RESEARCH ARTICLE

Seasonal shifts and complementary use of pollen sources by two bees, a lacewing and a ladybeetle species in European agricultural landscapes

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Funding information
Schweizerischer Nationalfonds zur Förderung der Wissenschaftlichen Forschung, Grant/Award Number: 6571210; Deutsche Forschungsgemeinschaft, Grant/Award Number: EN 979/3-1

Handling Editor: Lucas Garibaldi

Abstract

1. Continuous availability of food resources, such as pollen, is vital for many insects that provide pollination and pest control services to agriculture. However, there is a lack of knowledge about the shared or complementary use of floral resources by such species, which hampers more effective landscape management to simultaneously promote them in agroecosystems.

2. Here, we simultaneously quantified pollen use by a bumblebee (Bombus terrestris) and a mason bee (Osmia bicornis), two bee species recognized as important crop pollinators, as well as a lacewing (Chrysoperla carnea) and a ladybeetle species (Harmonia axyridis), both common predators of crop aphids, throughout the season in 23 agricultural landscapes in Germany and Switzerland.

3. Pollen diets were more diverse and similar among C. carnea and H. axyridis compared to the two bee species, but all four species shared key pollen types early in the season such as Acer, Quercus, Salix and Prunus. All species exhibited a pronounced shift in pollen sources from primarily woody plants (mainly trees) in spring to primarily herbaceous plants in summer. The majority of pollen (overall ≥64%) came from non-agricultural plants even in crop-dominated landscapes.

4. Synthesis and applications. Our results highlight the importance of trees as pollen sources for many insect species, particularly early in the season. Our findings support incentives that promote heterogeneous agricultural landscapes including both woody and herbaceous semi-natural habitats, ensuring phenological complementarity of floral resources for insect species that can provide pollination and pest control services to agriculture. The identified key plant species can help to design and optimize agri-environment schemes to promote these functionally important insects.

KEYWORDS
bumblebee, floral resources, foraging habitat, landscape resources, mason bee, pollen diet, resource specialization

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1 | INTRODUCTION

Insects critically contribute to biodiversity in agroecosystems and provide ecosystem services sustaining crop production, such as crop pollination and pest control. Those services are of paramount economic and non-monetary value for human well-being (IPBES, 2016; Losey & Vaughan, 2006). Bees and other crop pollinators, as well as many natural enemies of crop pests such as syrphids, lacewings and ladybeetles, feed on nectar or pollen provided by flowering plants. Floral resources dominate the diet of bees, regardless of their development stage. The larvae of natural enemies, on the other hand, are predators primarily feeding on animal prey including major agricultural pests such as aphids, whilst adults regularly consume nectar and pollen as a sole food source (e.g. most lacewings) or to supplement their diet with key nutrients lacking in insect-only diets, in particular during periods of prey scarcity (e.g. ladybeetles; Lundgren, 2009). Hence, adequate floral resources may enhance these functionally important insects in agricultural landscapes (e.g. Carvell et al., 2017; Isaacs, Tuell, Fiedler, Gardiner, & Landis, 2009; Wäckers & Van Rijn, 2012; Williams, Regetz, & Kremen, 2012). The loss and degradation of semi-natural vegetation – and the concomitant loss of floral resources – is considered a principal cause of the decline of pollinators and pest enemies and the services they provide (Benton, Vickery, & Wilson, 2003; IPBES, 2016; Scheper et al., 2014). Therefore, promoting adequate floral resources at the right place and time is critical for successful habitat management (Isaacs et al., 2009; M’Gonigle, Ponisio, Cutler, & Kremen, 2015; Sutter, Jeanneret, Burtalou, Bocci, & Albrecht, 2017) and can be highly effective to promote pest control (Tsuchi, Albrecht, Entling, & Jacot, 2015; Wäckers & Van Rijn, 2012) and pollination services (Blaauw & Isaacs, 2014) provided by insects.

Most common crop pollinators and pests’ natural enemies consuming floral resources are dietary generalists; that is, they collect floral resources from multiple plant taxa from both crop and non-crop habitats (e.g. Villenave, Thierry, Al Mamun, Lodé, & Rat-Morris, 2005; Walther-Hellwig & Frankl, 2000). However, even the diets of generalist flower visitors are usually dominated by certain floral resource types (e.g. Sutter et al., 2017; Wäckers & Van Rijn, 2012). Only a subset of resources offered by the flowering plant community in agroecosystems is accessible, available at the right time, of adequate chemical composition and attractive to different insect species. For instance, spatial resource use and accessibility depend on species’ mobility and foraging ranges (e.g. Walther-Hellwig & Frankl, 2000). Furthermore, mass-flowering crops may offer abundant floral resources, but are ephemeral and only available during short periods. Spatio-temporal complementarity and thus continuous provisioning of floral resources by non-crop vegetation is, therefore, vital to ensure population persistence of pollinators and many important pest enemies, and the ecosystem services they provide (Schellhorn, Gagic, & Bommarco, 2015).

Yet, there is a lack of knowledge about the spatio-temporal dynamics of floral resource use (but see e.g. Grab, Blitzer, Danforth, Loeb, & Poveda, 2017) by co-occurring pollinators and pest enemies, which hampers more effective landscape management to simultaneously promote them in agroecosystems. Regarding pollen – a main source of protein for many functionally important insects – it remains unclear what proportions of pollen consumed by different species come from crop and non-crop plants or from different vegetation types such as woody (e.g. woodlots or hedgerows) or herbaceous (e.g. grasslands, herbaceous field margins) habitats. Information regarding the extent of overlap or divergence in pollen use by multiple species over time is also missing. Thus, to increase the effectiveness of measures to concomitantly promote service providing insects, we need to simultaneously examine how different plant resources contribute to their requirements, and how similar or complementary their use of floral resources is (Rollin et al., 2013; Shackelford et al., 2013). Such knowledge is a prerequisite to better assess and predict the distribution of these functionally important insect taxa in agricultural landscapes, and to guide scientists and land managers in identifying and promoting habitats and specific floral resources that are vital to sustain them.

The main objective of this study was to compare the spatio-temporal use of pollen resources between two pollinator and two aphid enemy insect species commonly found in Central European agricultural landscapes and that rely on pollen for at least part of their life cycle. The bumblebee Bombus terrestris and the mason bee Osmia bicornis were used as pollinator model taxa. These two species rely on pollen for development and adult survival, and are among the most abundant wild bees in the studied agroecosystems (e.g. Kleijn et al., 2015; Westphal et al., 2008) that provide pollination services in a wide range of crops, such as pumpkin (Pfister, Eckter, Schirmel, Cresswell, & Entling, 2017), field beans and oilseed rape (Garratt et al., 2014), or fruit trees and strawberries (Gruber, Eckel, Everaars, & Dormann, 2011; Klatt et al., 2014). Aphid enemies that rely on pollen include hoverflies, ladybeetles and lacewings. We selected the lacewing Chrysoperla carnea s.l. due to its high abundance in annual cropping systems (McEwen, New, & Whittington, 2007; Pfister, Schirmel, & Entling, 2017). The ladybeetle Harmonia axyridis, despite being invasive in Europe (Roy et al., 2012), was also included as a model taxon since it became one of the most dominant aphid predators in European agroecosystems in the past years (e.g. Pfister, Schirmel, et al., 2017; Stutz & Entling, 2011). For C. carnea and H. axyridis adults, pollen is either an obligatory (Chrysoperla) or complementary (Harmonia) part of their diet, which can be vital in particular during periods of prey scarcity (Berkevens et al., 2010; Lundgren, 2009).

We addressed the following questions:

1. What are the main pollen types used by the studied bee, lacewing and ladybeetle species?
2. To what extent does pollen use overlap among the four species? Do they share important pollen taxa that could be targeted by habitat management schemes?
3. What is the relative importance of different pollen sources (i.e. woody/herbaceous plants, crop/non-agricultural plants)? How does the use of pollen sources change over the season and does...
the pollen use of the four insect species show similar temporal dynamics?
4. How does the landscape context influence the use of different pollen sources?

2 | MATERIALS AND METHODS

2.1 | Study regions and pollen sampling design

The study was conducted in 2016 in southwestern Germany and northeastern Switzerland (see Appendix S1.A). In each region, 11 (Germany) to 12 (Switzerland) landscape sectors of 500 m radius were selected. The selected landscapes represent the typical range in the proportion of the two major land-use types characteristic for the study regions: agricultural land (e.g. arable crops, managed grassland, some horticulture; 38%–90%, mean = 68%, SD = 16%) and woody habitat (woodlots and hedgerows; 0%–51%, mean = 11%, SD = 12%; see Appendix S1.A; Figure S1). In each landscape sector three (Germany) to five (Switzerland) sampling points were selected: one central sampling point, as well as 2 to 4 further ones randomly spread across the landscape (see Appendix S1.B).

Samples of insects and pollen were collected approximately every 2 weeks from beginning of April–mid-June (see Table S2). At each sampling point, adults of C. carnea and H. axyridis were sampled using sticky traps. For each species, up to five individuals per sampling round and landscape sector were randomly selected for pollen analysis. Pollen collected by B. terrestris was obtained from the pollen sacs of up to 10 worker bees per sampling round and landscape sector upon return to colonies experimentally established at each landscapes’ central sampling point (“Mini hive”; purchased from Biobest and containing 30–40 workers). Pollen collected by O. bicornis was obtained from up to five brood cell provisions per sampling round of experimentally established trap nests at each sampling point (Switzerland) or central sampling point (Germany; see Appendix S1.B; Figure S2).

2.2 | Pollen samples processing and analysis

Elytra and/or wings of sampled C. carnea and H. axyridis individuals were removed, and insects were thoroughly rinsed with ETOH to remove pollen from the exoskeleton to minimize the potential of including in the analysis pollen grains that were not consumed by the insects. Subsequently, insects were crushed and acetolysis was performed following Jones (2012).

All pollen samples belonging to the four insect species were treated chemically with acetolysis and KOH, and mounted in glycerine following standard palynological methods (Moore, Webb, & Collinson, 1991). Pollen grains were identified under a light microscope (400× magnification) based on palynological keys (Beug, 2004; Moore et al., 1991) and a photo atlas (Reille, 1992), as well as using the reference collection of the Institute of Plant Sciences of the University of Bern. Pollen grains were identified at species whenever possible, or at subgenus, genus, or family level (hereafter pollen types (=t); see Table S1). About 1,070 samples were available for analysis (see Table S2). For each sample, we identified and counted up to 100 pollen grains whenever possible (i.e. between 30 and 100 grains).

2.3 | Data analysis

All statistical analyses were conducted using R 3.4.1 (R Core Team, 2017). To account for unequal numbers of pollen grains between samples or insect species, and as we were interested in pollen composition comparisons, data were always standardized to proportions (i.e. relative contributions with total 100), and analyses were performed using pollen types percentages. For analyses of temporal dynamics and to facilitate comparisons across the two study regions, four sampling periods were defined based on accumulated Growing Degree Days (GDD) (see Table S3).

To describe samples’ pollen types diversity, we used pollen type richness (number of pollen types), as well as the Simpson diversity index, which represents the probability that two grains randomly selected from a sample will belong to different types \((1 – D, \text{ with } D = \sum p^2, p \text{ being the proportion of pollen grains belonging to one pollen type}). Mean pollen type richness and mean Simpson diversity were compared for the four insect species with Kruskal-Wallis and Dunn tests.

To assess the degree of exclusiveness or overlap in pollen use among the four studied insect species, complementary specialization \(d^\prime\) (Blüthgen, Menzel, & Blüthgen, 2006) was calculated for each species for each sampling period (R package "bipartite 2.08"; Dormann, Fründ, Blüthgen, & Gruber, 2009). The index \(d^\prime\) measures how strongly the pollen types collected by a species deviate from that of other species (Kämper et al., 2016). The measure ranges from 0 (complete overlap in pollen types use, i.e. “opportunistic” species sharing all their pollen types with other species) to 1 (exclusive pollen types use, i.e. “specialized” species; Blüthgen, Fründ, Vázquez, & Menzel, 2008; Junker et al., 2013). Mean \(d^\prime\) along the season was compared for the four species with Student t test. We also calculated \(H_{2}^\prime\), which describes the average degree of complementary specialization for the four insect species (i.e. network specialization; Blüthgen et al., 2006). Network specialization equals the weighted sum of the specialization of its nodes (i.e. weighted sum of \(d^\prime\) of all species). It also ranges from 0 (pollen types used by the four species completely overlap; “maximum niche overlap”; Schleuning et al., 2012) to 1 (each species uses a unique set of pollen types; “maximum exclusiveness” or “maximum niche divergence”; Blüthgen et al., 2008; Schleuning et al., 2012). The species-level index \(d^\prime\) was used to compare the specialization levels of the four studied species within networks, while \(H_{2}^\prime\) index was used for comparing the different networks across the season. The two study regions were analysed together, and for each sampling period all samples belonging to one insect species were pooled. Only pollen types that accounted for more than 1% of the total number of pollen grains were considered for the analyses.
To examine the importance of pollen from woody plants, pollen types were classified as "woody" or "herbaceous" (see Table S1). Pollen types that could not be identified at the species level potentially including both woody and herbaceous plants (5% ± 10% of the data) were excluded from these analyses. For each insect species and sampling period, samples within a landscape were pooled. We used generalized linear mixed models (GLMMs) with a binomial error distribution to test the impacts of the fixed factors sampling period, species (B. terrestris, O. bicornis, C. carnea, H. axyridis), study region (Germany/Switzerland) and their interactions and the random factor landscape sector, on the proportion of pollen from woody plants used by the insects. An observation level term was added as a second nested random effect to account for overdispersion (Lee & Nelder, 2000). Models were fitted with the R package “lme4 1.1-13” (Bates, Maechler, Bolker, & Walker, 2015). As there was no significant three-way interaction in the full model (p = .64), indicating consistent temporal patterns of pollen use across species in both regions, data of both regions were finally analysed together. Predictors’ significance was tested with Wald chi-square tests. We also examined the importance of non-agricultural plants by classifying pollen types as "non-agricultural" (i.e. associated plant taxa can be unambiguously classified as non-agricultural plants, which includes plants from semi-natural habitats and crop weeds) and “potential crop” (i.e. associated plant taxa could potentially include crop or sown grassland plant species; see Table S1). We used similar GLMMs to those previously described, with the proportion of pollen from non-agricultural plants as the response variable, and sampling period, insect species and their interactions as fixed factors. In this case, as the significant three-way interaction of the full model indicated distinct patterns among the two study regions, they were analysed separately.

To examine the influence of the landscape context on pollen use, we calculated landscape metrics using ArcGIS 10.5 (ESRI) based on 2016 land-cover maps of the study regions. Maps were digitized based on photo interpretation and were completed and validated with ground surveys. Land use was classified into five categories: woody semi-natural habitats (e.g. woodlands, hedgerows), woody crops (including vines and orchards), herbaceous crops (e.g. cereals), grasslands and “other land use” (including settlements). We calculated two metrics within each landscape sector of 500 m radius: the surface of woody semi-natural habitats, and the total surface of woody land-use types (including woody semi-natural habitats and woody crops). We used as a basis the GLMMs previously described, including pollen proportion from woody plants or from non-agricultural plants used by insects as the response variable, and sampling period, insect species and their interactions as fixed factors, and landscape sector and observation-level term as random effects. We complexified those models by adding a landscape metric and interactions with sampling period and insect species as additional fixed factors. The total surface of woody habitat was used as a landscape metric for the first model including the pollen proportion from woody plants as the response variable, whereas the surface of woody semi-natural habitat was used for the second model including the pollen proportion from non-agricultural plants as the response variable. We confirmed that we had no remaining spatial autocorrelation in the models by checking residuals against spatial coordinates with correlogram plots using the ncf package in R (Bjornstad, 2019).

3 RESULTS

3.1 Main types and diversity of pollen used

A total of 140 different pollen types were identified: 91 types were used by B. terrestris, 54 by O. bicornis, 99 by C. carnea and 82 by H. axyridis (see Table S1). Most individual samples contained at least two different pollen types (see Figure S3). Individual samples of pollen used by C. carnea and H. axyridis had a roughly three times higher pollen type richness than those of the two bee species (Dunn test, p < .001), and this pattern was consistent across the season (Figure 1). Simpson diversity of pollen types was also higher in samples from C. carnea and H. axyridis (0.54 ± 0.24 and 0.66 ± 0.18, respectively) than in those from O. bicornis (0.27 ± 0.22) and B. terrestris (0.17 ± 0.22; Dunn test: p < .001). Results were similar when pooling samples at the landscape level (see Figure S4).

Bombus terrestris collected mainly pollen from insect-pollinated plants (83% ± 25%) in contrast to H. axyridis which was mainly using pollen from wind-pollinated plants (67% ± 19%), whereas O. bicornis and C. carnea used pollen from both, insect- and wind-pollinated plants (see Table S4). More precisely, early in the season, B. terrestris collected Salix, Prunus t. (=type), Acer and Brassicaceae pollen (presumably oilseed rape; see Table S5), accounting for more than 80% of the pollen collected. Later in the season, mainly Rubus, Papaver rhoeas t., Trifolium (mainly Trifolium repens t.) and Tilia pollen were collected by this species (Figure 2). O. bicornis collected mainly Acer and Quercus pollen early in the season, accounting for more than 65% of the pollen collected, whereas Acer and Ranunculus acris t. (probably Ranunculus sp.) dominated in summer samples. Prunus t., Betula, Salix, Carpinus, Acer, Fagus, Quercus and Brassicaceae pollen covered more than 60% of the pollen diet of C. carnea early in the season, whereas Poaceae species dominated the pollen diet during the summer months. Finally, H. axyridis consumed mainly Betula, Fagus, Carpinus, Quercus, Acer and Pinus pollen early in the season, covering almost 60% of the pollen diet, whereas half of the pollen consumed in summer belonged to Urtica and Poaceae species (Figure 2).

3.2 Overlap in pollen use among insect species

Complementary specialization at the species level was on average twice as high in the two bee species compared to C. carnea and H. axyridis (Figure 3; mean d’ of 0.65 and 0.30 for the two bees and the two aphid enemy species, respectively; Student t test: p < .001). At the network level, the degree of complementary specialization (H2S; i.e. mean complementary specialization of all four insect species) was low to intermediate, ranging from 0.33 to 0.53 across the sampling season (mean = 0.46; Figure 3). Overlap in pollen use
between the four insect species was highest in May (GDD 100–200; \(H_2' = 0.33\), i.e. less pronounced niche complementarity), primarily due to a relatively high proportion of shared pollen types from woody plants such as Acer, Quercus, Fagus, Prunus t. and Salix, as well as a fairly general use of Brassicaceae pollen (Figure 3). Further key pollen types shared by at least two species included Betula early in the season (April, GDD 0–100), and Poaceae, Tilia, Papaver rhoeas t. and Ranunculus acris t. later in the season (June to mid-July, GDD 201–600).

### 3.3 | Relative importance of different pollen sources, temporal shifts and landscape effects

For all insect species, the proportion of pollen from woody plants (collected from trees and shrubs) was high early in the year (April and May, GDD 0–200), but decreased significantly later in the season (June, GDD 201–400), indicating a shift from woody to herbaceous pollen sources (Table 1; Figures 2–4). The proportion of pollen from woody plants remained low until mid-July (GDD 401–600) for most species, but tended to increase again for B. terrestris because of the importance of Tilia pollen for this species in the late season. There was no significant relationship between the proportion of woody habitats in the landscapes and the proportion of pollen from woody plants collected by the insects (\(p > .05\); see Table S6).

There was no clear temporal trend in the proportion of pollen from non-agricultural plants used by studied insect species. Throughout the season, O. bicoris used higher percentages (82 ± 17%) of pollen from non-agricultural plants than H. axyridis (63 ± 15%), B. terrestris (57 ± 14%) or C. carnea (56 ± 9%; Table 2). At any time of the season and across all insect species, however, a significant part of the pollen diet was from non-agricultural plants (min ≥40%, mean ≥64%; Table 2). There was no significant relationship between the proportion of woody semi-natural habitats in the landscapes and the proportions of pollen from non-agricultural plants collected by the insects (\(p > .05\); see Table S6).

### 4 | DISCUSSION

Our findings reveal: (a) a higher diversity and lower pollen diet specialization of the two potential aphid enemies Chrysoperla carnea and Harmonia axyridis compared to the two studied bee species; (b) some important pollen plant taxa (e.g. Acer) shared by all four species, in particular early in the season; (c) the importance of woody plants (primarily trees) as pollen sources early in the season and a pronounced shift from woody to herbaceous pollen sources during the season for all studied insect species; (d) a generally high proportion of pollen from weeds and non-agricultural sources used by all four species.

#### 4.1 Composition and diversity of pollen used

Our findings are in agreement with the general expectation that bees (O. bicoris and B. terrestris) – exclusively relying on pollen as protein source for offspring provisioning – more selectively use pollen taxa of high nutritional quality that can be collected at relative low energy costs (e.g. mass-flowering plants), while natural enemies (C. carnea and H. axyridis) are more opportunistic in their pollen use.

In spring as well as in summer, Osmia bicoris collected pollen mainly from a very limited number of plants: Acer and Quercus early in the year, and Ranunculus acris type (probably Ranunculus...
sp.), Acer, Juglans and Papaver in summer. The similar pollen use of O. bicornis in different years and regions (Coudrain, Rittiner, Herzog, Tinner, & Entling, 2016; Radmacher & Strohm, 2010) indicates clear preferences among plants. These plants include mass-flowering trees such as Quercus, which is wind-pollinated but provides abundant and high-quality pollen for bees (Roulston,
(a) GDD 0 to 100; $H' = 0.45$

(b) GDD 101 to 200; $H' = 0.33$

(c) GDD 201 to 400; $H' = 0.53$

(d) GDD 401 to 600; $H' = 0.52$
**FIGURE 3** Pollen use network of the four insect species at each sampling period. Growing Degree Days (GDD) 0–100 correspond approximately to the month of April, 101–200 to May, 201–400 to June and 401–600 to end of June to mid-July (see Table S3). H₁’ measures network specialization; it ranges from 0 for the most generalized (i.e. maximum niche overlap) to 1 for the most specialized network (i.e. high exclusiveness, or maximum niche divergence). Upper bars represent insect species and lower bars the average proportion of pollen types used across all insect species (see Table S1 for more information on pollen types). Brown colours represent pollen from woody plants, green colours those from herbaceous plants, and white those for which this information was not available or which comprise both woody and herbaceous plants. The width of the arrows between upper and lower bars represents the proportion of a pollen type used by an insect species. Number of samples is given in brackets next to species names, and values of species-level complementary specialization (d’) are shown below. A high d’ value indicates a high degree of specialization in pollen use of an insect species (high “exclusiveness”), whereas insect species sharing many pollen types with other taxa receive small d’ values (i.e. “opportunistically”).

**TABLE 1** Analysis of deviance table (Type II Wald chi-square tests) of a generalized linear mixed model with binomial error structure testing for the effects of sampling period, insect species and their interaction on the proportion of pollen from woody plants used by insects

<table>
<thead>
<tr>
<th>Predictor</th>
<th>χ²</th>
<th>df</th>
<th>p(χ²)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Insect species</td>
<td>3.85</td>
<td>3</td>
<td>.279</td>
</tr>
<tr>
<td>Sampling period</td>
<td>143.9</td>
<td>3</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>Insect species:Sampling period</td>
<td>15.59</td>
<td>9</td>
<td>.076</td>
</tr>
</tbody>
</table>

Note: Pollen use by Bombus terrestris, Osmia bicornis, Chrysoperla carnea and Harmonia axyridis was assessed during four sampling periods (roughly April, May, June and July; see Table S3). Significant values are highlighted in bold.

Cane, & Buchmann, 2000), and some abundantly flowering, pollen-rich herbaceous plants including Ranunculus, which has been shown to be toxic to other bee species but not for O. bicornis (Sedivy, Müller, & Dorn, 2011).

**FIGURE 4** Mean proportion of pollen from woody plant taxa collected per landscape sector for each insect species and sampling period. The four sampling periods (expressed in Growing Degree Days) correspond roughly to the months of April, May, June and July (see Table S3). Abbreviations: BT, Bombus terrestris; OB, Osmia bicornis; CC, Chrysoperla carnea; HA, Harmonia axyridis

4.2 | **Relative importance of pollen sources**

Despite the observed differences in pollen use among the studied insect species, our results reveal several general patterns. First, all four species used high proportions of pollen from non-agricultural plants (overall ≥64%), despite the fact that most of the studied landscapes were dominated by agricultural land (ranging from 38% to 90%; mean = 68%). The percentage of pollen from non-agricultural plants could exceed 90% in the case of O. bicornis and was generally high early in the year, underpinning the crucial role of floral resources offered by weeds and non-crop vegetation to sustain functionally important insect populations outside crops’ flowering periods (Requier et al., 2015). These percentages should be interpreted as conservative estimates since all pollen types that potentially include crop or sown grassland species were not considered. Evidence from North American agroecosystems support these findings for generalist pollinators (e.g. Russo & Danforth, 2017; Williams & Kremen, 2007).

Another important finding is that woody plant species play a crucial role in providing the studied insect species with...
been identified as a critical period during the life history of many species, such as intraguild predation (e.g. H. axyridis may prey upon C. carnea and conversely; Pell, Baverstock, Roy, Ware, & Majerus, 2008), which may lead to potential management conflicts. It should also be noted here that the invasive ladybeetle H. axyridis, despite being an effective aphid enemy, should not be specifically promoted by landscape management, because it threatens native biodiversity (and native ladybeetles in particular; Roy et al., 2012). However, our data show that due to its broad range of pollen use, and the strong dietary overlap with C. carnea, there appears to be little scope to selectively enhance either species by pollen resource provision. We hypothesize that the dominance of H. axyridis over native aphid predators can neither be countered nor exacerbated through pollen resource provisioning, but this topic requires further investigation.

### 4.3 | Seasonal shifts in pollen sources used

The four studied insect species exhibited similar temporal dynamics of pollen use during the season, characterized by a pronounced shift from woody to herbaceous pollen sources. This corroborates evidence of previous single-taxon studies focusing for example, on honeybees (Requier et al., 2015) or lacewings (Chrysoperla ssp.; Villenave et al., 2005). This pattern is likely at least partly driven by plant phenology: in most European agricultural landscapes, many dominant flowering trees and shrubs bloom relatively early and contribute more to floral resource availability early rather than later in the season (Williams et al., 2012; but see e.g. *Tilia* as an important exception). Yet, our results highlight the critical role of phenological complementarity among habitat types in providing food resources for multiple functionally important insect species throughout the year, and thus the importance of maintaining heterogeneous agricultural landscapes (Benton et al., 2003; Mallinger, Gibbs, & Gratton, 2016), including both species-rich woody and herbaceous habitats.

However, the high diversity of insects inhabiting agricultural landscapes that we could not investigate points to some limitations of our study. Further research is needed to confirm to what extent our findings could be generalized to natural bumblebee colonies and other important taxonomic groups of pollinators and crop pest enemies. Different functional groups such as parasitic wasps may show contrasting patterns of floral resource use (Patt, Hamilton, & Lashomb, 1997). Finally, monitoring over several years may help identify potential inter-annual variation in the observed pollen use patterns (e.g. due to varying phenologies of flowering plants and insects).

<table>
<thead>
<tr>
<th>Pollen Source</th>
<th>GDD 0–100</th>
<th>GDD 101–200</th>
<th>GDD 201–400</th>
<th>GDD 401–600</th>
<th>Whole sampling season (mean ± SD)</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Bombus terrestris</em></td>
<td>52%</td>
<td>68%</td>
<td>40%</td>
<td>69%</td>
<td>57% ±14%</td>
</tr>
<tr>
<td><em>Osmia bicorns</em></td>
<td>95%</td>
<td>82%</td>
<td>92%</td>
<td>58%</td>
<td>82% ±17%</td>
</tr>
<tr>
<td><em>Chrysoperla carnea</em></td>
<td>63%</td>
<td>63%</td>
<td>46%</td>
<td>51%</td>
<td>56% ±9%</td>
</tr>
<tr>
<td><em>Harmonia axyridis</em></td>
<td>77%</td>
<td>74%</td>
<td>50%</td>
<td>51%</td>
<td>63% ±15%</td>
</tr>
<tr>
<td>Mean</td>
<td>72%</td>
<td>72%</td>
<td>57%</td>
<td>57%</td>
<td>64% ±16%</td>
</tr>
</tbody>
</table>

Note: Numbers represent percentages of pollen types that can be unambiguously classified as from non-agricultural plants (i.e. pollen types potentially including crops and sown grassland plants are not included; see Table S1). The four sampling periods (expressed in Growing Degree Days) correspond roughly to the months of April, May, June and July (see Table S3).
5 | CONCLUSIONS

Our findings have several important implications for the promotion of multiple key insect species that can provide crop pollination and pest control services in agroecosystems. Among the most important findings is that pollen use overlapped only little among the four studied species, with the exception of Chrysoperla carnea and Harmonia axyridis. Nevertheless, the diet generally shifted from woody to herbaceous pollen collected from mostly weeds and non-agricultural vegetation sources. This strongly supports incentives to maintain or restore heterogeneous agricultural landscapes (Benton et al., 2003; Martin et al., 2019). Heterogeneous landscapes should be characterized by a high diversity of plants and vegetation types, including both woody and herbaceous vegetation providing complementary floral resources throughout the year (phenological completeness), a factor proposed to be critical for effective provision of pollination and pest control services in agroecosystems (Schellhorn et al., 2015).

Such incentives may include floral enhancement measures, such as the establishment of flower-rich hedgerows or sown flower strips promoted through agri-environmental schemes, which mitigate the isolation of semi-natural areas, and have a high potential to enhance floral-resource consuming insects and the ecosystem services they provide (e.g. Blaauw & Isaacs, 2014; M’Gonigle et al., 2015; Sutter et al., 2017; Tschumi et al., 2015). Our results, however, highlight an important role of trees for the nutrition of all four studied insect species in the early (Acer, Quercus, Salix, Fagus or Prunus) or late (Tilia) season, questioning whether schemes based on herbaceous plants alone can always provide enough adequate pollen resources to meet the pollen dietary requirements of multiple functionally important insect species (see also Wood, Holland, & Goulson, 2017). The identified early flowering trees are promising targets for habitat management measures to simultaneously promote both crop pollinators and pest enemies and the multiple ecosystem services they can provide in European agroecosystems.

ACKNOWLEDGEMENTS

This research was funded by the Swiss National Science Foundation (SNSF, grant number 6571210), and the German Research Foundation (DFG, grant number EN 979/3-1). We thank Julius Krebs, Jo Marie Reiff, Felix Wedel, Katrin Magin, Stephan Bosshart, Jonas Rudin, Dagmawi Dereje, Ives Nolle and Nicolas Hefti who have contributed to the field work; Martin Zuber, Jacqueline van Leeuwen, Sandra O. Brugger and Fabian Rey for their valuable help with pollen sample preparation and identification; and the farmers for field work allowance.

AUTHORS’ CONTRIBUTIONS

C.B., M.A., L.A., M.H.E., L.M. and F.H. conceived and designed the experiment. C.B., L.A. and L.M. performed the field surveys. C.B., P.E., E.G. and W.T. contributed to pollen sample preparation and identification. C.B. and M.A. performed the analysis. C.B. wrote the first draft of the manuscript, and all authors contributed substantially to writing. All authors gave final approval for publication.

DATA AVAILABILITY STATEMENT

Data are available from the Dryad Digital Repository https://doi.org/10.5061/dryad.6836p06 (Bertrand et al., 2019).

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**SUPPORTING INFORMATION**

Additional supporting information may be found online in the Supporting Information section at the end of the article.

**How to cite this article:** Bertrand C, Eckerter PW, Ammann L, et al. Seasonal shifts and complementary use of pollen sources by two bees, a lacewing and a ladybeetle species in European agricultural landscapes. *J Appl Ecol*, 2019;00:1–12. https://doi.org/10.1111/1365-2664.13483