval  $[x - 10\%, x + 10\%]$  (Le Bagousse-Pinguet et al. 2019). It is recommended to focus on intermediate

1114 thresholds since care should be taken to avoid over-interpreting high or low thresholds (Lefcheck et  
 1115 al. 2015) and intermediate thresholds have been shown to provide an effective measure of  
 1116 multitrophic diversity in agricultural landscapes (Byrnes et al. 2014). We chose to focus our analyses  
 1117 on the threshold of 60% after checking that the distribution of T60.landscape allows developing  
 1118 robust linear statistics (Fig.S3.3).  
 1119



1120 **Figure S3.3.** Distribution of the threshold-based multitrophic diversity calculated at the landscape  
 1121 level for thresholds between 20 and 90%.  
 1122

1123 A high multidiversity value based on a threshold of 60% means that most taxa are associated  
 1124 with species richness levels higher than 60% of the regional maximum (SRmax.region) observed in  
 1125 our study. Note that (i) T60.landscape was highly correlated with the averaged multidiversity index in  
 1126 our dataset and other threshold-based multidiversity indices (Table S3.4) (ii) our results were not  
 1127 sensitive to the threshold selected (Fig. S5.1 in SI 5).  
 1128

1129 **Table S3.4.** Correlation between average-based multidiversity (M), various threshold-based  
 1130 multidiversity indices calculated at the landscape level (T) and species richness for each taxa. Colors  
 1131 correspond to increasing correlation values (from orange to dark red).  
 1132

	M	T20	T30	T40	T50	T60	T70	T80	Plant	Bee	Syrphid	Butterfly	Carabid	Spider	Bird
M	1	0.48	0.60	0.71	0.80	0.86	0.88	0.86	0.51	0.59	0.39	0.54	0.56	0.64	0.37
T20	0.48	1	0.92	0.77	0.65	0.58	0.52	0.47	0.08	0.19	0.23	0.21	0.07	0.22	0.61
T30	0.60	0.92	1	0.93	0.79	0.69	0.62	0.56	0.15	0.31	0.23	0.28	0.21	0.31	0.59
T40	0.71	0.77	0.93	1	0.93	0.82	0.74	0.66	0.23	0.45	0.27	0.34	0.33	0.40	0.54
T50	0.80	0.65	0.79	0.93	1	0.94	0.85	0.74	0.32	0.54	0.28	0.39	0.41	0.48	0.50
T60	0.86	0.58	0.69	0.82	0.94	1	0.95	0.84	0.38	0.57	0.28	0.44	0.45	0.54	0.46
T70	0.88	0.52	0.62	0.74	0.85	0.95	1	0.95	0.42	0.54	0.29	0.45	0.46	0.59	0.43
T80	0.86	0.47	0.56	0.66	0.74	0.84	0.95	1	0.42	0.48	0.29	0.43	0.45	0.57	0.44
Plant	0.51	0.08	0.15	0.23	0.32	0.38	0.42	0.42	1	0.04	0.01	0.22	0.21	0.18	0.00
Bee	0.59	0.19	0.31	0.45	0.54	0.57	0.54	0.48	0.04	1	0.25	0.24	0.19	0.30	0.12
Syrphid	0.39	0.23	0.23	0.27	0.28	0.28	0.29	0.29	0.01	0.25	1	0.07	0.06	0.06	-0.06
Butterfly	0.54	0.21	0.28	0.34	0.39	0.44	0.45	0.43	0.22	0.24	0.07	1	0.14	0.20	0.03
Carabid	0.56	0.07	0.21	0.33	0.41	0.45	0.46	0.45	0.21	0.19	0.06	0.14	1	0.34	-0.02
Spider	0.64	0.22	0.31	0.40	0.48	0.54	0.59	0.57	0.18	0.30	0.06	0.20	0.34	1	0.15
Bird	0.37	0.61	0.59	0.54	0.50	0.46	0.43	0.44	0.00	0.12	-0.06	0.03	-0.02	0.15	1

1134 Data for bee species richness were only available for 183 landscapes. To determine whether  
 1135 this affected our results, we also calculated the multidiversity index across six taxa (all groups except  
 1136  
 1137

1138 bees). As there was no difference in results obtained with six or seven taxa, we here only present  
1139 results for the multidiversity index calculated across seven taxa within 435 landscapes.

1140  
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1142 **References**

1143

1144 Allan E, et al. (2014) Interannual variation in land-use intensity enhances grassland multidiversity. *Proc Natl Acad Sci U S A* 111(1):308–313.

1146 Bertrand C, Burel F, Baudry J (2016) Spatial and temporal heterogeneity of the crop mosaic influences carabid  
1147 beetles in agricultural landscapes. *Landsc Ecol* 31(2):451–466.

1148 Bibby, C.J., et al. (2005) *Bird Census Techniques*. Academic Press, London, UK.

1149 Byrnes JEK, et al. (2014) Investigating the relationship between biodiversity and ecosystem multifunctionality:  
1150 challenges and solutions. *Methods Ecol Evol* 5(2):111–124.

1151 Concepción ED, et al. (2016) Contrasting trait assembly patterns in plant and bird communities along  
1152 environmental and human-induced land-use gradients. *Ecography* 40(6):753–763.

1153 Fahrig L, et al. (2015) Farmlands with smaller crop fields have higher within-field biodiversity. *Agric Ecosyst*  
1154 *Environ* 200:219–234.

1155 Flynn DFB, et al. (2009) Loss of functional diversity under land use intensification across multiple taxa. *Ecol Lett*  
1156 12(1):22–33.

1157 Geiger, F., Bengtsson, J., Berendse, F., Weisser, W. W., Emmerson, M., Morales, M. B.,... & Eggers, S. (2010).  
1158 Persistent negative effects of pesticides on biodiversity and biological control potential on European  
1159 farmland. *Basic and Applied Ecology*, 11(2), 97–105.

1160 Kormann U, et al. (2015) Local and landscape management drive trait-mediated biodiversity of nine taxa on small  
1161 grassland fragments. *Divers Distrib* 21(10):1204–1217.

1162 Le Bagousse-Pinguet Y, et al. (2019) Phylogenetic, functional, and taxonomic richness have both positive and  
1163 negative effects on ecosystem multifunctionality. *Proc Natl Acad Sci*:201815727.

1164 Lefcheck JS, et al. (2015) Biodiversity enhances ecosystem multifunctionality across trophic levels and habitats.  
1165 *Nat Commun* 6:6936.

1166 Pollard, E., Yates, T.J. (1993). *Monitoring butterflies for ecology and conservation*. Chapman et Hall, London.

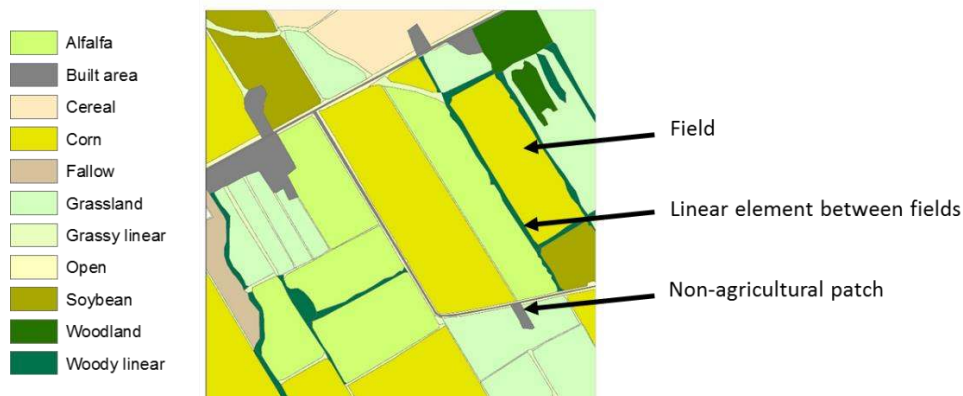
1167 Zavaleta ES, Pasari JR, Hulvey KB, Tilman GD (2010) Sustaining multiple ecosystem functions in grassland  
1168 communities requires higher biodiversity. *Proc Natl Acad Sci U S A* 107(4):1443–1446.

## SI 4 – Land cover mapping and landscape metrics

### 4.1. Land cover mapping

Land cover was mapped based on remotely-sensed data and ground-truthing. All cover types, including fields, linear elements between fields and non-agricultural cover types, were mapped as polygons ('patches') (Fig. S4.1). We here refer to 'cover types' rather than 'habitats' because 'habitat' refers to the specific ecological requirements of a given species while 'cover type' refers to a category of land cover without any assumption on species use. This is important in the present study where we assume that many farmland species are likely to use several cover types (landscape complementation).

Agricultural cover types included: cereal, fallow, alfalfa, clover, ryegrass, rice, corn, sunflower, sorghum, millet, moha, oilseed rape, mustard, pea, bean, soybean, linseed, orchard, almond, olive, vineyard, mixed vegetables, sugar beet, asparagus, carrot, onion, parsnip, potato, tomato, melon, strawberry, raspberry, wild bird cover, grassland (including temporary and permanent grassland managed for production purpose) and other crops (unknown or rare crops). We chose to include managed grassland within agricultural cover types because we were interested in assessing the role of spatial heterogeneity within the farmed part of the landscape. We considered grasslands where more than 50% of the biomass was removed as agricultural cover whereas those where less than 50% of the biomass was removed were considered as non-agricultural cover. Linear elements between fields were classified either as woody, grassy, water (e.g. ditches) or tracks. Non-agricultural cover types included woodland (including woody linear elements), open land (e.g. shrubland, grassy linear elements), wetland and built-area (including roads).



**Figure S4.1.** Example of land cover map used to calculate variables within each landscape (1km x 1km).

### 4.2. Landscape metrics

It is well known that different taxa and even species are likely to respond to the landscape structure at different spatial scales. Since our aim was to assess the overall effects of crop diversity and mean field size on a range of contrasted taxa, we chose to calculate landscape variables within a 1x1 km because this spatial extent represent the best compromise between highly mobile taxa (e.g. birds) and taxa with more limited dispersal abilities (e.g. plants or spiders; Kormann et al. 2015).

#### 4.2.1. Number of crop types sampled

The number of crop types sampled ranged from one to three. The diversity of crop types available in the landscape and the number of crop types sampled within each landscape were not heavily correlated ( $r=0.45$ ).

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#### 4.2.2. Crop compositional heterogeneity

We used the diversity of crop types available in the landscape (hereafter ‘crop diversity’) as a measure of crop compositional heterogeneity. We measured crop diversity using the Shannon diversity index, a widely used metric of landscape heterogeneity (e.g. Bertrand et al. 2016; Bosem Baillod et al. 2017):  $H' = -\sum_{i=1}^n p_i \ln p_i$  where  $p_i$  is the proportion of crop type  $i$  in the agricultural mosaic. Note that this metric assumes that all agricultural cover types (defined in 4.1) are considered equally different. This variable does not take into account within-field crop heterogeneity, e.g. intercropping patterns.

The diversity of crop types available in the landscape and the number of crop types sampled within each landscape were not heavily correlated ( $r=0.45$ ).

#### 4.2.3. Crop configurational heterogeneity

We used mean field size (ha) as a measure of crop configurational heterogeneity. We chose this metric over total field perimeter length per landscape (e.g. Bosem Baillod et al. 2017) because it is directly related to our hypotheses (see SI 1). Moreover it is easier to base practical recommendations for future agricultural policies on mean field size rather than on total field perimeter length. Fields were only mapped within the 1 km<sup>2</sup> landscape. As a result, for fields located partly outside of the 1 km<sup>2</sup> landscape, only their area contained within the landscape was considered in calculating mean field size. This may lead to a slight underestimation of mean field size.

#### 4.2.4. Semi-natural cover proportion

We calculated the sum of woodland (including woody linear elements), open land (e.g. shrubland, grassy margins) and wetland cover (including ponds, rivers, ditches) in the landscape.

#### 4.2.5. Total length of semi-natural linear elements

We assessed the total length of vegetation occurring in semi-natural linear elements between fields (SNL, in meters) by calculating half the sum of all semi-natural linear elements located between two fields (e.g. hedgerows, grassy margins). Note that semi-natural linear elements located along roads or urban areas were not included in the calculation of SNL. SNL and mean field size were highly correlated (see Table S5.5. in SI 5).

#### 4.2.6. Latitude and longitude

We calculated the latitude and longitude of the center of each landscape using the WGS 1984 World Mercator projection system.

### 4.3. Descriptive statistics for the 435 landscapes selected

The 435 landscapes selected across eight regions of Europe and North America had the following characteristics (mean  $\pm$  sd; see also Table S4.1): 1.94 $\pm$ 0.56 crop types sampled, 81.3 $\pm$ 9.6 % of agricultural cover, 12.7 $\pm$ 8.9 % of semi-natural cover, 5631 $\pm$ 3822 m of linear semi-natural elements between fields, mean field size 2.99 $\pm$ 2.02 ha and a Shannon diversity index of agricultural cover types of 1.03 $\pm$ 0.39 (Fig S4.3). These gradients are representative of most Western European agricultural landscapes (Herzog et al. 2006) and most American agricultural landscapes (Yan & Roy 2016).

**Table S4.1.** Descriptive statistics for each landscape variable (mean, median, 25th and 75th quartiles, min and max): number of crop types sampled (Crop nb), diversity of crop types available in the landscape (Crop diversity), mean field size (ha), the percentage of semi-natural cover types (SNC), and the length of semi-natural linear elements (SNL).

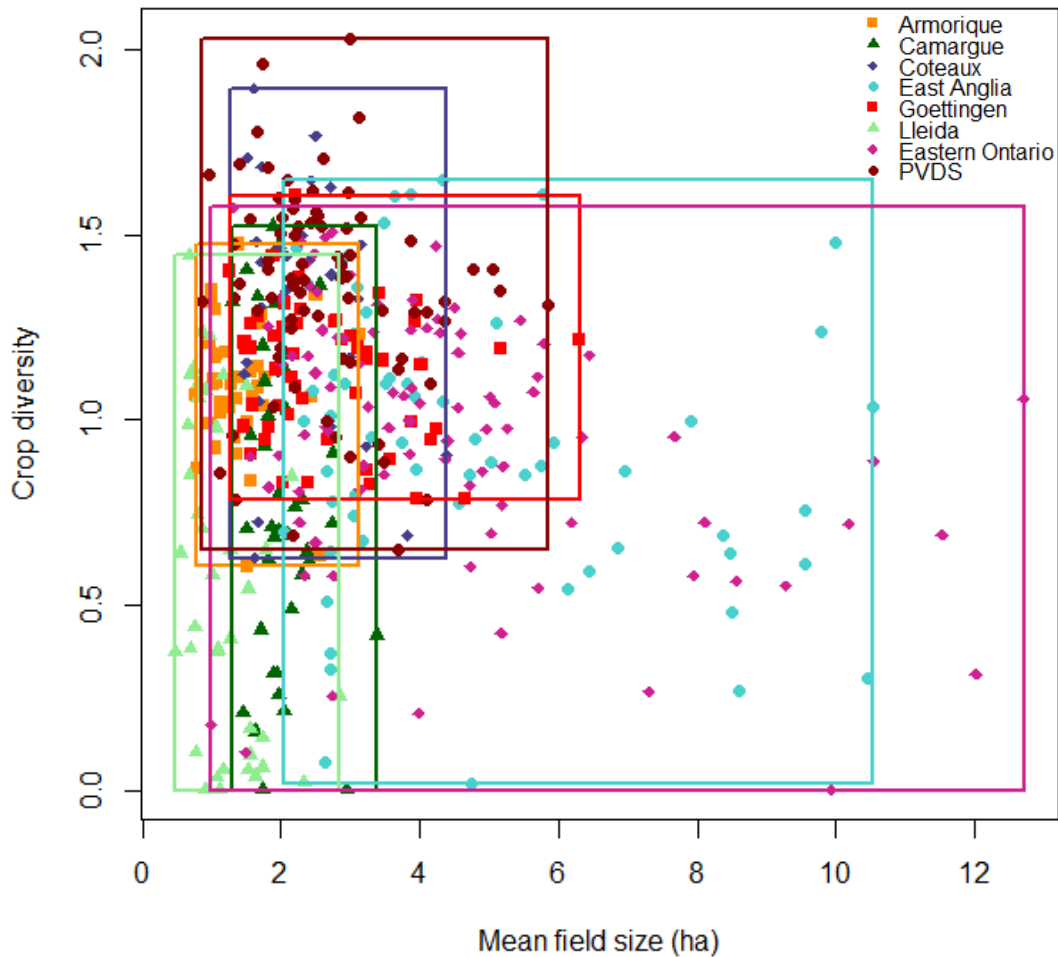
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Crop nb	Crop diversity	Mean field size (ha)	SNC (%)	SNL (m)
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Min	1	0.0	0.48	0.0	0
1st quartile	2	0.8	1.71	6.0	3108
Median	2	1.09	2.43	10.9	4824
Mean	1.94	1.03	2.99	12.7	5632
3rd quartile	3	1.31	3.69	17.6	7370
Max	3	2.03	12.71	49.5	27989

1260



1261

1262 **Figure S4.3.** Variation in crop diversity and mean field size (ha) across the eight regions. Points  
 1263 correspond to selected landscapes (N= 435) and boxes corresponds to the range of crop diversity and  
 1264 mean field size sampled within each region (orange=Armorique, dark green=Camargue, dark  
 1265 blue=Coteaux, light blue=East Anglia, light red=Goettingen, light green=Lleida, pink=Eastern Ontario,  
 1266 dark red=PVDS).

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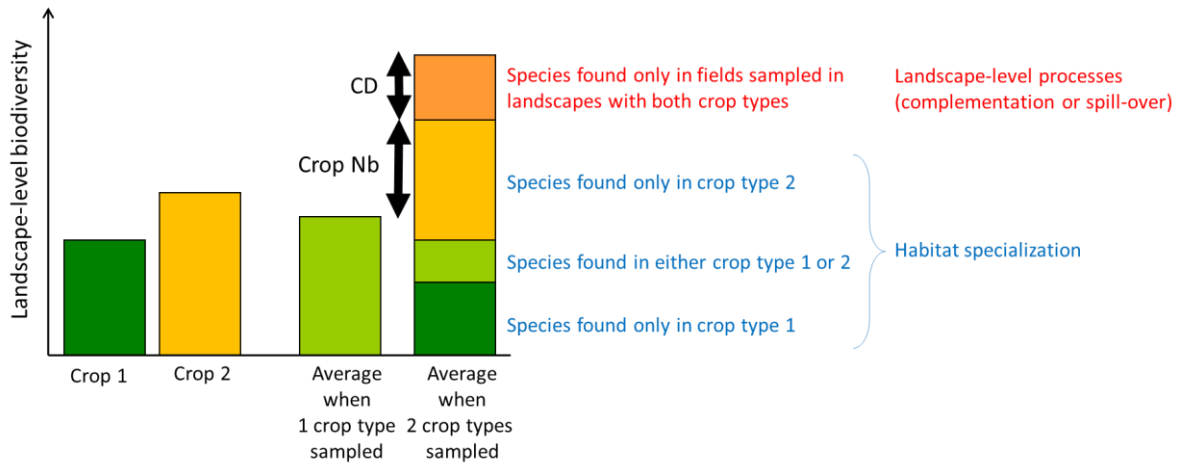
#### 1269 4.4. Effects of the number of crop types sampled vs. the diversity of crop types in the landscape

1270

1271 Biodiversity may increase with increasing crop diversity if different crop types can serve as  
 1272 habitat for different specialized species (single habitat specialization; Fig. S4.4). In that case, sampling  
 1273 more crop types will result in higher observed landscape-level multitrophic diversity. Biodiversity  
 1274 may also increase with crop diversity if different crop types provide different resources required for

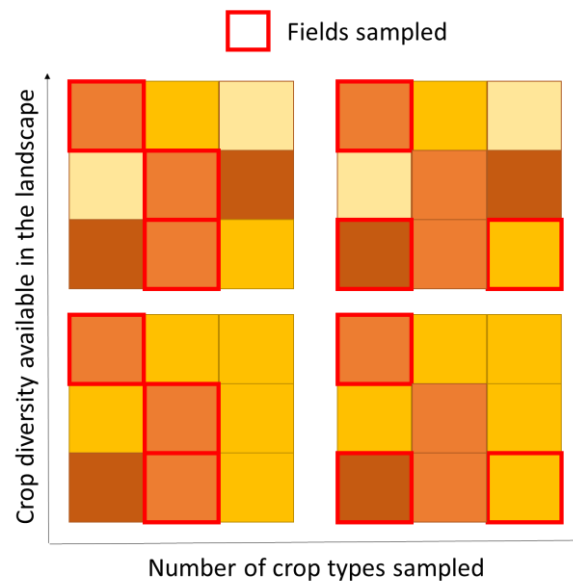


1275 single species (landscape complementation). In that case, sampling the same number of crop types in  
 1276 landscapes with higher crop diversity will result in higher landscape-level multitrophic diversity.  
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 1279 **Figure S4.4.** Roles of habitat specialization, landscape complementation or spill-over in the potential  
 1280 positive effect of crop diversity on multitrophic diversity (see SI 1). Black arrows represent the effect  
 1281 of our two explanatory variables (CD = increasing the diversity of crop types in the landscape; Crop  
 1282 Nb = increasing the number of crop types sampled).  
 1283

1284 Since the diversity of crop types available in the landscape and the number of crop types  
 1285 sampled within each landscape were not heavily correlated ( $r=0.45$ ), we were able to disentangle the  
 1286 role of these two mechanisms (Fig. S4.5).  
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 1289 **Figure S4.5.** Representation of our sampling design allowing us to take into account the potential  
 1290 contribution of habitat specialization and landscape complementation/spillover to the positive effect  
 1291 of crop diversity on multitrophic diversity.  
 1292

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 1294 **References**

1295  
 1296 Bertrand C, Burel F, Baudry J (2016) Spatial and temporal heterogeneity of the crop mosaic influences carabid  
 1297 beetles in agricultural landscapes. *Landsc Ecol* 31: 451–466.

1298 Bosem Baillod A, Tschardtke T, Clough Y, Batáry P (2017) Landscape-scale interactions of spatial and temporal  
1299 cropland heterogeneity drive biological control of cereal aphids. *J Appl Ecol*, 54: 1804-1813.  
1300 Herzog F, et al. (2006) Assessing the intensity of temperate European agriculture at the landscape scale. *Eur J*  
1301 *Agron* 24(2): 165–181.  
1302 Kormann U, et al. (2015) Local and landscape management drive trait-mediated biodiversity of nine taxa on small  
1303 grassland fragments. *Divers Distrib* 21(10):1204–1217.  
1304 Yan, L. & Roy, D.P. (2016) Conterminous United States crop field size quantification from multi-temporal Landsat  
1305 data. *Remote Sensing of Environment*, 172: 67–86.

1306 **SI 5 – Complementary analyses**

1307

1308 **5.1. Details of the model selection and model averaging for multitrophic diversity**

1309

1310 We first tested the effect of crop heterogeneity on multitrophic diversity (Model 1).

1311

1312 *Model 1: lmer (T60.landscape ~ CD \* MFS \* SNC + CD<sup>2</sup> + MFS<sup>2</sup> + SNC<sup>2</sup> + CropNb + Lat + Lon + (1|*  
 1313 *Region/Year))*

1314

1315 The model selection approach based on  $\Delta AICc < 2$  resulted in the selection of 10 sub-models  
 1316 (Table S5.1). Using a  $\Delta AICc$  of 7 did not change the results of the model averaging or results on  
 1317 variable importance. All models included crop diversity (CD), mean field size (MFS), semi-natural  
 1318 cover (SNC), the number of crops sampled per landscape (Crop nb) and the interaction between crop  
 1319 diversity and semi-natural cover (CD x SNC). The AICc of the Null model was 3709 while the AICc of  
 1320 the best model was 3667, i.e. a  $\Delta AICc$  of 42, suggesting that the best selected models were far more  
 1321 parsimonious than the null model including only Region and Year as random effects.

1322

1323 **Table S5.1.** List of all sub-models selected and used for the model averaging approach for model 1.

Sub-model	1-1	1-2	1-3	1-4	1-5	1-6	1-7	1-8	1-9	1-10	Null
Lat											
Lon											
CD											
CD <sup>2</sup>											
MFS											
MFS <sup>2</sup>											
SNC											
Crop nb.											
CD x SNC											
MFS x SNC											
CD x MFS x SNC											
Marginal R <sup>2</sup>	0.13	0.13	0.12	0.15	0.15	0.12	0.14	0.12	0.14	0.13	0
Conditional R <sup>2</sup>	0.36	0.37	0.35	0.38	0.38	0.36	0.37	0.37	0.37	0.36	0.23
df	10	11	9	11	12	10	10	11	11	11	–
AICc	3667.5	3668.09	3668.16	3668.21	3668.75	3668.75	3668.86	3669.36	3669.39	3669.56	3709.7
delta	0	0.59	0.66	0.7	1.24	1.24	1.35	1.85	1.89	2.05	42.23
weight	0.17	0.13	0.12	0.12	0.09	0.09	0.09	0.07	0.07	0.06	–

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1327 **5.2. Influence of selected threshold on parameter estimates for multitrophic diversity**

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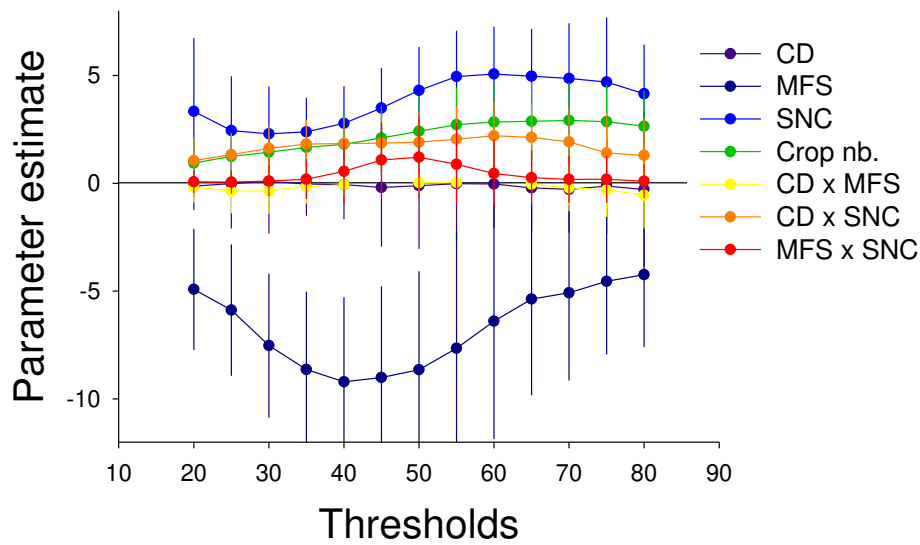
1329 To test whether the choice of threshold for computing the multitrophic diversity index  
 1330 impacted our conclusions, we ran model 2 for all thresholds from T20 to T80 (i.e. proportion of taxa  
 1331 for which the species richness is equal to or higher than 20% to 80% of the regional maximum species  
 1332 richness per landscape).

1333

1334 Parameters estimates were consistent across the range of thresholds (Fig. S5.1). Moreover,  
 1335 variations in parameter estimates suggests that increasing mean field size may be particularly  
 1336 effective to reach intermediate multidiversity thresholds (i.e. between 30 and 50% of regional  
 1337 maximum) whether increasing semi natural cover may be effective to reach higher multidiversity  
 1338 threshold (i.e. above 50% of regional maximum).

1338

1339 This comparison confirms the validity of choosing T60.landscape, i.e. the proportion of taxa  
 1340 for which the species richness is equal or higher than 60% of the regional maximum species richness  
 per landscape.



1341 **Figure S5.1.** Parameter estimates based on model 1 for different thresholds. Thresholds correspond  
 1342 to the % of SR max used to calculate the multidiversity index. In this paper, we present model  
 1343 outcomes for a threshold of 60%, i.e. we use the proportion of taxa that exceeded 60% of the  
 1344 maximum species richness.  
 1345  
 1346

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 1348 **5.3. Variation in the response of multitrophic diversity among regions**  
 1349

1350 To test whether the effects of crop diversity, mean field size and semi-natural cover on  
 1351 multitrophic diversity measured at the landscape level (T60.landscape) varied significantly among  
 1352 regions we added random effects for region on the slopes of crop diversity, mean field size, semi-  
 1353 natural cover as well as the interaction between crop diversity and semi-natural cover (model 2). We  
 1354 assumed that the effects of region on the intercept and slopes were uncorrelated. To test whether  
 1355 Region had a significant effect on the slope of either crop diversity, mean field size, semi-natural  
 1356 cover as well as the interaction between crop diversity and semi-natural cover, we used the function  
 1357 exactRLRT from package RLRsim.

1358 *Model 2: Imer (T60.landscape ~ CD \* MFS \* SNC + CD<sup>2</sup> + MFS<sup>2</sup> + SNC<sup>2</sup> + CropNb + Lat + Lon +*  
 1359 *(1|Region/Year) + (0+CD|Region)) + (0+MFS|Region) + (0+SNC|Region) + (0+CD:SNC|Region))*

1360 **Table S5.2.** Comparison of model 1 and model 2 (i.e. model including a random effect of region on  
 1361 slope). Parameter listed are those retained in the model selection procedure. Parameter estimates  
 1362 and confidence intervals are based on the model averaging approach. ° p<0.1; \* p<0.05; \*\* p<0.01;  
 1363 \*\*\* p<0.001.  
 1364

	model 1	model 2
Crop diversity (CD)	-0.03 [-2.07 ; 2.01]	-0.16 [-2.22 ; 1.9]
Mean field size (MFS)	-6.39 [-11.85 ; -0.94] *	-5.22 [-11.29 ; 0.85] °
Semi-Natural Cover (SNC)	5.07 [2.87 ; 7.26] ***	4.35 [0.79 ; 7.91] *
Nb of Crops sampled	2.84 [1.07 ; 4.62] ***	3.05 [1.29 ; 4.8] ***
Latitude	1.5 [-3.55 ; 6.55]	
Longitude	3.73 [2.47 ; 9.93]	-2.39 [-8.39 ; 3.62]
MFS <sup>2</sup>	3.78 [-0.67 ; 8.23] °	3.78 [-2.26 ; 9.83]
SNC <sup>2</sup>		-2.39 [-8.39 ; 3.62]
CD :SNC	2.20 [0.64 ; 3.76] **	2.06 [0.29 ; 3.82] *
MFS :SNC	1.15 [-0.66 ; 2.96]	1.51 [-0.44 ; 3.46]

1365

1366 The random effect of region on the slope of MFS was significant in model 2 (RLRT = 3.28,  
1367 p=0.02) whereas the effects on CD (RLRT=0, p=1), SNC (RLRT=0.04, p=0.33) and CD:SNC (RLRT=0.19,  
1368 p=0.24) were not (Fig. 4). This result confirms that the regional context can modulate the effect of  
1369 mean field size on multitrophic diversity, but that the positive effects of increasing CD, when SNC is  
1370 high enough, and decreasing MFS remain valid across all regions (Table S5.2).

1371

#### 1372 5.4. Results on the species richness of taxonomic groups

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1374 We tested the effects of crop heterogeneity on the species richness of taxonomic groups  
1375 (Model 3). To do this, we fitted a similar model, using the landscape-level species richness of  
1376 taxonomic groups (SR) as the response variable. To reflect that species pools vary between taxa, we  
1377 added Taxon as a random effect.

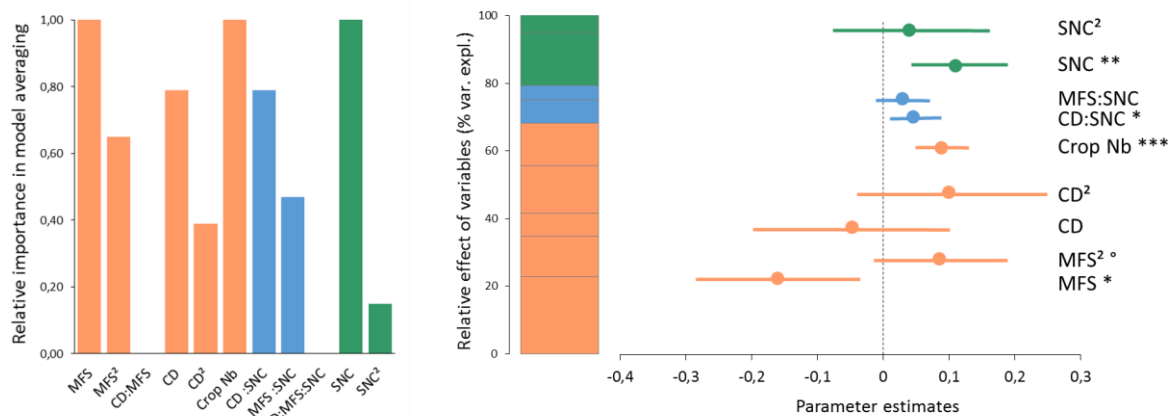
1378

1379 *Model 3: lmer (SR ~ CD\*MFS\*SNC + CD<sup>2</sup> + MFS<sup>2</sup> + SNC<sup>2</sup> + CropNb + Lat + Lon + (1|Region/Year) +*  
1380 *(1|Taxon))*

1381

1382 The effects of crop heterogeneity on the species richness of taxonomic groups were similar  
1383 to their effects on multitrophic diversity (Fig. S5.2).

1384



1385

1386 **Figure S5.2.** Response of the species richness of taxonomic groups to the diversity of crop types  
1387 available within the landscape (CD), the number of crops sampled (Crop Nb), mean field size (MFS),  
1388 semi-natural cover (SNC), and interaction terms (CD:SNC, MFS:SNC, see further details in Methods),  
1389 based on data collected in 435 landscapes located in eight agricultural regions. Covariates (Lon, Lat)  
1390 were excluded from the figure for simplicity. Importance of each variable in the model averaging  
1391 approach (model 3), estimated as the proportion of models where the variable was selected. The  
1392 relative effect of each variable corresponds to the ratio between its parameter estimate and the sum  
1393 of all parameter estimates (i.e. the % of variance explained). Parameter estimates and confidence  
1394 intervals, based on a model averaging approach applied to model 3 (Methods). ° p<0.1; \* p<0.05; \*\*  
1395 p<0.01; \*\*\* p<0.001. Variables are grouped in three components: orange = crop heterogeneity (MFS,  
1396 MFS<sup>2</sup>, CD, CD<sup>2</sup>, MFS:CD, Crop Nb), green = semi-natural cover (SNC, SNC<sup>2</sup>), blue = interactive effects  
1397 between crop heterogeneity and semi-natural cover (CD:SNC, MFS:SNC, CD:MFS:SNC).

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1399 To test whether the effects of crop diversity, mean field size and semi-natural cover on the  
1400 species richness of taxonomic groups varied significantly among taxa we added random effects for  
1401 Taxon on the slopes of crop diversity, mean field size, semi-natural cover as well as the interaction  
1402 between crop diversity and semi-natural cover (model 4). We assumed that the effects of Taxon on

1403 the intercept and slopes were uncorrelated. To test whether Taxon had a significant effect on the  
 1404 slope of either crop diversity, mean field size, semi-natural cover or the interaction between crop  
 1405 diversity and semi-natural cover, we used the function exactRLRT from package RLRsim.

1406 *Model 4: Imer (SR ~ CD \* MFS \* SNC + CD<sup>2</sup> + MFS<sup>2</sup> + SNC<sup>2</sup> + CropNb + Lat + Lon + (1|Taxon) +*  
 1407 *(1|Region/Year) + (0+CD|Taxon)) + (0+MFS|Taxon) + (0+SNC|Taxon) + (0+CD:SNC|Taxon))*

1408 **Table S5.3.** Comparison of model 3 and model 4 (i.e. model including a random effect of taxa on  
 1409 slopes). Parameter listed are those retained in the model selection procedure. Parameter estimates  
 1410 and confidence intervals are based on the model averaging approach. ° p<0.1; \* p<0.05; \*\* p<0.01;  
 1411 \*\*\* p<0.001.

	model 3	model 4
Crop diversity (CD)	-0.05 [-0.2 ; 0.11]	-0.05 [-0.21 ; 0.1]
Mean field size (MFS)	-0.16 [-0.28 ; -0.04] *	-0.14 [-0.26 ; -0.03] *
Semi-Natural Cover (SNC)	0.11 [0.04 ; 0.18] **	0.11 [0.06 ; 0.17] ***
Nb of Crops sampled	0.09 [0.05 ; 0.13] ***	0.09 [0.05 ; 0.13] ***
Latitude	0.07 [-0.03 ; 0.16]	0.06 [-0.03 ; 0.16]
CD <sup>2</sup>	0.1 [-0.04 ; 0.24]	0.08 [-0.07 ; 0.23]
MFS <sup>2</sup>	0.08 [-0.02 ; 0.19] °	0.07 [-0.03 ; 0.17]
SNC <sup>2</sup>	0.04 [-0.08 ; 0.16]	0.01 [-0.11 ; 0.13]
CD :SNC	0.04 [0.01 ; 0.08] *	0.05 [0.002 ; 0.09] *
MFS :SNC	0.03 [-0.01 ; 0.07]	0.03 [-0.01 ; 0.07]

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1414 The random effect of taxa on the slope of CD (RLRT = 1.94, p=0.06), MFS (RLRT=0.05, p=0.34),  
 1415 SNC (RLRT=0.26, p=0.24) and CD:SNC (RLRT=0.35, p=0.22) were not significant in model 4 (Fig. 5).  
 1416 This result confirms that the effects of crop heterogeneity on species diversity vary only marginally  
 1417 among taxa, and that the positive effects of decreasing mean field size, increasing the number of  
 1418 crop sampled, increasing semi-natural cover, and when semi-natural cover is high, increasing crop  
 1419 diversity, remain valid across all taxa (Table S5.3).

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## 1422 5.5. Correlations and alternative mechanisms at the landscape level

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1424 Crop diversity and mean field size are likely to be correlated with several variables, including  
 1425 the overall composition of the crop mosaic, the proportion of grassland in the mosaic or the length of  
 1426 semi-natural vegetation occurring between fields. Disentangling the role of crop heterogeneity from  
 1427 the effects of these other variables is necessary in order to infer potential mechanisms explaining the  
 1428 positive effect of crop heterogeneity on multitrophic diversity. In the present study, some of these  
 1429 additional variables were correlated among themselves, or with our variables of interest. Exploring  
 1430 their role sometimes required running models using a data subset for which relevant variables were  
 1431 uncorrelated. As a result, we could not include all these variables in a single model and present these  
 1432 analyses as separate, complementary analyses.

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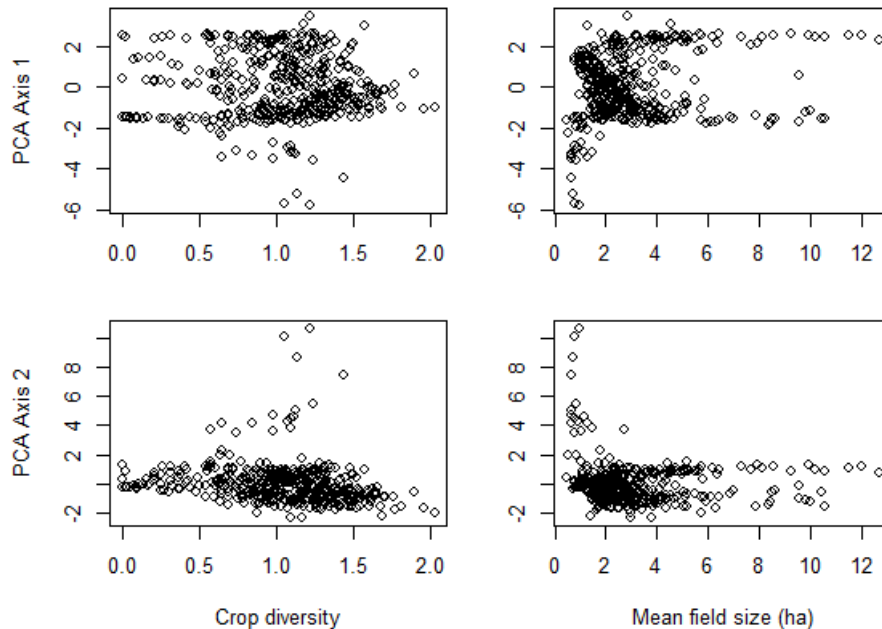
### 1434 5.5.1. Role of the identity of crops in the agricultural mosaic

1435 The identity of crop types in the mosaic may vary along the gradients of crop diversity and  
 1436 mean field size. For instance, landscapes with small fields may be composed of more biodiversity-  
 1437 friendly crops. Such a correlation would represent a potential bias in our study and hamper our  
 1438 ability to test the effects of crop heterogeneity on multitrophic diversity.

1439 We investigated the correlation between each crop heterogeneity gradient and the identity  
 1440 of crop types in the mosaic for 435 landscapes from 8 regions. We conducted a Principal Components

1441 Analysis on the matrix of percentage cover per agricultural cover type per landscape. The first axis  
 1442 represented 40% of the variance, while the second axis represented 19% of the variance.

1443 The Pearson correlations between crop diversity and the first two axes of the PCA were weak  
 1444 (axis 1:  $r=-0.03$ ,  $p=0.56$  and axis 2:  $r=-0.19$ ,  $p<0.001$ ), as were the Pearson correlations between mean  
 1445 field size and the first two axes of the PCA (axis 1:  $r=0.21$ ,  $p<0.001$  and axis 2:  $r=-0.12$ ,  $p=0.01$ ; Fig.  
 1446 S5.3).



1448 **Figure S5.3.** Relationships between the two crop heterogeneity gradients and the identity of crop  
 1449 types in the mosaic (axes 1 and 2 of the Principal Components Analysis).

1452 We added the scores of landscapes along axes 1 and 2 of the PCA to model 1 and compared  
 1453 the outcomes of the obtained model (model 3) with those of model 1.

1454 *Model 1:  $Imer(T60.landscape \sim CD * MFS * SNC + CD^2 + MFS^2 + SNC^2 + CropNb + Lat + Lon + (1 |$*   
 1455 *Region/Year))*

1456 *Model 5:  $Imer(T60.landscape \sim CD * MFS * SNC + CD^2 + MFS^2 + SNC^2 + CropNb + Lat + Lon + Axis1 +$*   
 1457 *Axis 2 + (1 | Region/Year))*

1460 The average model selected based on model 5 included the same variables as the average  
 1461 model selected based on model 1, plus variable PCA Axis 1. Parameter estimates and significance for  
 1462 variables of interest remained unchanged (Table S5.4). This result suggests that the effects of CD, in  
 1463 combination with SNC, and MFS cannot be explained by the composition of crop types occurring in  
 1464 the mosaic.

1465 **Table S5.4.** Comparison of estimates for model 1 and model 5 – mosaic crop composition (i.e. model  
 1466 taking into account the composition of crop types in the mosaic). Parameter listed are those retained  
 1467 in the model selection procedure. Parameter estimates and confidence intervals are based on the  
 1468 model averaging approach. °  $p<0.1$ ; \*  $p<0.05$ ; \*\*  $p<0.01$ ; \*\*\*  $p<0.001$ .

	model 1	model 5 – mosaic crop composition
Crop diversity (CD)	-0.03 [-2.07 ; 2.01]	-0.06 [-2.1 ; 1.96]
Mean field size (MFS)	-6.39 [-11.85 ; -0.94] *	-6.44 [-11.88 ; -1.01] *
Semi-Natural Cover (SNC)	5.07 [2.87 ; 7.26] ***	5.07 [2.88 ; 7.27] ***
Nb of Crops sampled	2.84 [1.07 ; 4.62] ***	2.84 [1.06 ; 4.62] **

Latitude	1.5 [-3.55 ; 6.55]	1.5 [-3.55 ; 6.55]
Longitude	3.73 [2.47 ; 9.93]	3.73 [-2.47 ; 9.93]
MFS <sup>2</sup>	3.78 [-0.67 ; 8.23] °	3.73 [-0.72 ; 8.19]
CD :SNC	2.20 [0.64 ; 3.76] **	2.21 [0.65 ; 3.77] **
MFS :SNC	1.15 [-0.66 ; 2.96]	1.15 [-0.66 ; 2.96]
PCA axis 1		1.5 [-3.55 ; 6.55]

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### 5.5.2. Role of the proportion of grassland in the crop mosaic

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The identity of some ecologically important crop types in the mosaic may vary along the gradients of crop diversity and mean field size. In this study, we chose to include managed grassland within agricultural cover types because we were interested in assessing the role of spatial heterogeneity within the farmed part of the landscape. In our dataset, grassland cover was only moderately correlated with crop diversity ( $r=-0.001$ ,  $p=0.97$ ) and mean field size ( $r=-0.21$ ,  $p<0.001$ ). However, we were aware that the proportion of grassland in the crop mosaic, in particular permanent grassland, may have a strong positive effect on biodiversity (Öckinger & Smith 2007).

We added the proportion of grassland to model 1 (using data collected in 435 landscapes from 8 regions) and compared the outcomes of the following model (model 6) with those of model 1.

*Model 6: Imer (T60.landscape ~ CD \* MFS \* SNC + CD<sup>2</sup> + MFS<sup>2</sup> + SNC<sup>2</sup> + CropNb + Lat + Lon + Grassland + (1 | Region/Year))*

Model selection based on model 6 included the same variables as for model 1, plus Grassland, which had a marginally significant positive effect. However, parameter estimates and significance for other variables of interest remained unchanged (Table S5.5). This result suggests that the effects of CD, in combination with SNC, and MFS cannot be explained by the proportion of grassland in the mosaic.

**Table S5.5.** Comparison of model 1 and model 6 – grassland (i.e. complete model taking into account the proportion of grassland in the mosaic). Parameter listed are those retained in the model selection procedure. Parameter estimates and confidence intervals are based on the model averaging approach. °  $p<0.1$ ; \*  $p<0.05$ ; \*\*  $p<0.01$ ; \*\*\*  $p<0.001$ .

	model 1	model 6 – grassland
Crop diversity (CD)	-0.03 [-2.07 ; 2.01]	0.18 [-1.9 ; 2.26]
Mean field size (MFS)	-6.39 [-11.85 ; -0.94] *	-6.2 [-11.83 ; -0.59] *
Semi-Natural Cover (SNC)	5.07 [2.87 ; 7.26] ***	5.07 [2.88 ; 7.27] ***
Nb of Crops sampled	2.84 [1.07 ; 4.62] ***	2.73 [0.94 ; 4.52] **
Latitude	1.5 [-3.55 ; 6.55]	
Longitude	3.73 [2.47 ; 9.93]	4.07 [-2.34 ; 10.47]
MFS <sup>2</sup>	3.78 [-0.67 ; 8.23] °	3.98 [-0.48 ; 8.44] °
CD :SNC	2.20 [0.64 ; 3.76] **	2.25 [0.69 ; 3.81] **
MFS :SNC	1.15 [-0.66 ; 2.96]	1.33 [-0.51 ; 3.16]
Grassland		1.87 [-0.26 ; 4.00] °

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### 5.5.3. Role of semi-natural vegetation occurring between fields

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Mean field size (MFS in ha) and the length of semi-natural linear elements between fields (SNL) or the length of hedgerows (H) were strongly correlated, particularly in some regions (e.g. Armorique, Table S5.6). As a result, we could not include both MFS and SNL (or MFS and H) in our models and disentangle their effects on multitrophic diversity.



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**Table S5.6.** Pearson correlation coefficients among explanatory variables across and within regions. CD = crop diversity, MFS = mean field size, SNC= proportion of semi-natural cover, SNL= length of semi-natural linear elements between fields, H = length of hedgerows between fields. N = number of landscapes. Correlations between H and CD or SNC were low and are not shown here for simplicity.

	CD-MFS	CD-SNC	CD-SNL	MFS-SNC	MFS-SNL	MFS-H	SNC-SNL	N
All regions	-0.13	-0.27	-0.30	-0.02	-0.44	-0.37	0.13	435
Armorique	-0.03	0.09	0.10	-0.01	-0.71	-0.67	-0.06	40
Camargue	-0.20	-0.25	0.11	-0.06	-0.55	-0.17	-0.59	40
Coteaux	-0.27	-0.22	0.51	-0.31	-0.57	-0.50	-0.24	32
East Anglia	-0.18	0.21	0.18	-0.16	-0.34	-0.23	-0.41	60
Goettingen	-0.17	0.15	0.05	0.15	-0.43	-0.10	-0.10	52
Lleida	-0.40	-0.14	0.16	-0.15	-0.50	-0.23	-0.20	40
Eastern Ontario	-0.34	-0.13	0.27	-0.40	-0.53	-0.43	-0.08	93
PVDS	-0.16	-0.08	-0.02	-0.37	-0.51	-0.57	0.29	78

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To test whether our results for MFS were likely due to the correlation with SNL or H, we selected a subset of landscapes for which explanatory variables, in particular MFS and SNL as well as MFS and H, were uncorrelated i.e. with a Pearson correlation coefficient <0.56 for each pair of explanatory variables, within each region (Table S5.7).

**Table S5.7.** Pearson correlation coefficients among explanatory variables, across and within regions, within the subset of landscapes (274 landscapes) used to test for the influence of SNL and H on our results for the effects of crop heterogeneity. CD = crop diversity, MFS = mean field size, SNC= proportion of semi-natural cover, SNL= length of semi-natural linear elements between fields, H = length of hedgerows between fields. N = number of landscapes.

	CD-MFS	CD-SNC	CD-SNL	MFS-SNC	MFS-SNL	MFS-H	SNC-SNL	N
All regions	-0.15	-0.30	-0.40	-0.08	-0.27	-0.28	0.30	274
Armorique	-0.02	0.29	0.40	-0.06	-0.04	-0.15	-0.33	20
Camargue	-0.25	-0.19	-0.14	-0.56	-0.05	-0.15	-0.09	20
Coteaux	0.31	-0.38	0.20	-0.46	0.06	-0.12	-0.52	20
East Anglia	-0.15	-0.04	0.35	-0.32	-0.18	-0.31	-0.40	43
Goettingen	-0.26	0.10	0.10	-0.02	-0.22	-0.01	-0.07	45
Lleida	-0.33	0.08	-0.51	-0.37	0.24	-0.20	0.08	20
Eastern Ontario	-0.18	-0.07	-0.03	-0.43	-0.21	-0.32	-0.32	44
PVDS	-0.16	-0.15	-0.08	-0.41	-0.28	-0.46	0.29	62

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We built a model similar to model 1 including both SNL and MFS in order to disentangle their effects on multitrophic diversity:

*Model 7: lmer (T60.landscape ~ CD \* MFS \* SNC + CD<sup>2</sup> + MFS<sup>2</sup> + SNC<sup>2</sup> + CropNb + Lat + Lon + SNL + (1|Region/Year))*

Model selection based on model 7 included the same variables as for model 1 (except Latitude and SNC<sup>2</sup>), plus SNL. SNL was marginally significant. Parameter estimates and significance for variables of interest remained unchanged (Table S5.8). This results does not confirm the general assumption that the positive effect of MFS is only due to the positive effect of the amount of SNL.

Our variable SNL included a variety of semi-natural linear elements (e.g. hedgerows, grassy margins) that may not play the same role for biodiversity. Therefore, we built another model similar

1539 to model 7 including the length of hedgerows (Hedgerow) instead of SNL in order to test whether the  
 1540 effect of MFS on multitrophic diversity may be due to the increase in the length of hedgerows:

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 1542 *Model 8: lmer (T60.landscape ~ CD \* MFS \* SNC + CD<sup>2</sup> + MFS<sup>2</sup> + SNC<sup>2</sup> + CropNb + Lat + Lon +*  
 1543 *Hedgerows + (1 | Region/Year))*

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 1545 Model selection based on model 8 included the same variables as for model 1 (except SNC<sup>2</sup>  
 1546 and MFS:SNC), plus Hedgerows. Hedgerows were non-significant. Parameter estimates and  
 1547 significance for variables of interest remained unchanged (Table S5.8). This results does not confirm  
 1548 the general assumption that the positive effect of MFS is only due to the positive effect of the  
 1549 amount of SNL or hedgerows. Instead, this result lends support to the idea that agricultural  
 1550 landscapes with smaller fields provide better access to different field types for species that require  
 1551 landscape complementation.

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 1553 **Table S5.8.** Comparison of models 1, 7 (with SNL) and 8 (with Hedgerows) based on the uncorrelated  
 1554 subset of landscapes. Parameter listed are those retained in the model selection procedure.  
 1555 Parameter estimates and confidence intervals are based on the model averaging approach. ° p<0.1; \*  
 1556 p<0.05; \*\* p<0.01; \*\*\* p<0.001.

	model 1 (subset)	model 7 – SNL	model 8 – Hedgerows
Crop diversity (CD)	-0.14 [-2.9 ; 2.62]	0.39 [-2.39 ; 3.17]	-0.03 [-2.8 ; 2.74]
Mean field size (MFS)	-9.9 [-18.1 ; -1.68] *	-8.92 [-17.24 ; -0.61] *	-8.28 [-16.94 ; 0.38] °
Semi-Natural Cover (SNC)	3.09 ; 0.15 ; 6.03] *	3.16 [0.25 ; 6.07] *	3.17 [0.21 ; 6.14] *
Latitude		2.94 [-3.03 ; 8.9]	
Longitude	2.61 [-2.01 ; 8.89]	2.06 [-4.5 ; 8.62]	2.74 [-4.1 ; 9.58]
MFS <sup>2</sup>	6.71 [-0.07 ; 13.49] °	6.54 [-0.16 ; 13.24] °	6.33 [-0.44 ; 13.11] °
SNC <sup>2</sup>		2.71 [0.14 ; 5.34] *	2.6 [-0.03 ; 5.24] °
Nb of Crops sampled	3.87 [1.58 ; 6.17] ***	4.28 [1.98 ; 6.58] ***	3.86 [1.57 ; 6.15] **
CD :SNC	1.85 [-0.28 ; 3.98] °	1.79 [-0.31 ; 3.89] °	1.83 [-0.29 ; 3.96] °
MFS :SNC	0.66 [-2.01 ; 3.32]	0.83 [-1.81 ; 3.47]	
SNL		3.64 [-0.06 ; 7.34] °	
Hedgerows			2.69 [-0.22 ; 5.56] °

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## 1559 5.6 Correlations and alternative mechanisms at the field level

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1561 Crop diversity and mean field size are also likely to be correlated with several variables at the field  
 1562 level, including the identity of crops sampled, the local land-use intensity (e.g. herbicide use,  
 1563 ploughing frequency). Disentangling the role of crop heterogeneity from the effects of these other  
 1564 variables is also necessary in order to infer potential mechanisms explaining the positive effect of  
 1565 crop heterogeneity on multitrophic diversity. This required running models at the field level, using a  
 1566 data subset for which co-variable data were available. As a result, we could not include all these  
 1567 variables in a single model and therefore present these analyses as separate, complementary  
 1568 analyses.

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### 1570 5.6.1. Role of the identity of sampled crop types

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1572 We tried to limit correlations between the two crop heterogeneity gradients and the identity  
 1573 of sampled crop types. In some cases, correlations were impossible to avoid because some crops  
 1574 occurred or were dominant only in some regions (e.g. rice in Camargue, almond and olive in Lleida)  
 1575 or some landscapes (e.g. landscapes with low crop compositional heterogeneity). As a result,  
 1576 different types of crop sampled were associated with significantly different values of crop diversity or  
 1577 mean field size (Table S5.9).

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**Table S5.9.** Analysis of variance showing the relationship between the two heterogeneity gradients (crop diversity and mean field size) and sampled crop type within each region. Since sampled crop type is a categorical variable, correlation coefficient cannot be used. We therefore used the function aov in R, crop diversity and mean field size being the response variables and sampled crop type being the predictor variable. Values correspond to the F value of the function aov in R. \* p<0.05; \*\* p<0.01; \*\*\* p<0.001.

	Crop diversity	Mean field size
All regions	5.78***	9.28***
Armorique	1.95	0.29
Camargue	8.54**	0
Coteaux	1.16	0.59
East Anglia	3.35***	1.29
Goettingen	0	0
Lleida	9.43***	2.18
Eastern Ontario	2.57*	2.61**
PVDS	0.35	0.53

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To evaluate whether the sampled crop type influenced our results, we built a model similar to model 1 but using multidiversity calculated at the field level as the response variable (T60.field). We compared models with and without adding crop type as a random effect (using data collected in 1305 fields in 435 landscapes from 8 regions). Crop type was added as a random effect because we were not interested in estimating the specific effect of each particular crop type. Note there were enough crop types (16) to estimate the random effect adequately.

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*Model 9: lmer (T60.field ~ CD \* MFS \* SNC + CD<sup>2</sup> + MFS<sup>2</sup> + SNC<sup>2</sup> + Lat + Lon + (1 | Region/Year/Landscape))*

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*Model 10: lmer (T60.field ~ CD \* MFS \* SNC + CD<sup>2</sup> + MFS<sup>2</sup> + SNC<sup>2</sup> + Lat + Lon + (1 | Region/Year/Landscape) + (1 | Crop type))*

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To test whether crop type had a significant effect on field-level multitrophic diversity, we used a restricted likelihood-ratio test based on simulated values from the finite sample distribution available in the function exactRLRT from package RLRsim. We then compared the estimates and p-values associated with models 9 and 10 to determine whether any effects of crop type influenced our conclusions regarding the effects of crop heterogeneity on multitrophic diversity.

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Although we detected a significant effect of crop type on field-level multitrophic diversity (RLRT = 125.43, p-value < 0.001), adding crop type as a random effect in the model did not change the outcome of model selection or the significance of variables of interest (Table S5.8). This result suggests that variations in the identity of crops sampled do not explain the effects of CD, in combination with SNC, and MFS on multitrophic diversity detected in our study.

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**Table S5.10.** Comparison of models built at the field level for multitrophic diversity (model 9 – field level, i.e. without sampled crop type as a random effect; model 10 – sampled crop id, i.e. with sampled crop type as a random effect). Parameter listed are those retained in the model selection procedure. Parameter estimates and confidence intervals are based on the model averaging approach. ° p<0.1; \* p<0.05; \*\* p<0.01; \*\*\* p<0.001.

	model 9 (field)	model 10 (field) – sampled crop ID
Crop diversity (CD)	0.78 [-0.79 ; 2.36]	0.25 [-2.08 ; 2.58]
Mean field size (MFS)	-3.14 [-6.57 ; 0.28] °	-2.44 [-4.77 ; -0.10] *
Semi-Natural Cover (SNC)	3.14 [-1.12 ; 7.4]	3.79 [0.98 ; 6.60] **
Latitude	0.97 [-3.4 ; 5.33]	
Longitude	3.63 [-1.68 ; 8.93]	1.2 [-4.88 ; 7.28]
CD <sup>2</sup>		0.67 [-4.25 ; 5.6]
MFS <sup>2</sup>	2.07 [-1.52 ; 5.66]	1.19 [-2.38 ; 4.76]
SNC <sup>2</sup>	2.9 [-1.27 ; 7.06]	2.05 [-2.08 ; 6.18]
CD :SNC	1.35 [0.08 ; 2.63] *	1.39 [0.14 ; 2.63] *
MFS :SNC	1.55 [0.09 ; 3.00] *	1.91 [0.47 ; 3.34] **
CD :MFS		0.2 [-1.12 ; 5.56]

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### 5.6.2. Role of crop heterogeneity in cereal fields

To further assess the role of crop identity, we applied model 9 to the subset of data collected in cereal fields. Indeed, cereal is the most widespread crop type sampled in our dataset and the only one present in all regions. We therefore applied model 6 on 615 fields in 334 landscapes in our 8 regions (after removing the random effect of landscape since most landscape contain only one cereal field). This analysis confirms that decreasing MFS and, when SNC is high enough, increasing CD have positive effects on multitrophic diversity in cereal crop fields (Table S5.11).

**Table S5.11.** Comparison of models built at the field level for multitrophic diversity (model 9) with the complete dataset and with the cereal subset. Parameter listed are those retained in the model selection procedure. Parameter estimates and confidence intervals are based on the model averaging approach. ° p<0.1; \* p<0.05; \*\* p<0.01; \*\*\* p<0.001.

	model 9 (field) – complete dataset	model 9 (field) – cereal subset
Crop diversity (CD)	0.78 [-0.79 ; 2.36]	-2.78 [-8.62 ; 3.06]
Mean field size (MFS)	-3.14 [-6.57 ; 0.28] °	-4.51 [-9.24 ; 0.23] °
Semi-Natural Cover (SNC)	3.14 [-1.12 ; 7.4]	3.16 [0.26 ; 6.06] *
Latitude	0.97 [-3.4 ; 5.33]	
Longitude	3.63 [-1.68 ; 8.93]	2.03 [-0.87 ; 4.94]
MFS <sup>2</sup>	2.07 [-1.52 ; 5.66]	3.62 [-0.19 ; 7.43] °
SNC <sup>2</sup>	2.9 [-1.27 ; 7.06]	1.49 [-3.09 ; 6.08]
CD :SNC	1.35 [0.08 ; 2.63] *	1.76 [0.17 ; 3.36] *
MFS :SNC	1.55 [0.09 ; 3.00] *	3.31 [1.73 ; 4.9] ***
CD :MFS		0.46 [-1.17 ; 2.09]

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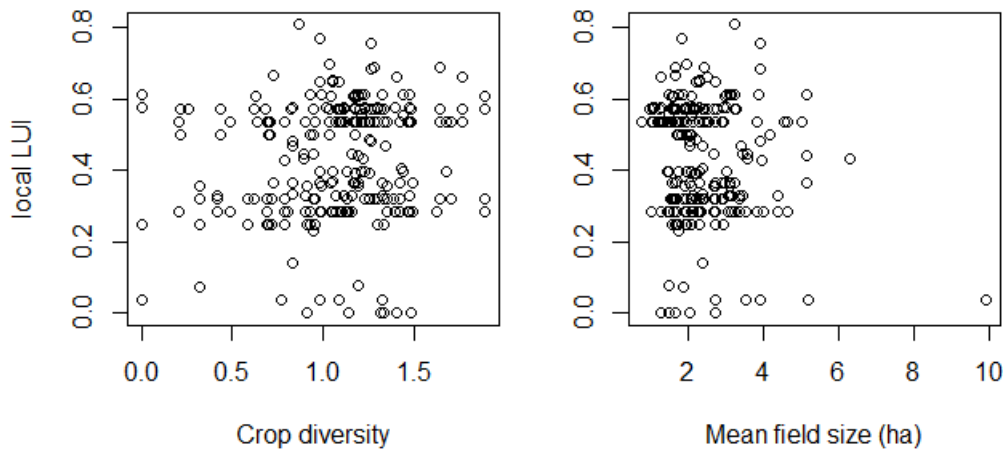
### 5.6.3. Role of field-level Land-Use Intensity

Land-use intensity may be correlated with crop heterogeneity in some regions. For instance, landscapes with larger mean field sizes may be associated with higher fertilizer inputs (Levers et al. 2016, Roschewitz et al. 2005). Such correlations could hamper our ability to draw conclusion on the effects of crop heterogeneity on multitrophic diversity.

We conducted farmer surveys to collect data on land use intensity of the sampled fields. Information included ploughing (0=no/1=yes), use of fertilizer (0=no/1=yes), frequency of herbicide use (from 0 to 7) and frequency of insecticide use (from 0 to 6) in 324 fields located in 132 landscapes across five regions (Armorique, Camargue, Coteaux, Goettingen and Eastern Ontario). We calculated a local Land-Use Intensity index (local LUI) based on the normalized mean of these four variables (after scaling each variable) following a formula similar to the one developed by Herzog et

1648 al. (2006):  $LUI = \frac{1}{4} (\text{scale(ploughing)} + \text{scale(fertilizer)} + \text{scale(herbicide)} + \text{scale(insecticide)})$ . This local  
 1649 LUI index therefore varies between 0 (low intensity) and 1 (high intensity).

1650 The Pearson correlation between local LUI and crop diversity was weak and not significant  
 1651 ( $r=0.10$ ;  $p=0.12$ ). The Pearson correlation between local LUI and mean field size was negative (i.e.  
 1652 opposite to expectation;  $r= -0.27$ ;  $p<0.001$ ; Fig. S5.4).  
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 1655 **Figure S5.4.** Relationship between the two crop heterogeneity gradients and Land-Use Intensity  
 1656 (LUI).  
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1658 We added local LUI to model 10 and compared the outcomes of model 10 and model 11  
 1659 using the data subset for which Field LUI data was available.  
 1660

1661 *Model 11:  $lmer(T60.field \sim CD * MFS * SNC + CD^2 + MFS^2 + SNC^2 + Lat + Lon + Field\ LUI + (1|$*   
 1662 *Region/Year/Landscape) + (1|Crop type))*

1663  
 1664 Model selection based on model 11 included almost the same variables as for model 10, plus  
 1665 Field LUI, which had a significant negative effect. Parameter estimates for model 10 using the LUI  
 1666 data subset differ slightly from parameter estimates due to the fact that more complex interactions  
 1667 were included. However, we checked that the overall shape of the relationships do not differ much  
 1668 between the model based on the whole dataset and the model based on the LUI dataset. More  
 1669 importantly, parameter estimates and significance for other variables of interest remained very  
 1670 similar between model 10 and model 11 (Table S5.12). This result suggests that the effects of mean  
 1671 field size and crop diversity cannot be explained by variations in field-level land-use intensity. It is  
 1672 interesting to note that we observe here a significant negative interaction between crop diversity  
 1673 and mean field size which is consistent with the ‘landscape complementation’ hypothesis, i.e. the  
 1674 fact that multitrophic diversity benefit more from increasing crop diversity when fields become  
 1675 smaller and can be reached more easily. However, the fact that this relationship was not observed in  
 1676 other models calls for further investigations.  
 1677

1678 **Table S5.12.** Comparison of models built at the field level for multitrophic diversity with and without  
 1679 field-level land use intensity (LUI). Parameter listed are those retained in the model selection  
 1680 procedure. Parameter estimates and confidence intervals are based on the model averaging  
 1681 approach. °  $p<0.1$ ; \*  $p<0.05$ ; \*\*  $p<0.01$ ; \*\*\*  $p<0.001$ .  
 1682

	model 10 (field level LUI subset)	model 11 (field level LUI subset) - LUI
Crop diversity (CD)	18.1 [5.35 ; 20.85] **	16.14 [3.42 ; 28.86] *
Mean field size (MFS)	8.81 [0.31 ; 17.31] *	8.32 [-0.41 ; 17.05] °

Semi-Natural Cover (SNC)	17.69 [6.26 ; 29.12] **	19.11 [7.9 ; 30.33] ***
Latitude	4.38 [0.95 ; 7.82] *	5.91 [1.72 ; 10.09] **
Longitude	2.98 [-0.19 ; 6.15] °	
CD <sup>2</sup>	-15.54 [-27.25 ; -3.83] **	-14.25 [-25.88 ; -2.61] *
MFS <sup>2</sup>	-12.27 [-21.8 ; -2.7] *	-13.33 [-22.78 ; -3.88] **
SNC <sup>2</sup>	-15.76 [-27.97 ; -3.54] *	-17.9 [-29.89 ; -5.91] **
CD :SNC	-4.8 [-8.53 ; -1.06] *	-5.2 [-8.86 ; -1.55] **
MFS :SNC	2.55 [-0.77 ; 5.86]	
CD :MFS	-4.06 [-7.55 ; -0.57] *	-3.8 [-6.71 ; -0.87] *
CD :MFS :SNC	1.6 [-0.99 ; 4.19]	
Field LUI		-2.53 [-4.79 ; -0.26] *

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### 5.7. Moving window modeling approach for Crop heterogeneity × Semi-natural cover interaction

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We used a moving window modeling approach (Humpries et al. 2010; Berdugo et al. 2018) to identify potential discontinuities in the response of multitrophic diversity measured at the landscape level (T60.landscape) to crop diversity and mean field size along the gradient of semi-natural cover. To do so, we ordered all landscapes (n = 435) along the gradient of semi-natural cover (%) and selected the first 75 landscapes with the lowest semi-natural cover. Using this subset, we ran the model obtained from the averaging approach applied to model 1 (Fig. 2A main text) after excluding semi natural cover and its interactions with CD and MFS, such as:

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*Model 12: lmer (T60.landscape ~ CD\*MFS + MFS<sup>2</sup> + CropNb + Lat + Lon + (1 | Region/Year))*

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We then extracted and stored the model coefficient for crop diversity (CD), mean field size (MFS) and the confidence intervals (CIs). We then removed the landscape with the lowest value of semi-natural cover from the subset of 75 landscapes, added the landscape scoring the next higher value, ran model 12 and extracted model coefficients and CIs. We repeated this loop as many times as landscapes remained along the entire gradient of semi-natural cover (n = 286 subsets, see R code below). We saved all coefficients and confident intervals for each step and plotted them against the gradient of semi-natural cover (Fig. S5.5).

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Consistently with our multiple regression analyses (Fig. 2A in main text), this moving window analysis showed that the effect of crop diversity and mean field size on multitrophic diversity changes along the gradient of semi-natural cover (Fig. S5.5 A and B). The effect of crop diversity is positive for high values of semi-natural cover, neutral as semi-natural cover decreases and negative for the low values of semi-natural cover. The effect of mean field size is neutral for the high values of semi-natural cover and negative for low values of semi-natural cover.

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However, this analysis reveals that changes in the effect of crop diversity and mean field size on multitrophic diversity are not smooth but instead show abrupt transitions when semi-natural cover decreases. For crop diversity, there is an abrupt change at 11.2% of semi-natural cover where the effect of crop diversity shifts abruptly from positive to neutral and one at 4.5% where the effect of crop diversity shifts from neutral to negative. For mean field size, there is one abrupt change at 8% where the effect of mean field size shifts abruptly from neutral to negative. This analysis allows identifying three thresholds that can be used to guide recommendations on how to manage the three main components of agricultural landscape heterogeneity, namely crop diversity, mean field size and the amount of semi-natural cover (see main text for more details).

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R Code for the Moving Window Analysis (the code provided only concerns crop diversity)

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```

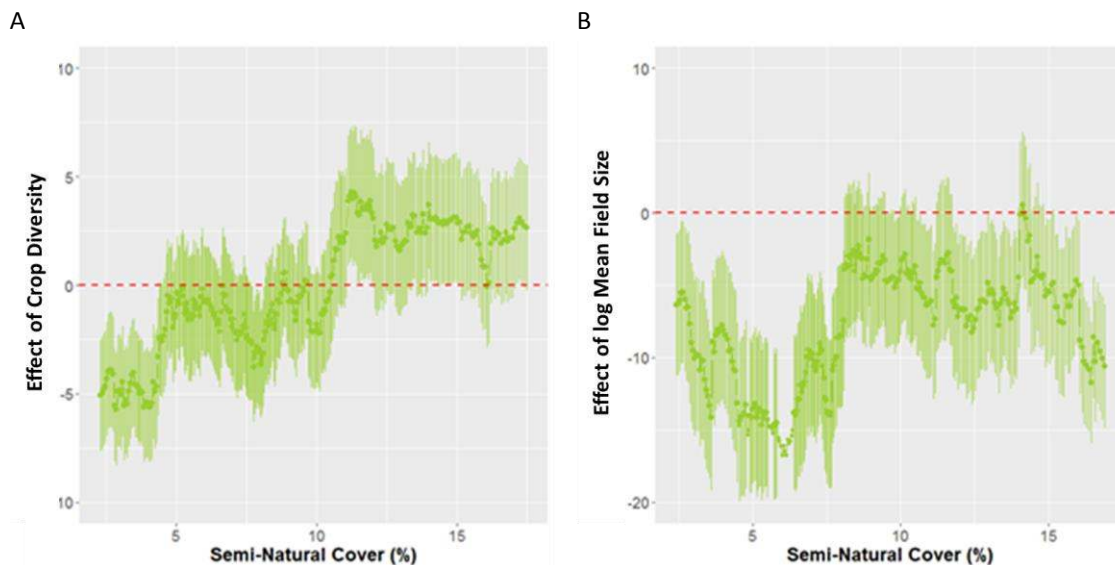
1724 ##### moving window function
1725 WindowSKR <- function(df,Factor,X,Y,formul,n=10){
1726   myvars<-c(Factor,X,Y)
1727   dftemp = df[myvars]
1728   dftemp = dftemp[order(dftemp[Factor]),]
1729   tt=length(unlist(dftemp[Factor]))-n
1730   i = 1
1731   mdl <- lmer(data = dftemp, formula = formul)
1732   res<- matrix(data = NA,nrow = 1,ncol = length(fixef(mdl))+1)
1733   ci<-res
1734   library(lme4)
1735   while(tt>n){
1736     dfi <- dftemp[i:(i+n),]
1737     Fact <- mean(unlist(dfi[Factor]))
1738     mdl <- lmer(data = dfi, formula = formul, na.action = na.fail,REML ="TRUE")
1739     #dist<- mean(unlist(dfi[X]))+1-mean(unlist(dfi[Y]))
1740     res <- rbind(res,c(Fact,fixef(mdl)))
1741     cii <- (abs(confint(mdl)[-c(1,2),1]-confint(mdl)[-c(1,2),2]))/2
1742     ci<-rbind(ci,c(Fact,cii))
1743     tt=tt-1
1744     i=i+1
1745   }
1746   res<- as.data.frame(res)
1747   ci<-as.data.frame(ci)
1748   colnames(res)<-c("MWfactor",names(fixef(mdl)))
1749   colnames(ci)<-c("MWfactor",names(fixef(mdl)))
1750   RES<-list(res=res,ci=ci)
1751   return(RES)
1752 }
1753
1754 ##### uploading libraries
1755 library(jsonlite)
1756 library(ggplot2)
1757 library(tidyr)
1758 library(boot)
1759 library(lme4)
1760
1761 ##### running moving window analysis
1762 formul<-T60.landscape~ Crop_SHDI+Crop_MFS + sampled.crop.nb + MFS2 + Lon + Lat + (1|Region/Year) -1
1763 RES <- WindowSKR(df,"Seminat_Cover",c("Crop_SHDI","MFS2","Crop_MFS", "Seminat_Cover",
1764 "sampled.crop.nb", "Region", "Year", "Lon", "Lat"),"T60.landscape",formul,n=75)
1765
1766 ##### plotting results of the moving window analysis
1767 dfres=data.frame(MWfactor<-RES$res$MWfactor, Effect<-RES$res$Crop_SHDI, CI<-RES$ci$Crop_SHDI)
1768 limits <- aes(ymax = Effect + CI, ymin=Effect - CI)
1769 p1<-ggplot(data = dfres,aes(x = MWfactor,y = Effect), ylim = c(1,4))+
1770   geom_line(col = "olivedrab3")+
1771   geom_point(col = "olivedrab3")+
1772   geom_pointrange(limits,col = "olivedrab3")+
1773   xlab("Semi-Natural Cover (%)"+

```

```

1774 ylab("Effect of Crop Diversity")
1775 p1 + theme(axis.text=element_text(size=14), axis.title.x = element_text(size=18, face="bold"), axis.title.y =
1776 element_text(size=18, face="bold"))
1777
1778

```



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1781 **Figure S5.5.** Effect of crop diversity (A) and mean field size (B) on multitrophic diversity for different levels of  
1782 semi-natural cover. Parameter estimates and confidence intervals are based on a moving window analysis (see  
1783 detailed description in SI5). The red line indicates a null effect. Each dot and CI correspond to the estimate  
1784 values of CD or MFS for the average semi-natural cover of a given window along the semi-natural cover  
1785 gradient. Due to the low number of landscapes with semi-natural cover >17.5% (Table S4.1), we only represent  
1786 the gradient between 0 and 17.5% of semi-natural cover on these figures.

1787  
1788 **References**

1789 Berdugo M, et al. Aridity preferences alter the relative importance of abiotic and biotic drivers on  
1790 plant species abundance in global drylands. *J Ecol* 0(0). doi:10.1111/1365-2745.13006.  
1791 Herzog F, et al. (2006) Assessing the intensity of temperate European agriculture at the landscape  
1792 scale. *Eur J Agron* 24(2):165–181.  
1793 Humphries NE, et al. (2010) Environmental context explains Lévy and Brownian movement patterns  
1794 of marine predators. *Nature* 465(7301):1066–1069.  
1795 Levers C, Butsic V, Verburg PH, Müller D, Kuemmerle T (2016) Drivers of changes in agricultural  
1796 intensity in Europe. *Land Use Policy* 58(Supplement C):380–393.  
1797 Öckinger E & Smith HG (2007) Semi-natural grasslands as population sources for pollinating insects in  
1798 agricultural landscapes. *Journal of applied ecology* 44(1): 50-59.  
1799 Roschewitz I, Thies C, Tscharntke T (2005) Are landscape complexity and farm specialisation related  
1800 to land-use intensity of annual crop fields? *Agric Ecosyst Environ* 105(1–2):87–99.  
1801