



## Think twice before using narrow buffers: Attenuating mowing-induced arthropod spillover at forest – grassland edges



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### ABSTRACT

Spillover between agricultural land and natural habitats is recognised as an important mechanism shaping biodiversity and ecosystem functions. Its spatio-temporal patterns and magnitude are thoroughly described in the literature and it is often stated that spillover should be considered in conservation planning. In fact studies that implement and test active interventions to modulate spillover are scarce. Therefore, we studied the spillover of spiders and carabids between hay meadows and natural forests after mowing and tested whether leaving unmown buffer strips in the edges can mitigate undesirable aspects of mowing-induced spillover. We found that mowing affected the assemblages both in the meadows and forests and, interestingly, changes were more profound in the forests. Mowing reduced the spillover of forest assemblages into meadows. Mowing also led to the retraction of forest assemblages from the peripheral zones of forests but did not trigger an influx of meadow assemblages into the forests. Wide (10 m) unmown buffers attenuated or completely offset most of these effects. Leaving narrow (5 m) buffers had unexpected consequences, as they did not function only as buffers but as facilitators of forest-ward spillover from meadows, potentially compromising ecological interactions such as predation or competition in forests. We conclude that using wide buffers can be recommended as a refinement of standard management practices in hay meadow–forest mosaics. Narrow buffers should be applied with great caution and should generally be avoided if the forest-specific assemblages are of conservation interest.

### 1. Introduction

Semi-natural grasslands are decreasing in area and naturalness across Europe due to management intensification, abandonment and transformation to other land use types (Báldi et al., 2013; Dengler et al., 2014). Best management practices for the preservation of remaining grasslands include extensive grazing with various livestock and/or mowing once or a few times a year (Tälle et al., 2016; Török et al., 2016). Extensive grazing with appropriate livestock may mimic pre-historical disturbance regimes of wild megaherbivores (Vera, 2002; Sutherland, 2002) and can create heterogeneous vegetation (Marty, 2005; Tölgyesi et al., 2015), which supports high arthropod diversity (Lambert et al., 2007; Woodcock et al., 2006). Conversely, mowing, which is nowadays performed by high-power machines, leads to more homogenous habitat structure and arthropod assemblages (Cizek et al., 2012). Mowing machines also mechanically kill a large number of animals from insects (Thorbeck and Bilde, 2004) to birds (Vadász and Lóránt, 2015). The physical environmental conditions on freshly mown meadows may be beyond the limit of tolerance of the survivors and they

are also more visible to predators (Opatovsky and Lubin, 2012). To counteract the negative effects of mowing, several recommendations have been proposed in the frame of agri-environmental schemes (AES). These include the careful timing of mowing, increased cut height, the combined application of mowing and grazing, and leaving unmown strips or patches, where animals can find shelter and plants can set seed (Buri et al., 2013; Cizek et al., 2012; Humbert et al., 2012a,b; Lebeau et al., 2015; van Klink et al., 2017).

All recommendations listed above aimed to mitigate the negative effect of mowing on grassland biodiversity. However, processes in one habitat can affect the peripheral zones of adjacent habitats through edge effects (Murcia, 1995). Accordingly, the mowing of grasslands can affect the biota of the neighbouring habitats. These effects should be taken into account, particularly if these habitats are of high conservation value (such as forest fragments) and/or if the grassland and non-grassland patches form a natural mosaic or a fragmented landscape. In such landscapes the edge to patch interior ratio is high, thus the potentially affected proportion of non-grassland habitats is also high (Cook et al., 2002; Madeira et al., 2016).

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A frequently studied manifestation of edge effects is the spillover of arthropods, i.e. their periodic/annual movement or foraging across habitat edges (Rand et al., 2006, Tschardt et al., 2012). Spillover can be traced back to various ecological reasons. Differences in productivity can lead to the movement of organisms from a productive habitat to a less productive neighbouring habitat in a passive, diffusion-like manner, corresponding to the source-sink dynamics described by Dunning et al. (1992). Conversely, an abrupt decline in habitat quality can trigger active emigration from a patch into a neighbouring one (Holt and Hochberg, 2001; Thorbek and Bilde, 2004). Landscape complementation (i.e. the need of an organism for both neighbouring habitats to complete its life-cycle) and landscape supplementation (i.e. if the neighbouring habitats provide alternatives for certain resources) can also explain spillover (Dunning et al., 1992; Fahrig et al., 2011; Tschardt et al., 2012). Spillover has been detected between various habitat pairs such as adjacent arable fields and natural grasslands (Madeira et al., 2016; Rand and Louda, 2006), natural forests and forest plantations (Lucey and Hill, 2012), grasslands and coniferous forests (Lacasella et al., 2015) and even between fields of different crop types (Duflo et al., 2016; Macfadyen and Muller, 2013), indicating that it is a wide-spread phenomenon. The relevance of spillover for conservation issues is emphasised in most of these studies (e.g. Pryke and Samways, 2012; Schneider et al., 2013), as they claim that the influx of arthropods from the neighbouring habitats can import new ecosystem functions (Blitzer et al., 2012; Rand et al., 2006) but can disrupt some functions by increasing competition or predation (Ries and Sisk, 2004).

In relation to forest-grassland edges it has been found that spillover of forest assemblages into grasslands is more pronounced than spillover of grassland assemblages into forests (Boetzel et al., 2016; Lacasella et al., 2015) as the higher biomass of forests can maintain a higher abundance of arthropods, which then spillover into grasslands according to the source-sink dynamics (Dunning et al., 1992). Mowing, however, causes a rapid deterioration of habitat quality in the grassland, actively driving arthropods out of the affected habitats (Eyre et al., 2013; Opatovsky and Lubin, 2012). Thus, we expect that forest assemblages will show an abrupt decline in their spillover rate into freshly mown grasslands and grassland assemblages will likely be seeking shelter in adjacent forest patches, leading to increased spillover into forests immediately after mowing. Although some seasonal landscape complementation or supplementation can exist between the forests and meadows (Dunning et al., 1992; Fahrig et al., 2011), the effects of modern mowing techniques may cause perturbations in spillover that were unprecedented in earlier times. Thus, there is a need to assess the outlined cross-edge effects of mowing and measures should be taken to offset them if necessary. To date, little effort has been made to address this issue. We therefore designed a field experiment in hay meadow-hardwood grove mosaics in Hungary to study the changes of arthropod assemblages in the peripheral zones of adjacent meadow and forest patches immediately after mowing. We chose spiders and carabid beetles as test organisms because they usually have distinct assemblages in forest and grassland habitats (Noreika and Kotze, 2012), are mobile enough to cross edges, are sensitive to environmental variation at small spatial scales, and occupy an intermediate trophic level, which allocates a key role to them in the maintenance of ecosystem functions (Lacasella et al., 2015; Pearce and Venier, 2006). We focussed only on the peripheral zones of forests and grasslands because spillover is expected to be more pronounced there than farther away from the forest edges (Boetzel et al., 2016). Specifically, we asked the following questions:

- (1) Does mowing alter the species composition of spider and carabid assemblages in the peripheral zones of forests and meadows, and can these effects be prevented by leaving unmown buffer strips?
- (2) Does mowing cut back on the spillover of forest assemblages, and can it be restored by buffer strips?
- (3) Does mowing trigger a quick influx of meadow specific

assemblages, and can this be attenuated by buffer strips?

## 2. Material and methods

### 2.1. Study area

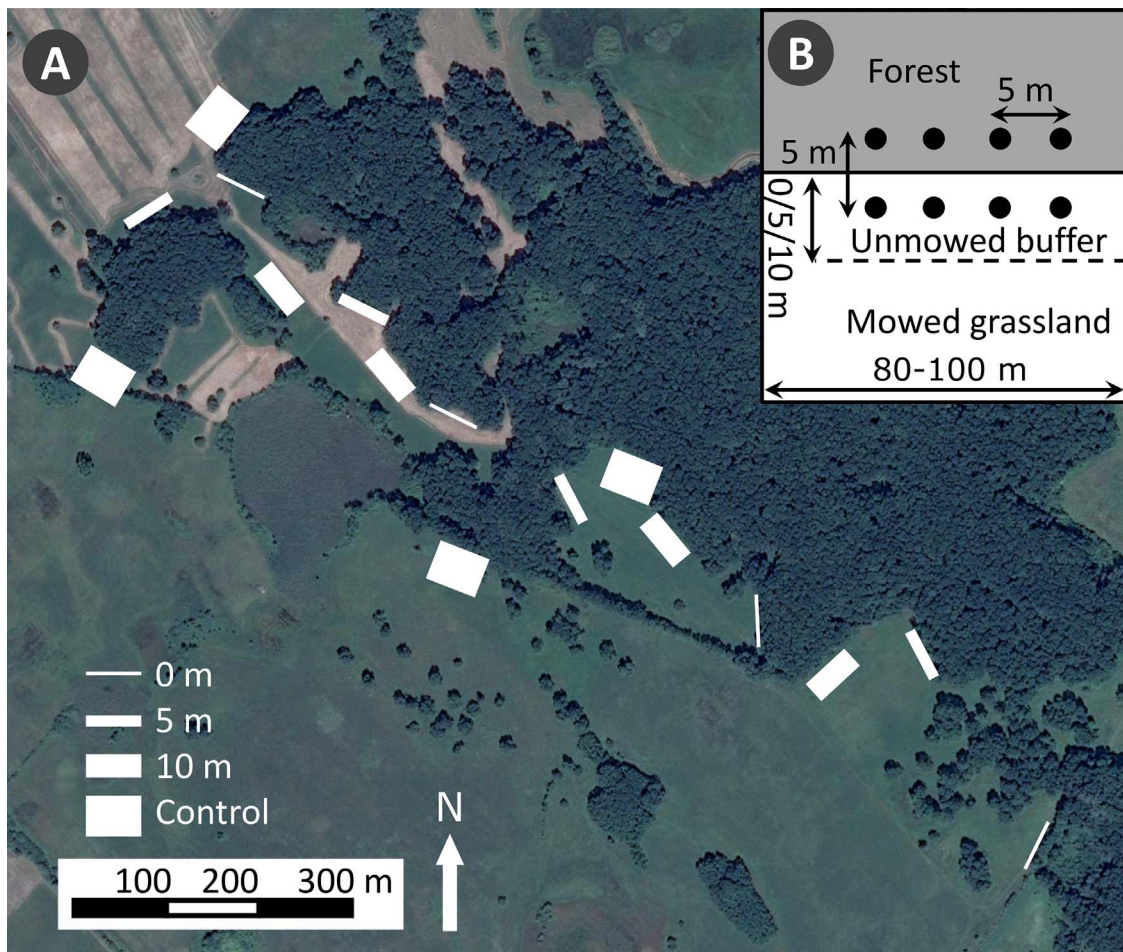
We studied a hay meadow-hardwood grove mosaic in the Kiskunság National Park, central Hungary (N46.725 E19.347, 98 m a.s.l.). The climate is continental with a sub-Mediterranean influence; the annual precipitation is 500–600 mm and the mean annual temperature is 10–11 °C (Tölgyesi et al., 2016). Extra water, supplied by groundwater seeping from the adjacent Danube-Tisza Sandy Ridge maintains a vegetation type with high water demand. Hay meadows are characterised by tall grasses, such as *Molinia caerulea* and *Deschampsia caespitosa* and form a mosaic with forest patches, characterised by *Fraxinus angustifolia* subsp. *danubialis* and *Quercus robur*. Both forests and meadows provide habitat for several plants (e.g. *Gladiolus palustris*), invertebrates (e.g. *Isophya costata* and *Phengaris teleius*) and birds (e.g. *Crex crex* and *Haliaeetus albicilla*) of community interest in the European Union. Both habitats are also listed in the Habitats Directive (European Union 1992). The forest patches are part of a strict forest reserve, in which forestry activities have been banned for 60 years. The meadows are used for extensive hay production and are mown once a year, mostly after mid-July to avoid damage to ground-nesting birds. Some strips or larger blocks (10–15% of the total area) are left unmown every year for wildlife refuge and to allow the seed production of late summer plants. Meadows are usually mown up to the edges of the forests. In some years in winter, shrubs (e.g. *Cornus sanguinea* and *Frangula alnus*) are cut back if they show considerable encroachment towards the meadows. As a result, forest edges are stable in position and have not changed notably since the first military mapping of the region in 1783 (Molnár et al., 1997).

### 2.2. Data collection

In July 2014, we prepared a special mowing plan for the site managers. In four locations, mowing was performed as usual, without leaving any unmown fringe along the forest edges. In other locations, five or ten meters of unmown buffers were left, with four spatial replicates for each width. In four further locations, no mowing was allowed (control). Each of these locations corresponded to an 80–100 m long straight forest edge (Fig. 1A). In each location, we installed eight pitfall traps (128 in total), of which four traps were placed in a line in the peripheral zone of the meadows, approx. 2.5 m from the forest edges, and four traps parallel to them in the peripheral zone of the forests, approx. 2.5 m from the forest edge (Fig. 1B). We used 0.5 l plastic glasses as traps with an upper diameter of 8 cm. The trapping fluid was ethylene glycol diluted with water (1/2, v/v) and a few drops of detergent were also added. The traps were installed immediately after mowing and were open for seven days. We chose this short period because grassland vegetation starts to regenerate from approximately one or two weeks onward, which would have obscured the short-term perturbations of the arthropod assemblages the research was designed to measure.

### 2.3. Data processing

We applied permutational multivariate analysis of variance (perMANOVA) with 1000 permutations to identify the effects of mowing and buffer width on the composition of spider and carabid assemblages. Calculations were performed separately on the untransformed species-abundance matrices of the meadows and forests. Traps were handled separately but their nested arrangement was accounted for in the analysis. If a result was significant, we carried out pair-wise perMANOVAs and adjusted the resulting *P*-values with the FDR method. For a visual representation of compositional patterns, we



**Fig. 1.** Mowing (A) and sampling (B) design of the study. No mowing was performed in the control sites, while 0, 5 or 10 m wide unmown buffer zones were left along the forest edges of the other sampling sites. The dimensions of the white rectangles indicating different treatment groups are not to scale. Eight traps were placed in each sampling site (black dots), four in the peripheral zone of the forests and four in that of the meadows.

performed principal component analysis (PCA). Prior to this analysis, the four traps of each location were pooled and the resulting data were square-root transformed.

We prepared generalised linear mixed-effects models (GLMMs) on the untransformed total abundance values of the arthropods using ‘treatment type’ as the fixed factor (four levels: control and 0 m, 5 m and 10 m buffer widths) and ‘location’ as the random factor to account for the spatial non-independence of traps within each location. We used negative binomial error term because the data were overdispersed. Finally, we sorted the species according to their habitat preference into forest specialist, open habitat specialist and generalist species groups following Buchar et al. (2002) for spiders and Freude et al. (2004) and Hurka (1996) for carabids, and prepared similar GLMMs on their abundance. Separate models were made for the meadow and forest data. The significance of the models was assessed by comparing the full model to a null model containing only the random factor. If a model explained a significant proportion of the variability of the data, pairwise comparisons of the factor levels were also considered and the resulting *P*-values were adjusted with the FDR method.

We made all calculations in R environment (R Core Team, 2014). The perMANOVAs and the PCAs were performed with the ‘adonis’ and ‘rda’ functions of the *vegan* package (Oksanen et al., 2016), respectively. We used the ‘glmer.nb’ function of the *lme4* package (Bates et al., 2015) for the GLMMs. Comparisons of full and respective null models were made with the ‘anova’ function of the built-in *stats* package. Pairwise comparisons were assisted by the ‘relevel’ function and the adjusted *P*-values were calculated with the ‘p.adjust’ function.

### 3. Results

#### 3.1. Spiders

We collected a total of 3035 spiders of 97 species, which included 25 forest specialist, 41 open habitat specialist and 31 generalist species. We caught 1332 individuals in the forests (63 species) and 1703 (79 species) in the meadows. According to the perMANOVAs, the composition of spider assemblages was affected by treatment type, and the effect was significant for both habitats (Table 1). Pairwise comparisons of the forest data revealed that the spider assemblage of the ‘0 m’ group differed from all other groups. Significant compositional difference was also detected between the ‘5 m’ and the ‘10 m’ buffer types. The control did not differ from the ‘5 m’ and ‘10 m’ groups. Similarly, the separation of the ‘0 m’ group was detectable on the PCA scatter plot as well, while the other groups showed a considerable overlap. The separation of the ‘5 m’ and ‘10 m’ groups was not clear using the two most important components (Fig. 2A). In the meadows, we found significant differences between the control and all the three treatment groups. Furthermore, the ‘0 m’ group differed from the ‘5 m’ and the ‘10 m’ groups; that is, difference was detected for all group pairs but the ‘5 m’ and ‘10 m’ groups. These findings also appear on the PCA scatter plot, with a considerable overlap between groups ‘5 m’ and ‘10 m’ and a clearer separation between all other group pairs (Fig. 2B).

The GLMMs of the forest data revealed that treatment significantly affected the total abundance as well as the abundance of forest and open habitat specialist spiders but not that of the generalist ones



**Table 1**

perMANOVA results of the spider and carabid assemblages of the peripheral zones of the forests and meadows. 'Total' refers to the full perMANOVAs and the other results are pairwise comparisons. C: control; \* $P < .05$ ; \*\* $.01 < P < .05$ ; \*\*\* $P < .001$ ;  $P$ -values were corrected for multiple comparisons with the FDR method.

Forest	<i>F</i>	<i>R</i> <sup>2</sup>	<i>P</i>	Meadow	<i>F</i>	<i>R</i> <sup>2</sup>	<i>P</i>
<b>SPIDERS</b>							
Total	2.54	0.094	< .001***	Total	3.34	0.111	< .001***
C – 0 m	2.32	0.071	.012*	C – 0 m	3.24	0.093	< .001***
C – 5 m	1.84	0.056	.062	C – 5 m	2.36	0.068	.014*
C – 10 m	1.21	0.039	.279	C – 10 m	2.56	0.079	.005**
0 m – 5 m	2.30	0.070	.012*	0 m – 5 m	2.13	0.059	.042*
0 m – 10 m	2.07	0.064	.018*	0 m – 10 m	3.98	0.120	.004**
5 m – 10 m	2.89	0.085	.007**	5 m – 10 m	1.25	0.039	.246
<b>CARABIDS</b>							
Total	2.01	0.074	.014*	Total	2.18	0.069	.014*
C – 0 m	2.48	0.077	.014*	C – 0 m	1.11	0.035	.418
C – 5 m	1.87	0.059	.110	C – 5 m	2.55	0.072	.021*
C – 10 m	1.45	0.047	.238	C – 10 m	1.32	0.045	.364
0 m – 5 m	1.30	0.044	.238	0 m – 5 m	1.78	0.055	.107
0 m – 10 m	1.23	0.041	.273	0 m – 10 m	1.08	0.038	.418
5 m – 10 m	1.19	0.038	.273	5 m – 10 m	1.03	0.034	.438

(Fig. 3A–D, Table 2). Regarding total abundance, only the '0 m' group had lower values than the control, and the '5 m' group had higher values than the '0 m' and the '10 m' groups. Forest specialist spiders were fewer in the '0 m' group than in the other groups, which, in turn, did not differ from each other. We detected no significant difference between the control and the other groups regarding open habitat species but we caught significantly more individuals in the '5 m' group than in the '0 m' and '10 m' groups.

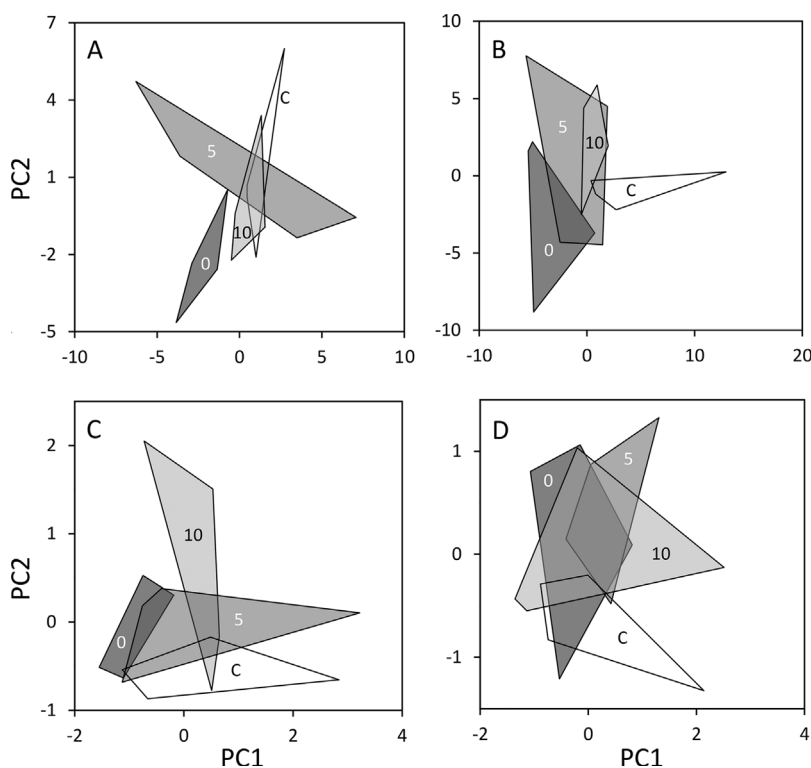
In the meadows, treatment affected only the abundance of forest specialist species, whereas the total abundance and the abundance of open habitat specialist and generalist species showed no significant treatment effect (Fig. 3E–H, Table 2). Both the '0 m' and '5 m' groups had fewer forest specialists than the control and the difference was also significant between the '0 m' and the '10 m' groups, with the latter being more abundant in forest specialist spiders.

### 3.2. Carabid beetles

We caught a total of 1148 carabid individuals of 57 species, which included 12 forest specialist, 23 open habitat specialist and 22 generalist species. There were 542 individuals in the forests (37 species) and 606 (50 species) in the meadows. Treatment had significant effect on the composition of carabid assemblages in both habitats (Table 1). The pairwise comparisons of the forest data indicated that the control differed from the '0 m' group, while the other comparisons yielded nonsignificant results. The PCA scatterplot of forest carabids (Fig. 2C) also indicated a separation between the control and the '0 m' group. There was no overlap between the '0 m' and the '10 m' groups either but the compositional difference was not confirmed statistically with the corresponding pairwise perMANOVA. In the meadows, we detected significant difference only between the control and the '5 m' group, which was in line with the patterns of the PCA scatter plot (Fig. 2D), showing a separation between these groups and a considerable overlap between other group pairs.

Regarding carabids in the forests, all GLMMs were significant (Fig. 3I–L, Table 2). Total abundance was significantly lower in the '0 m' group than in all other groups, and the '5 m' group tended to have more individuals than the control but we detected only marginal significance. The other pairwise comparisons yielded nonsignificant results. Forest specialist carabids were less abundant in the '0 m' group than in the other groups, whereas these latter groups did not significantly differ from each other. Open habitat specialists were more abundant in the control and the '5 m' group than in the '0 m' and '10 m' groups. We detected no difference between the control and the '5 m' group and between the '0 m' and the '10 m' groups. We caught more generalist species in every treated group than in the control and there were more carabids in the '5 m' group than in the '0 m' group.

In the meadows, treatment had no detectable effect on either the total abundance of carabids or the abundance of forest and open habitat specialist species. However, we detected a marginal significance for generalist species; thus, we did the pairwise comparisons. These indicated significantly more carabids in the '5 m' group than in the control and the '10 m' groups (Fig. 3M–P, Table 2).



**Fig. 2.** Scatterplots of principal component analysis of spiders in the forests (A) and meadows (B) and carabids in the forests (C) and meadows (D). Vertices of the polygons represent forest edge sites (pooled data of four traps). C: unmowed control; 0: zero buffer; 5: narrow (5 m) buffer; 10: wide (10 m) buffer. Variances covered by the first two axes: 2A: 29.3 and 24.4%, 2B: 35.3 and 32.1%, 2C: 37.9 and 14.1%, 2D: 24.5 and 12.9%.

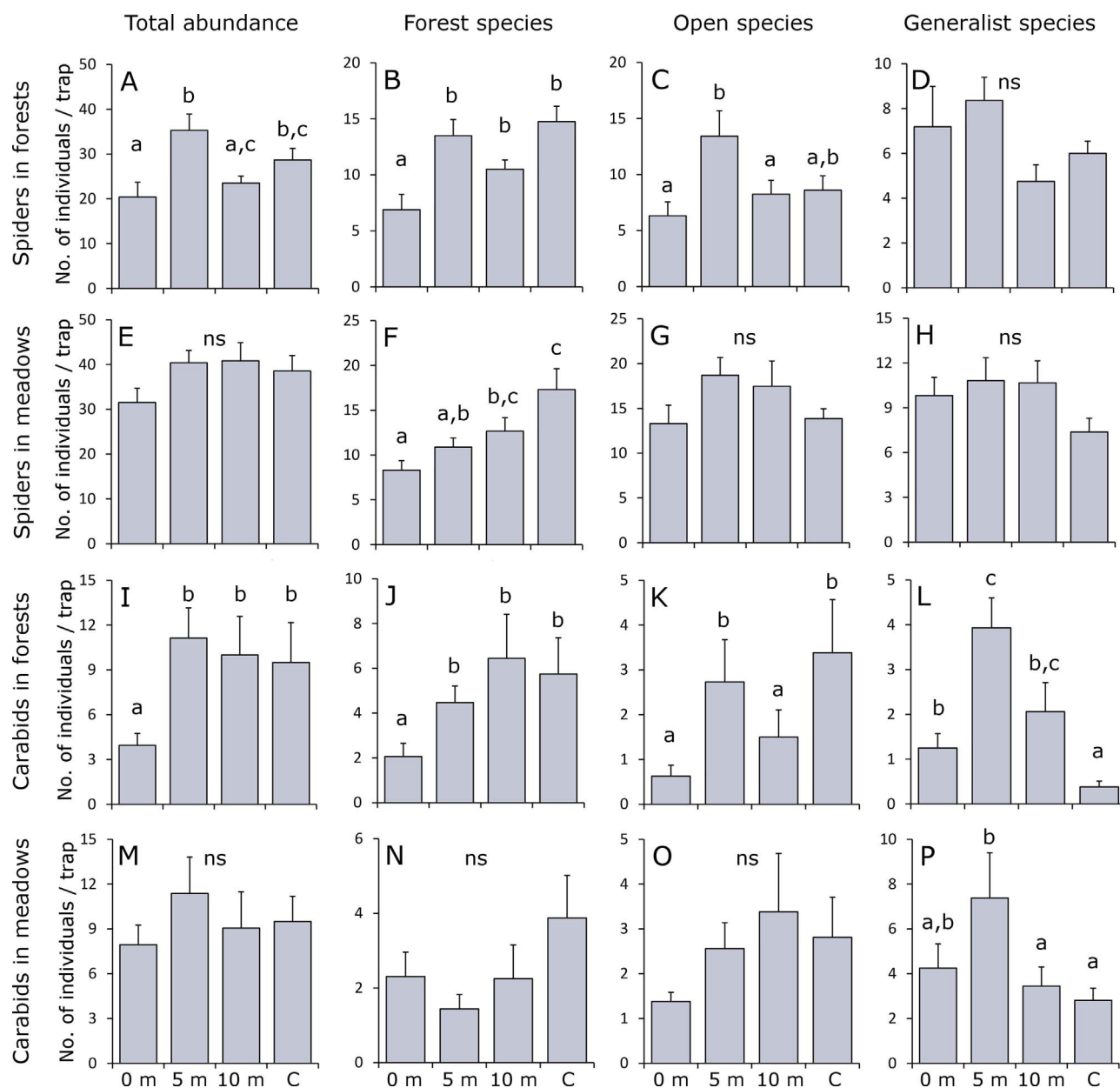


Fig. 3. The total abundance of spiders and the abundance of forest specialist, open habitat specialist and generalist spiders in the forests (A–D, respectively) and in the meadows (E–H, respectively), and the total abundance of carabids and the abundance of forest specialist, open habitat specialist and generalist carabids in the forests (I–L, respectively) and in the meadows (M–P, respectively). Whiskers are standard errors, lowercase letters identify significantly different groups, ‘ns’ indicates nonsignificant treatment effects and ‘C’ stands for control.

## 4. Discussion

### 4.1. Compositional patterns

We found that mowing resulted in significant changes in the species composition of both habitats, indicating a strong cross-edge effect of mowing. It appears that leaving a narrow (5 m) buffer may offset the compositional perturbations in the forests, although a marginal compositional difference was still detectable for spiders. In the meadows, even the wide buffer was not enough to avoid all compositional changes for spiders.

Interestingly, some of our results suggest that the effect of the narrow buffer zone is not intermediate between the zero buffer (0 m) and the wide buffer, as (i) the composition of the narrow buffer group and the control differed in a case, where the zero buffer did not differ from the control (see carabids in the meadows) and (ii) in another case

the control did not differ significantly from either the narrow or the wide buffers but the narrow and the wide buffers did differ from each other (see spiders in the forest). The effects of buffers are therefore likely to be more complex than anticipated. The models prepared for the total and trait-specific abundances did, however, shed light on the main trends.

### 4.2. Abundance patterns in the meadows

We found that treatment did not affect the total abundance of either arthropod taxon in the meadows but the abundance of forest specialist spiders decreased due to mowing. This decrease in abundance was less severe when a narrow buffer was applied and was no more detectable with the wide one as compared to the unknown control. This reaction of the forest specific species can be explained by their sensitivity to dry and warm microclimate (Magura et al., 2001; Noreika and Kotze,

**Table 2**  
 Test results of the GLMMs prepared for the spider and carabid assemblages of the forest and meadow sites. Pairwise results are provided if treatment had significant or marginally significant effect. \* $P < .05$ ; \*\* $.01 < P < .05$ ; \*\*\* $P < .001$ ;  $P$ -values were corrected for multiple comparisons with the FDR method.

	FOREST						MEADOW					
	Total abundance	Forest spp.	Open spp.	Generalist spp.	Total abundance	Forest spp.	Open spp.	Generalist spp.	Total abundance	Forest spp.	Open spp.	Generalist spp.
SPIDERS	$Chi^2$	$Chi^2$	$Chi^2$	$Chi^2$	$Chi^2$	$Chi^2$	$Chi^2$	$Chi^2$	$Chi^2$	$Chi^2$	$Chi^2$	$Chi^2$
Treatment	10.85	15.37	7.92	4.01	2.85	.416	11.35	.010	.686	5.12	.163	
	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$
C-0 m	.026*	-4.31	-1.31	.189	. . .	. . .	-4.02	<.001***	. . .	. . .	. . .	. . .
C-5 m	.177	-0.26	1.86	.063	. . .	. . .	-2.59	.010*	. . .	. . .	. . .	. . .
C-10 m	.190	-1.87	-0.20	.843	. . .	. . .	-1.77	.077	. . .	. . .	. . .	. . .
0 m-5 m	<.001***	3.98	3.17	.002**	. . .	. . .	1.45	.147	. . .	. . .	. . .	. . .
0 m-10 m	.349	2.51	1.13	.257	. . .	. . .	2.22	.027*	. . .	. . .	. . .	. . .
5 m-10 m	.008**	1.57	-2.09	.037*	. . .	. . .	0.79	.431	. . .	. . .	. . .	. . .
CARABIDS	$Chi^2$	$Chi^2$	$Chi^2$	$Chi^2$	$Chi^2$	$Chi^2$	$Chi^2$	$Chi^2$	$Chi^2$	$Chi^2$	$Chi^2$	$Chi^2$
Treatment	19.2	11.97	11.59	21.26	3.66	.301	5.19	.158	.273	7.12	.068	
	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	
C-0 m	-2.58	-2.98	-2.93	.003**	. . .	. . .	. . .	. . .	. . .	. . .	. . .	
C-5 m	1.92	0.29	-0.04	.970	. . .	. . .	. . .	. . .	. . .	. . .	. . .	
C-10 m	0.18	-0.02	-2.07	.037**	. . .	. . .	. . .	. . .	. . .	. . .	. . .	
0 m-5 m	4.65	2.98	2.84	.004**	. . .	. . .	. . .	. . .	. . .	. . .	. . .	
0 m-10 m	2.77	3.13	1.10	.270	. . .	. . .	. . .	. . .	. . .	. . .	. . .	
5 m-10 m	-1.76	-0.31	-1.99	.046*	. . .	. . .	. . .	. . .	. . .	. . .	. . .	
	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$
C-0 m				2.29	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-5 m				4.42	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-10 m				3.20	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-5 m				2.65	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-10 m				1.08	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
5 m-10 m				-1.65	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$
C-0 m	.010*	-2.98	-2.93	.003**	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-5 m	.055	0.29	-0.04	.970	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-10 m	.855	-0.02	-2.07	.037**	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-5 m	<.001***	2.98	2.84	.004**	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-10 m	.006**	3.13	1.10	.270	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
5 m-10 m	.080	-0.31	-1.99	.046*	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$
C-0 m				2.29	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-5 m				4.42	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-10 m				3.20	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-5 m				2.65	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-10 m				1.08	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
5 m-10 m				-1.65	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$
C-0 m	.010*	-2.98	-2.93	.003**	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-5 m	.055	0.29	-0.04	.970	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-10 m	.855	-0.02	-2.07	.037**	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-5 m	<.001***	2.98	2.84	.004**	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-10 m	.006**	3.13	1.10	.270	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
5 m-10 m	.080	-0.31	-1.99	.046*	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$
C-0 m				2.29	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-5 m				4.42	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-10 m				3.20	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-5 m				2.65	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-10 m				1.08	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
5 m-10 m				-1.65	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$
C-0 m	.010*	-2.98	-2.93	.003**	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-5 m	.055	0.29	-0.04	.970	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-10 m	.855	-0.02	-2.07	.037**	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-5 m	<.001***	2.98	2.84	.004**	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-10 m	.006**	3.13	1.10	.270	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
5 m-10 m	.080	-0.31	-1.99	.046*	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$
C-0 m				2.29	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-5 m				4.42	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-10 m				3.20	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-5 m				2.65	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-10 m				1.08	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
5 m-10 m				-1.65	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$
C-0 m	.010*	-2.98	-2.93	.003**	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-5 m	.055	0.29	-0.04	.970	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-10 m	.855	-0.02	-2.07	.037**	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-5 m	<.001***	2.98	2.84	.004**	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-10 m	.006**	3.13	1.10	.270	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
5 m-10 m	.080	-0.31	-1.99	.046*	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$
C-0 m				2.29	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-5 m				4.42	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-10 m				3.20	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-5 m				2.65	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-10 m				1.08	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
5 m-10 m				-1.65	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$
C-0 m	.010*	-2.98	-2.93	.003**	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-5 m	.055	0.29	-0.04	.970	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-10 m	.855	-0.02	-2.07	.037**	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-5 m	<.001***	2.98	2.84	.004**	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-10 m	.006**	3.13	1.10	.270	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
5 m-10 m	.080	-0.31	-1.99	.046*	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$	$z$
C-0 m				2.29	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-5 m				4.42	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-10 m				3.20	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-5 m				2.65	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-10 m				1.08	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
5 m-10 m				-1.65	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$	$P$
C-0 m	.010*	-2.98	-2.93	.003**	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-5 m	.055	0.29	-0.04	.970	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
C-10 m	.855	-0.02	-2.07	.037**	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-5 m	<.001***	2.98	2.84	.004**	. . .	. . .	. . .	. . .	. . .	. . .	. . .	. . .
0 m-10 m	.006**	3.13	1.10	.270	. . .	. . .	. . .	. . .				

because forests are not as far from their optimum than for the open habitat specialists. The narrow buffer further increased the influx of generalist species and even the wide buffer had a positive effect on their spillover compared with the control. These findings are in line with other studies claiming that generalists are the main contributors of spillover across habitat edges (e.g. Schneider et al., 2013; Torma and Gallé, 2011).

The facilitative effect of narrow buffers on spillover into forests can be either beneficial or undesirable. It is beneficial when the species that temporarily find shelter in the peripheral zones of forests are of conservation importance and their long-term persistence is supported by this mechanism, as a novel, management-assisted landscape complementation. Leaving unmown strips or patches away from forest fragments may potentially have similar beneficial effects (Humbert et al., 2012b; Lebeau et al., 2015). Conversely, the increased spillover of spiders and carabids can interfere with trophic interactions in the forests, creating higher predatory pressure on forest-specific prey assemblages and increasing competition with forest-specific predators (Boetzel et al., 2016; Lacasella et al., 2015; Rand et al., 2006). As a result, leaving either zero buffer or a wide buffer may be a better option than leaving a narrow buffer if the net effect of the spillover from grasslands is considered negative for landscape level biodiversity and ecosystem functions.

## 5. Conclusions

Although the detected trends of carabids and spiders were not completely symmetrical regarding either composition or abundance, none of the statistically confirmed differences were contrasting. In fact, the results of spiders and carabids supplemented each other to receive a coherent view on the effects of the treatments.

We conclude that unmown buffers along forest edges can, depending on their width, reduce or completely offset the effects of mowing on the composition of arthropod assemblages of forests. However, even wide buffers cannot fully preserve species composition in the peripheral zones of meadows. Regarding abundance change, spillover from forests is reduced by mowing but this can fully be reversed by applying wide buffers. Changes inside forests are more complex. Forest specialists withdraw from the peripheral zones but open habitat specialists do not necessarily enter the forests in higher numbers. The application of narrow buffers can facilitate the influx of open habitat specialists and generalists but this effect disappears when using wide buffers. On the basis of these findings, spillover should be considered when planning mowing in grassland-forest mosaics. We recommend using wide buffers but since this can also lead to shrub or tree encroachment, the application of wide buffers should be alternated with zero buffers both in space and time. Narrow unmown buffers along the forest edges should mostly be avoided as they can facilitate unwanted spillover. The actual width of effective buffers may vary according to different climatic regimes, different forest and grassland types, and management intensity; thus, choosing the right width may require local optimization. If, for instance, the grassland is more intensively used than in our case, generalist species may have a higher proportion and could pose a higher threat to the forest assemblages after mowing due to their higher mobility across edges. In such cases, buffers wider than 10 m may be necessary.

Future research should also test the application of buffers in preventing undesirable changes of spillover in forest-cropland mosaics. Although unharvested crop strips do not seem a feasible option, extensively used, permanent grassland buffers, as also recommended by Madeira et al., 2016, may be more appropriate for the purpose. Our results, however, indicate that simply deciding for the use of buffers is not enough. It may be necessary to test multiple buffer widths, because a too narrow buffer may not only have lower efficiency but may have an opposite effect compared to the intended one.

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