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Spontaneous flowering vegetation favours hoverflies and parasitoid wasps in

apple orchards but has low cascading effects on biological pest control

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Abstract

Floral resources support several ecosystem services in agroecosystems, such as pollination or

biological control. Many beneficial organisms feed on nectar or pollen for an important part of their

12 life cycle. Providing adequate and sufficient floral resources through the conservation of

spontaneous flowering vegetation may be a strategy to improve biological pest control. However, the

role of spontaneous flowering plant species has mainly been evaluated on the recruitment of natural

enemies. Cascading effects on pest regulation and damage reduction are rarely studied.

Here we evaluated the effect of spontaneous flowering vegetation on pest regulation in 18

Mediterranean apple orchards. We focused on two main apple pests, *Cydia pomonella* and *Dysaphis*

plantaginea, and on two groups of their natural enemies depending on floral resources: hoverflies

and parasitoid wasps. We combined generalised linear mixed models and piecewise structural

equation models to test for direct and indirect effects of spontaneous flowering plant species on

beneficial insects (hoverflies, parasitoid wasps, wild bees and honeybees) and on pest regulation. We

also tested for potential negative interactions between honeybees and hoverflies, parasitoids wasps

or wild bees.

 There was a positive and significant effect of insect-pollinated flowers on hoverflies, parasitoid wasps and bees, but small cascading effects on *D. plantaginea* or *C. pomonella* density and associated damages. There was no evident relationship between honeybees and hoverflies, parasitoid wasps or wild bees. The reduction of *D. plantaginea* infestation was partially mediated by hoverfly abundance. Furthermore, we observed effects of spontaneous flowering vegetation on pest regulation and damage reduction independent of hoverfly and parasitoid wasp abundances. These results highlight the relevance of conserving floral resources to support biological control of apple pests. Further studies are needed to better understand interactions between spontaneous flowering vegetation and crop management practices to promote sustainable pest regulation strategies. **Key-words:** resident vegetation, wildflower, hoverflies, hymenopteran parasitoids, conservation

biological control

1. Introduction

 Arthropods are a key component of agroecosystem functioning. Many arthropod species provide important services to agriculture, such as crop pollination or biological pest control (Dainese *et al.*, 2019; Requier *et al.*, 2023). However, the current decline of arthropods (Seibold *et al.*, 2019) may compromise such ecosystem services (Zhou et al., 2023). Several causes have been identified to explain arthropod decline and most are related to agricultural intensification (Sánchez-Bayo & Wyckhuys, 2019; Wagner *et al.*, 2021). Thus, there is a clear need for more sustainable farming systems that preserve ecosystem services. Conservation Biological Control (CBC) is an important approach to improve sustainability by providing habitats for natural enemies of pests (Shields *et al.*, 2019).

 Several taxa of natural enemies feed on floral resources such as nectar and pollen during an important part of their life cycle (Lu et al., 2014). Nectar availability is positively related to the longevity of adult parasitoid wasps (Hymenoptera) in general and specifically increases the fecundity of synovigenic parasitoids (Russell et al., 2015; Benelli et al., 2017). The effect of floral resources on predators depends on feeding strategies (He et al., 2021). Hoverflies (Diptera: Syrphidae) and lacewings (Neuroptera: Chrysopidae) are predators at the larval stage, whereas adults feed primarily on floral resources which strongly affect their longevity and fecundity (He et al., 2021). Other predators, such as adult ladybugs (Coleoptera: Coccinelidae), are omnivorous and feed on floral resources when access to other resources is limited (Wolf et al., 2018; He & Sigsgaard, 2019). To promote natural enemies, a common habitat management strategy is to establish flowering plant species within or around crop fields (Landis et al., 2000). Nectar is mainly provided by insect- pollinated plants (Roy *et al.*, 2017). Therefore, insect-pollinated flowers are mainly sought to favour beneficial insects for pest biological control (Albrecht *et al.*, 2021a; Kirmer *et al.*, 2018). Planting such flowering vegetation often improves pest regulation but the increase is highly variable across studies (Albrecht *et al.*, 2021b ; Fountain, 2022).

 Spontaneous vegetation meaning ruderal and weed vegetation that grow naturally in or around crop fields in field margins or inter-rows of permanent crops can also provide important resources to pest natural enemies. Spontaneous vegetation is usually less costly for farmers to manage, better adapted to local conditions and can be as effective as standard flowering seed mixtures to promote beneficial insects (Blaix et al., 2018 ; Araj *et al.*, 2019). Balfour & Ratnieks (2022) showed that the abundance and diversity of beneficial insects were twice as high on spontaneously occurring weeds than on plant species of a recommended mixture to attract pollinators. Mei *et al.* (2021) also found that the abundance of arthropod predators mainly relied on the cover and diversity of spontaneous flowering plant species in field margins and not on the establishment of flower strips. Yet, the effects of spontaneous flowering vegetation on biological control services are often evaluated based on the recruitment of natural enemies (*e.g.* Blaix & Moonen, 2022; Denis *et al.*, 2021; Rosa García & Miñarro, 2014). The effect of spontaneous flowering vegetation on pest abundance and damage reduction has not always been evaluated or results were inconsistent (Johnson et al., 2020). Cascading effects of floral resource availability on pest regulation and reducing crop damage need to be better understood.

 Flowering plant species provide resources for different arthropod taxa, also including pests and pollinators that may compete for nectar or pollen (Boetzl et al., 2021). Several studies highlighted competition between honey bees and wild bees to access floral resources (Wojcik et al., 2018). Competition may also occur between pollinators and natural enemies (Campbell et al., 2012; Jeavons et al., 2020) or between natural enemies (Jeavons et al., 2022). However, these interactions remain 80 poorly documented and their effects on biological control services are not well known (Jeavons et al., 2022).

 Here, we used Mediterranean apple orchards as a model system to study effects of spontaneous vegetation on biological control. Apple is the most harvested and exported fruit in Europe (Eurostat, 2017) and apple orchards are the most treated crop in France (Jacquet et al., 2022). Numerous insect pests attack apple trees throughout the growing season. The codling moth (CM), *Cydia pomonella*, Linneaus 1758 (Lepidoptera: Tortricidae) and the rosy apple aphid (RAA) *Dysaphis plantaginea*, Passerini 1860 (Hemiptera: Aphididae) are the major insect pests, causing high economic losses in 88 apple production. We focused on parasitoid wasps and hoverflies, two groups which include many species that are CM and RAA natural enemies. Several parasitoid species feed on codling moth (CM) at different stages of its development, such as *Ascogaster quadridentata* Wesmael (Hymenoptera: Braconidae), *Pristomerus vulnerator* Panzer (Hymenoptera: Ichneumonidae) or *Trichomma enecator* Rossi (Hymenoptera: Ichneumonidae) (Mills et al., 2005). Other parasitoid species such as *Aphidius spp.* and *Ephedrus spp.* (Hymenoptera: Braconidae) attack the rosy apple aphid (RAA) (Dib e al., 2010; Rodriguez-Gasol et al., 2019). Most hoverflies found in crop fields including apple orchards are aphidophagous (Rossi *et al.*, 2006 ; Dib et al., 2010 ; Wojciechowicz-Żytko & Wilk, 2023). Hoverflies and parasitoid wasps arrive early in the season and can therefore control early occurring pest insects (González *et al.*, 2022; Rodríguez-Gasol *et al.*, 2019).

 In apple orchards, many studies analysed the effects of increased cover and/or diversity of flowering plant species on natural enemy abundance as indicator of biological control. Flowering plantings, such as flower strips or cover crops, have often a positive effect on natural enemies (Herz et al., 2019). In comparison, the effects of spontaneous flowering vegetation are less documented (Herz et al., 2019). Among studies including vegetation effects on pest insects, most of them focused on a single apple pest, often an aphid species (e.g. Rodriguez-Gasol et al., 2019; Gontijo et al., 2013; Santos et al., 2018; Albert et al., 2017). Few studies analysed the integrative effects of non-crop vegetation on the dynamics of different pest species, on fruit damage, and on interactions between natural enemies and pollinators in apple orchards (but see Cahenzli et al., 2019 and Martínez-Sastre et al., 2021).

 The present study aimed to analyse the effects of spontaneous flowering vegetation on the recruitment of hoverflies, parasitoid wasps and bees and to evaluate cascading effects on CM and RAA infestation and damage. We hypothesised that the cover of resident flowering entomophilous species increases the abundance of these insect groups, whereas anemophilous species do not have a positive effect. We expected that an increase in hoverfly and parasitoid wasp abundances has a negative effect on pest density which in turn reduces the associated damages on apple fruits. We also tested for potential interactions between hoverflies, parasitoid wasps, wild bees and honeybees, which could negatively affect pest regulation. Finally, we combined mixed-effects models and structural equation models to assess complex relationships between spontaneous vegetation, the two natural enemy groups, apple pests and associated damages and to disentangle direct and indirect cascading effects.

2. Material and methods

2.1 Study sites

121 The study was conducted in the lower Durance valley (area about 200 km², central point: WGS84: 43°8' N, 3°9' E). The study area was an agricultural plain dominated by orchards of which apple production represents 87% of the orchard area. Orchards are bordered by windbreak hedgerows (dominated by cypress or poplar) to protect apple trees against strong north wind often occurring in the study area. Botanical and entomological surveys were conducted in 18 apple orchards for commercial production (Fig. 1): nine orchards were under organic management and the other nine were under integrated pest management (hereafter defined as conventional orchards). Organic and conventional orchards had similar average areas, and surrounding vegetation (Table A.1). The 129 average number of treatments was also similar in organic and in conventional orchards (27.1 \pm 4.08 130 and 23.3 ± 6.63 in 2021, respectively), with insecticides representing about one third of the treatments. One or two broad-spectrum insecticides included chlorantraniliprole, pyrethroid or emamectin in conventional orchards and spinosad or neem oil in organic orchards (see Bouvier et al. 2022 for detail). Herbicides were applied under apple trees (intra-row) once a year in most

 conventional orchards. None of the orchards were covered with a protection net against the codling moth.

2.2 Botanical and entomological surveys

 Surveys of spontaneous flowering vegetation and some beneficial insects (hoverflies, parasitoid wasps and bees) were carried out in three periods: late March, early May, and early June of 2021. These observation periods coincide with the hatching of RAA eggs laid in the previous autumn (late March to early April), with RAA infestation peak (early May), and with the egg laying of the first CM generation (May and June). In each orchard, two transects of 2 x 10 m along two different rows of apple trees were established (Fig. 1): one transect was in the orchard centre and the other one close 143 to a hedgerow, at the northern edge. The 2 m width was measured from the base of tree trunks and transects covered thus half of the row (below trees) and the inter-row (alley). The rows were usually tilled or treated with herbicides in the beginning of the season. The inter-rows were mown every four 146 to ten weeks. Regrowth in rows resulted in low vegetation cover with a dominance of annual species whereas inter-row vegetation was usually dense and dominated by perennial species of disturbed 148 grasslands. Within orchards, the distance between two transects was on average 20.6 (\pm 10.4) m (Table A.1).

2.2.1 Flower cover and diversity

 All flowering plant species were recorded, including grasses, in each transect and at each period. We additionally surveyed the cover of each non-flowering plant species at the June period. For each plant species, we estimated the cover as the vertical projection of above-ground plant organs and the percentage of flowering individuals. Flower cover was then calculated by multiplying total cover by the flowering percentage in each species. All recorded flowering plant species were divided into two main groups: wind-pollinated and insect-pollinated flowers (Table A.2). Insect-pollinated flowers provide nectar whereas wind-pollinated flowers are not nectariferous. We assigned pollination modes using the BiolFlor database (Klotz *et al.*, 2002). For both plant groups, we calculated total

 flower cover and Shannon diversity, using the relative flower cover of each species in a given transect.

2.2.2 Hoverflies, parasitoid wasps and bees

 We counted hoverflies, parasitoid wasps, wild bees and honeybees on the same day as the botanical surveys using two complementary methods. First, two observers directly recorded all individuals of the four insect groups in each transect for 5 minutes, moving slowly along and taking care not to shade the transect. Insects were recorded when they landed on or hovered over the transect. 166 Second, we used a hand net to collect parasitoid wasps from the vegetation (10 strokes per transect). Netting allowed us to more accurately measure the abundance of parasitoid wasps, which are more difficult to detect by direct observation than hoverflies and bees. Samples were kept in 70° ethanol in 8mL tubes and sorted at the laboratory to count all parasitoid wasps without further taxonomic investigation (Goulet *et al.*, 1993). Observations and captures were carried out at days with low wind speed (<15km/h), between 10am and 4pm and temperatures higher than 15°C. Counts from visual observations and net samples collected at the same day were added to estimate the abundances of hoverflies, parasitoid wasps, wild bees and honeybees.

2.3. Insect pests and associated damage

 We focused on two apple pest insects: the rosy apple aphid, *Dysaphis plantaginea* (RAA) and the codling moth, *Cydia pomonella* (CM). RAA infestation was measured by counting the number of infested and active shoots on five trees per row at the three botanical periods. In late June, RAA and CM damage were evaluated on 16 trees per row. We randomly selected 20 apples per tree and measured the proportion of apples with signs of CM damage (*i.e.* feeding holes, stings and frass) or RAA damage (*i.e.* distorted apples). Pest infestation and damage were analysed at medium canopy height at 1.5 m to 2m. In late June, we placed wide corrugated cardboard bands and wrapped them around the trunk of 10 trees per row to capture mature CM larvae. Cardboard traps were removed in October and we counted all diapausing CM larvae.

2.4. Statistical analyses

- All analyses were conducted using R version 4.2.0 (2022-04-22). All models included period and
- orchard as random factors to account for temporal and spatial autocorrelation (Goulnik et al., 2020).

2.4.1. Linear mixed models

 Vegetation management and disturbance may differ between organic and non-organic orchards and according the location within the orchard. Therefore, we analysed the effects of orchard management (organic *vs* conventional) and of the transect location within the apple orchard (centre *vs* edge) on vegetation cover and diversity using linear mixed models (*lmer*, package: lme4 version 1.1-31, Bates *et al.*, 2015). We used five response variables: total vegetation cover, the cover of wind-pollinated flowers, the cover of insect-pollinated flowers, the Shannon index of wind-pollinated flowers and the Shannon index of insect-pollinated flowers. Transect location within orchards and orchard management were considered as fixed factors. Flower cover was square-root transformed for both plant groups to meet assumptions of normality and homoscedasticity. Models were tested using a type II analysis of variance.

2.4.2. Generalized linear mixed models

 Generalized linear mixed models (GLMMs) were used (*glmer*, package: lme4) to assess (i) the effects 200 of vegetation on the abundance of the four groups of beneficial insects and (ii) the effects of hoverflies and parasitoid wasps on pest infestation and damage. In the first set of GLMMs, response variables were the abundance of hoverflies, of parasitoid wasps and of wild bees or honeybees. Explanatory variables were the cover and the diversity of wind-pollinated and insect-pollinated flowers. In the second set of GLMMs, response variables were the number of RAA colonies, the proportion of apples with RAA damage, the proportion of apples with CM damage and the number of diapausing CM larvae. In aphid-related models, explanatory variables were parasitoid wasp abundance and hoverfly abundance (fixed effects). Hoverflies were not included in codling moth-related models. All GLMMs included location and orchard management as fixed factors. We tested

 for correlation between all variables (Table A.3) and checked for collinearity between explanatory variables using variance inflation factor (VIF) analyses. All VIFs were lower than 2 (Table A.4 and A.5). All GLMMs were built with a Poisson distribution and log-link function. Model fit and over-dispersion were checked graphically (package: DHARMa version 0.4.6, Hartig, 2022). In the case of over- dispersion, an observation-level random factor was added (Harrison, 2014). For each GLMM, we used the *dredge* function (package: MuMIn version 1.47.1*,* Bartoń, 2022) to

obtain all possible subsets of the maximal model and calculated the Akaike weight (*wi*) of each

model. Then, a model-averaging approach was applied (Grueber *et al.*, 2011). The relative

importance of each explanatory variable corresponds to the sum of the weights of all the models in

which the variable was included. Explanatory variables were considered significant when their

relative importance was greater than 0.6 and when 95% confidence intervals of the estimates did not

include zero (Burnham & Anderson, 2002). All explanatory variables were standardized allowing a

comparison of size effects among averaged models.

2.4.3. Structural equation models

223 We used piecewise structural equation modelling (pSEM), to test for direct and indirect effects of spontaneous vegetation on beneficial insects (hoverflies, parasitoid wasps, wild bees and honeybees) and pest regulation using the *psem* (package: piecewiseSEM version 2.3.0, Lefcheck, 2023) and *glmer* functions. The pSEMs examine multivariate relationships between interrelated variables in a single causal pathway (Lefcheck, 2016). We constructed two separate pSEMs to avoid overparameterization using orchard and period as random factors to account for unexplained variation by the fixed factors. 229 The first pSEM aimed to evaluate potential competition effects between honeybees and the three 230 other insect groups. The second pSEM examined the cascading effects of spontaneous vegetation on CM and RAA regulation and assessed whether these effects were mediated by hoverflies and parasitoid wasps.

 Both initial pSEMs were *a priori* based on known relationships between flower cover and diversity, hoverflies, parasitoid wasps and pest regulation, and on GLMMs preliminary results (Fig. A.1). Flower cover and diversity of both plant groups were included as exogenous variables not depending on other variables (no paths leading to them). The first pSEM included the four groups of beneficial insects as endogenous variables, and tested for a possible negative effect of honeybees on three other beneficial groups (hoverflies, parasitoid wasps and wild bees). In the second pSEM, we did not include the two bee groups but we added four endogenous variables related to RAA and CM infestations or damages. We expected cascading effects, with nectariferous flowers affecting hoverflies and parasitoid wasps and these natural enemies affecting RAA and CM densities. We hypothesised that the number of RAA colonies in spring would affect the fruit damage caused by RAA in June. Furthermore, we expected that fruit damage caused by the first CM generation measured in June explain the number of CM diapausing larvae collected in October. Significant missing paths in the initial pSEMs were detected by the d-separation test and added to the initial model (Lefcheck *et al.*, 2016). The overall model fit was assessed using Fisher's C statistic (p>0.05). Finally, a jack-knife like approach was used to evaluate the robustness of the pSEMs. Eighteen sub-models were built, based on a set of 18 independent orchards by omitting one different orchard in each sub-model. We ran each sub-model separately and identified paths that remained significant in at least 15 out of the 18 sub-models (more than 80%).

3. Results

3.1. Flower cover and diversity

 The total cover of spontaneous vegetation was 67.4 (± SE: 3.4)% leaving 32.6% of bare soil. The vegetation cover was significantly higher in the centre than at the edge of the orchards (+23.4 %) (Table 1). Over the whole season, the cover of wind-pollinated flowers varied from 1.6% to 20.5% 256 and was on average 9.2 (\pm 1.2)% (Table A.6). It increased during the season and was 1.0 (\pm 0.5)% in 257 late March, 10.7 (\pm 1.7)% in early May and 16.0 (\pm 2.6)% in early June (Fig. A.2). The cover of insect258 pollinated flowers per transect varied from 0.0% to 10.5% and was on average 4.6 (± 0.6)% (Table 259 A.6). It remained constant throughout the season, with a cover of 5.30 (\pm 1.03)% in late March, 3.5 260 (\pm 0.7)% in early May and 5.1 (\pm 1.1)% in early June. Insect-pollinated flower cover was twice as high 261 in the centre of the orchard as in the orchard edge (Table 1). Wind-pollinated flower cover was 2.45 times higher in organic orchards than in conventional orchards (Fig. A.3), but we did not find any significant effect of orchard management on the cover of insect-pollinated flowers (Table 1).

 Over the whole season, we recorded 112 plant species of which 78 were flowering at least at one of 265 the three observation periods. We observed 6.7 (\pm 0.4) flowering species per transect and per period. In late March, the most frequently flowering species were insect-pollinated plants: *Veronica persica* (Scrophulariaceae), *Taraxacum officinale* (Asteraceae)*, Cardamine hirsuta* (Brassicaceae)*, Senecio vulgaris* (Asteraceae) and *Crepis sancta* (Asteraceae)*.* In early June, the most common flowering plants were wind-pollinated species: *Lolium rigidum* (Poaceae), *Plantago lanceolata* (Plantaginaceae), and *Lolium perenne* (Poaceae). The dominant insect-pollinated species in early June were clover species: *Trifolium repens* and *T. pratense* (Fabaceae). The Shannon index of wind- pollinated flowers increased from March to June. For insect-pollinated flowers, the Shannon index remained constant throughout the season (Fig. A.2). The Shannon index of wind-pollinated species was significantly higher (+ 46%) in organic orchards (Fig. A.4), but we did not find any significant

275 effect of the orchard management on the diversity of insect-pollinated species (Table 1).

3.2. Effects of spontaneous flowering vegetation on hoverflies, parasitoid wasps and bees

277 Over the whole season, we recorded 423 beneficial insects by combining the two sampling methods 278 (hoverflies: 93, parasitoid wasps: 144, honey bees: 61 and wild bees: 125). We counted 3.9 (± 0.5) insects per transect and per period. Beneficial insect abundance increased throughout the season, with a total of 31 individuals being recorded in late March, 144 in early May and 248 in early June (Table A.7). GLMM results showed that the flower cover of insect-pollinated species had a significant and positive effect on the abundance of all insect groups (Table 2), with the highest effect size for

 honeybees (Fig. 2). The diversity of insect-pollinated species had only a significant and positive effect on wild bees (Table 2). The cover of wind-pollinated species had a significant and positive effect on the abundance of hoverflies and parasitoid wasps. The diversity of wind-pollinated species also positively affected the abundance of hoverflies and wild bees (Table 2). We did not find a significant effect of location within the orchard or orchard management on either insect group (Fig. 2). Piecewise SEM showed that honeybee abundance did not have a significant effect on any other group of beneficial insects (hoverflies: *p* = 0.47, parasitoid wasps: *p* = 0.19, wild bees: *p* = 0.79; Fig 4. A.). PSEM also confirmed the positive and significant effect of insect-pollinated flower cover on hoverfly and parasitoid wasp abundance (hoverflies: *p* = 0.01; parasitoid wasps: *p* <0.01; Fig 4. A.). However, nectariferous flower cover explained only a small part of the variation in hoverfly and 293 parasitoid wasp abundances ($R²m < 0.15$; Fig. 4).

3.3. Effects of hoverflies and parasitoid wasps on pest regulation

295 We counted 0.77 (± 0.13) RAA infested shoots per tree and per period, varying from 0 to 28. The 296 infestation was very low in late March (0.08 \pm 0.03) and increased throughout the season (1.05 \pm 0.22 in early May and 1.22 ± 0.34 in early June). GLMMs showed that hoverfly abundance had a significant negative effect on RAA infestation, and we observed a positive and significant relationship between parasitoid wasp abundance and RAA infestation (Fig. 3 and Table 3). In late June, the proportion of 300 apples with RAA damage was 1.0 (\pm 0.4) % and varied from 0 to 8.9 %. None of the tested explanatory variables had a significant effect on RAA fruit damage (Fig. 3 and Table 3). The proportion of apples with CM damage was 5.5 (± 2.6) %, attaining up to 63.6%, and depended mainly 303 on orchard management (Fig. 3). In October, we collected 0.63 (\pm 1.66) CM larvae per tree. A low proportion of small CM larvae was observed (5.7 ± 3.4 % per orchard), indicating parasitism by *A. quadridendata* (Table A.6.2). However, we did not observe any significant effect of parasitoid abundance on CM damage or on the number of CM diapausing larvae (Fig. 3 and Table 3).

3.4. Effects of spontaneous flowering vegetation on insect pests and on associated damage

 PSEM showed significant effects of spontaneous vegetation on pest infestation and associated damage reduction that were independent of hoverfly and parasitoid wasp abundances (Fig. 4. B). The flower cover of insect-pollinated species had a significant negative effect on the proportion of CM damage (*p* < 0.001). The cover of wind-pollinated flowers was the best predictor of RAA infestation showing a significant negative effect (*p* < 0.001). In contrast, we observed a significant and positive effect of the insect-pollinated plant diversity (Shannon) on RAA infestation (*p* < 0.001). PSEM also showed a negative relationship between RAA infestation recorded in spring and the number of CM larvae collected in autumn (*p* = 0.04). However, this relationship was not robust among the observation sites, being significant in less than 80% of the sub-models (Fig. 4.B).

4. Discussion

 We analysed the effects of spontaneous flowering vegetation on the recruitment of hoverflies, parasitoid wasps, wild bees and honeybees in apple orchards and their cascading effects on the regulation of the codling moth and the rosy apple aphid. Our results showed that the cover of insect-321 pollinated flowers increased the abundances of these four insect groups. The cover of wind- pollinated flowers had also a positive effect on the abundances of hoverflies and parasitoid wasps. We observed a significant negative effect of hoverfly abundance on RAA infestation. The cover of insect-pollinated flowers had a negative effect on CM damage and the cover of wind-pollinated flowers reduced RAA infestation. However, this reduction of RAA infestation was only marginally mediated by hoverfly abundance.

The cover of insect-pollinated flowers had a positive effect on hoverflies, parasitoid wasps, wild bees

and honeybees, and insect-pollinated flower diversity also had a positive effect on wild bees.

Similarly, Bishop *et al.* (2023) found that wildflower cover in apple orchard field margins improved

natural enemy richness and abundance and Pollier *et al.* (2018) highlighted the importance of

flowering insect-pollinated plants close to wheat and oilseed rape fields. This is likely to be due to an

increase in available nectar resources (Mockford *et al.*, 2022). A lack of nectar in early spring is

 generally observed in farmland (Timberlake *et al.*, 2019), even with the addition of flower strips (Serée *et al.*, 2023). In our study, we observed a constant cover of insect-pollinated flowers throughout the survey periods including early spring. Some flowering plant species predominantly recorded in March (*e.g. Veronica persica*, *Taraxacum officinale*) are therefore promising candidate species to improve seed mixtures, as they provide accessible nectar for several groups of natural enemies including parasitoid wasps (Gardarin *et al.*, 2021; Serée *et al.*, 2023). We also found positive effects of wind-pollinated species on hoverflies and parasitoid wasps. Although we did not specifically analyse flower visits and pollen consumption, the positive effect on hoverflies may be explained by pollen provisioning. Several studies have reported evidence of hoverflies foraging on grass species (Saunders, 2018; Joseph *et al.*, 2020; Villa *et al.*, 2021). Furthermore, hoverfly and parasitoid females may forage on weeds, searching for alternative prey to lay their eggs (Rodriguez- Gasol et al., 2019). Finally, the high cover of wind-pollinated flowers may indicate low vegetation disturbance (Piqueray *et al.*, 2019), which may also benefit natural enemies (Horton *et al.*, 2003; Meyer *et al.*, 2019).

 Piecewise SEM showed that spontaneous flowering vegetation explained at most 22% of the variability in beneficial insect abundances and 18% of the variability in pest infestation. This result may be due to the relatively low variability of insect-pollinated flower cover between orchards (0 to 10%). Competition between flower visitors may also limit the effect of floral resources on pest biological control (Jeavons *et al.*, 2022), but, we did not observe any relationship between honeybee abundance and that of the three other beneficial insect groups studied here. Effect of spontaneous vegetation on pest biological control may further be limited by other farming practices in apple orchards such as mowing, tillage and pesticide treatment frequencies (McKerchar *et al.*, 2020). However, abundances of hoverflies, parasitoid wasps or bees did not significantly differ between conventional and organic apple orchards in our study. A more detailed description of the farming practices in each apple orchard independently of the organic *vs.* conventional management typology would help to precise their significance. Landscape composition may also affect natural enemy

 abundance, in particular the cover of semi-natural habitats, including hedgerows and their proximity to field crops (Maalouly *et al.* 2013; Daelemans *et al.*, 2022; Bishop *et al.*, 2023). The overall abundance of insects (hoverflies, parasitoid wasps and bees) was rather low, which may have limited the strength of observed effects. An increased observation time may be a solution to obtain higher abundances avoiding 0 values that reduce statistical power.

 Hoverfly abundance had a negative effect on RAA infestation. This confirms the potential of hoverflies to improve aphid biological control (Rodríguez-Gasol *et al.*, 2020). Hoverflies were not identified to species level but previous studies showed that most hoverfly species occurring in apple orchards are aphid predators (Rossi et al., 2006; Dib et al., 2010; Wojciechowicz-Żytko & Wilk, 2023). We did not find a significant relationship between parasitoid wasps collected on spontaneous vegetation and the abundance of codling moth larvae or their associated damage. We also recorded a very low number of small larvae (indicating codling moth parasitism). The RAA infestation was positively related to the abundance of parasitoids sampled in the vegetation. This observation suggests that RAA and parasitoid wasps are influenced by common factors not included in our analysis. The density of hedgerows in the monitored orchards that favours both parasitoid diversity (Bishop et al., 2023) and RAA infestation (Albert et al., 2017) may be such a common factor. We did not distinguish parasitoid wasps (as in Mateos-Fierro et al., 2021; Pollier et al., 2018; Rodriguez-Gasol et al., 2019), assuming that the overall abundance of parasitoids is positively related to the abundance of parasitoid species attacking the apple pests RAA and CM. However, it is possible that parasitoid wasp species collected on spontaneous vegetation were not involved in apple pest regulation. The identification of parasitoid wasps would thus be required to determine whether flowering plants in apple orchards attract parasitoid species that are really involved in pest regulation.

 Piecewise SEM showed effects of spontaneous flowering vegetation independent of hoverflies or parasitoid wasps. The cover of insect-pollinated flowers negatively affected CM damage, and wind pollinated flower cover had a negative effect on RAA infestation. These effects may be related to the activity of other groups of natural enemies than parasitoid wasps and hoverflies. Generalist predators such as earwigs (Dermaptera: Forficulidae), spiders (Araneae) highly contribute to RAA and CM regulation (Dib *et al.*, 2020; Unruh *et al.*, 2016). Additionally, ladybirds (Coleoptera: Coccinelidae) or flower bugs (Heteroptera: Anthocoridae) are involved in RAA regulation (Dib et al., 2010). All these predator groups may benefit from spontaneous flowering vegetation without directly consuming floral resources. For example, Mei *et al.* (2021) showed that spiders and ground beetles were positively affected by wildflower cover and diversity even though they do not consume nectar or pollen. Such generalist natural enemies benefit from shelter, favourable microclimate and structural diversity provided by a relatively undisturbed herbaceous layer (Ganser *et al.*, 2019). We also found a small but positive effect of the diversity of insect-pollinated flowers on RAA infestation. This result suggests that both variables, insect-pollinated flower diversity and RAA infestation, were influenced by common drivers not included in our analysis. However, it is also possible that a higher flower diversity may attract a higher number of natural enemy species increasing potential intraguild predation or competition (Daelemans *et al.*, 2022).

 To conclude, our results demonstrated that spontaneous vegetation rich in insect-pollinated plants improves the recruitment of hoverflies, parasitoid wasps, wild bees and honey bees in apple orchards. Positive effects of spontaneous vegetation on pest regulation and damage reduction were weak but significant. To further increase the regulation of apple pests, the preservation of spontaneous flowering vegetation should be combined with other farming practices, such as sowing plant species within or around orchards that favour apple pest natural enemies. Natural enemy activity generally decreases with increasing distance from such flowering plantings (Albert et al., 2017; Santos *et al.*, 2018). Spontaneous vegetation may act as a relay, promoting the movement of hoverflies and parasitoid wasps from field margins within the orchard and should be managed at the whole-field level, rather than only at the edges in agreement with Serée *et al.* (2023). Apple cultivars selected for their low susceptibility to RAA (Alhmedi et al., 2022) or CM (Joshi et al., 2015) and

- management practices (pruning, irrigation, fertilisation) that reduce tree vigour and RAA infestation
- (Simon et al., 2007; Rousselin et al., 2018) could additionally reduce apple damages. In order to
- develop guidelines for farmers, future research should therefore examine interactions between
- spontaneous flowering vegetation and farming practices to promote sustainable pest regulation.

Authorship contribution statement

- 415 LL, AB, FL and PF conceived the ideas and designed the methodology; LL, AB and PF collected the
- data; LL and RB analysed the data; LL wrote the first draft of the manuscript. All authors contributed
- critically to the drafts and gave final approval for publication

Declaration of Competing Interest

 The authors declare that they have no known competing financial interests or personal relationships 420 that could have appeared to influence the work reported in this paper.

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Figures

Figure 1. Location of the 18 orchards.

- Figure 2. Effect sizes (mean ± 95% CI) of flower cover and diversity, location within the orchard, and
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- **Tables**
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- Table 3. Averaged estimated effects (GLMM) of hoverfly and parasitoid abundances, location and
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 Figure 1. Location of the 18 apple orchards in south-eastern France within the long-term study zone "Basse 729 Vallée de la Durance" ([https://site-atelier-basse-vallee-durance.fr\)](https://site-atelier-basse-vallee-durance.fr/). In each orchard, we defined two transects at

the center and at the edge (2m x 10m) in which we conducted botanical and entomological surveys. Transects

731 were separated by 20.6 m (± 10.4) .

 Figure 2. Effect sizes (mean ± 95% CI) of flower cover and diversity, location within the orchard, and orchard management on beneficial insect abundance. Estimates were obtained by a model averaging approach. Triangles correspond to wild bee abundance, circles to parasitoid abundance, diamonds to hoverfly abundance, and squares to honeybee abundance. Significant positive effects are indicated in green.

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 Figure 3. Effect sizes (mean ± 95% CI) of natural enemies, location within the orchard, and orchard management on pest infestations and damages. Estimates were obtained by a model averaging approach. Circles correspond to rosy apple aphid (RAA) infestation (monitored from April to June), squares to the proportion of apples with RAA damage (recorded in June), triangles to the proportion of apples with codling moth (CM) damage (recorded in June), and diamonds to the number of codling moth diapausing larvae (monitored in October). Positive and negative significant effects are indicated in green and red, respectively.

 Figure 4. Final piecewise structural equation models (pSEM). Model **A** shows the effects of spontaneous flowering vegetation hoverflies, parasitoid wasps, honeybees and wild bees. Model **B** shows the effects of spontaneous flowering vegetation on natural enemies and cascading effects on pest infestation and associated damages. I-P flowers: Insect pollinated flowers; W-P flowers: Wind pollinated flowers; RAA: Rosy Apple Aphid; CM: Codling moth. Green and red arrows represent significant positive and negative pathways, respectively, with arrow thickness corresponding to standardised path coefficients. Coefficients in parentheses correspond to 767 pathways significant in less than 80% of sub-models. Dotted grey arrows indicate non-significant paths (p>0.05). 768 Double-headed arrows illustrate correlated errors. $R²m$ values show the proportion of variability explained by 769 fixed effects and R²c values show the proportion of variability explained by both fixed and random effects. Overall model fit is indicated using Fisher's C statistic. Summary statistics for the pSEM models are shown in Table A.8 and A.9.

772 **Table 1. Results of the linear mixed models (LMM) analysing the effects of location and orchard**

773 \ldots management on cover and diversity of insect-pollinated flowers and wind-pollinated flowers. χ^2 : test

774 statistic, df: degrees of freedom. Positive estimates for "Location" correspond to an increase towards

775 "Edge". Positive estimates for "Orchard management" correspond to an increase towards "Organic".

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 Table 2. Averaged estimated effects (generalized linear mixed models, GLMM) of spontaneous flowering vegetation on beneficial insect abundance. Σ*wi*: sum of the weights (i.e. relative importance). Positive estimates for "Location" correspond to an increase towards "Edge". Positive estimates for "Orchard management" corresponds to an increase towards "Organic". Significant variables are indicated in bold.

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 Table 3. Averaged estimated effects (generalized linear mixed models, GLMM) of hoverfly and parasitoid abundances, location and orchard management on pest numbers (RAA infestation and CM larvae) and their damages. Σ*wi*: sum of the weights (i.e. relative importance). Positive estimates for "Location" correspond to an increase towards "Edge". Positive estimates for "Orchard management" corresponds to an increase towards "Organic". Significant variables are indicated in bold. RAA: Rosy apple aphid, CM: Codling moth.

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Appendix

- Table A.1. Characteristics of the 18 apple orchards.
- Table A.2. Assignment of a pollination mode to each recorded flowering plant species.
- Table A.3. Combined Pearson and Spearman correlation matrix.

 Table A.4. Variance inflation factor (VIF) for the GLMM analysing the effects of spontaneous flowering vegetation on beneficial insect abundance.

 Table A.5. Variance inflation factor (VIF) for the GLMM analysing the effects of hoverfly and parasitoid abundances, location and orchard management on pest numbers (RAA infestation and CM larvae) and their damages.

- Table A.6. Summary statistics for all measured variables.
-

Table A.7. Number of recorded hoverflies, parasitoid wasps, wild bees and honeybees.

809 Table A.8. Summary statistics of the piecewise structural equation model (pSEM) explaining the effects

- 810 of spontaneous flowering vegetation on beneficial insect abundance.
- Table A.9. Summary statistics of the piecewise structural equation model (pSEM) explaining the effects
- of spontaneous flowering vegetation natural enemies and cascading effects on pest infestation and associated damage.
- Figure A.1. Initial pSEMs and associated hypotheses.
-
- Figure A.2. Evolution of flower cover (%), species richness and diversity from March to June.
- Figure A.3. Comparison of flower cover between organic and conventional orchards and between 818 transect location within the orchard.
- Figure A.4. Comparison of flower diversity (Shannon index) between organic and conventional orchards and between transect location within the orchard.

Table A.1. Characteristics of the 18 apple orchards.

Table A.2. Assignment of a pollination mode to each recorded flowering plant species. Flowering phenology was determined from the present study. Pollination mode was determined based on the BiolFlor database using information from 'pollen vector' and 'typical pollinator' (Mueller, 1881) traits. For 'selfpollinated' plant species, information from 'typical pollinator' trait were used to assign the species to insect-pollinated or wind-pollinated groups. For plant species not included in the BiolFlor database (NA), we used the information available for other species of the same genus to assign a group.

Table A.2 – *continues from previous page*

Table A.2 – *continues from previous page*

Table A.3. Combined Pearson and Spearman correlation matrix. The area in grey corresponds to Pearson correlations. Correlation coefficients are indicated (r of Pearson or rho of Spearman) and stars indicate significance level ('*': <0.05; '**': <0.01; '***': <0.001).

Table A.4. Variance inflation factor (VIF) for the GLMM analysing the effects of spontaneous flowering vegetation on beneficial insect abundance.

Table A.5. Variance inflation factor (VIF) for the GLMM analysing the effects of hoverfly and parasitoid abundances, location and orchard management on pest numbers (RAA infestation and CM larvae) and their damages.

Table A.6.1 Summary statistics for all measured variables. For the variables related to spontaneous vegetation (flower cover and diversity), given data are means and standard deviation (SD) of the three periods. I-P flower: insect-pollinated flower, W-P flower: wind-pollinated flower.

Table A.6.1. Summary statistics for all measured variables. For the variables related to spontaneous vegetation (flower cover and diversity), given data are means and standard deviation (SD) of the three periods. I-P flower: insect-pollinated flower, W-P flower: wind-pollinated flower. **–** *continues from previous page*

Table A.6.2 Summary statistics for all measured variables. For the variables related to beneficial insects (abundance of each group and Shannon diversity), given data are mean and standard deviation of the three survey periods. **–** *continues from previous page*

Table A.6.2. Summary statistics for all measured variables. For RAA infestation, given data are the mean and standard deviation of the three periods. For damages, given data are proportion recorded in late June. CM larvae corresponds to the number of codling moth diapausing larvae collected in October (sum per transect) **–** *continues from previous page*

Table A.7. Number of recorded hoverflies, parasitoid wasps, wild bees and honeybees. Mean (±SD) corresponds to the mean number of insects recorded per transect and per period.

Table A.8. Summary statistics of the piecewise structural equation model (pSEM) explaining the effects of spontaneous flowering vegetation on beneficial insect abundance. Each row corresponds to a path in the pSEM. SE: Standard Error; DF: Degree of freedom; Std Estimate: Standardized estimate. **Table A.9. Summary statistics of the piecewise structural equation model (pSEM) explaining the effects of spontaneous flowering vegetation on natural enemies and cascading effects on pest infestation and associated damage.** Each row corresponds to a path in the pSEM. SE: Standard Error; DF: Degree of freedom; Std Estimate: Standardized estimate.

We hypothesized that:

(1) Insect-pollinated flowers (cover and diversity) improve the abundance of beneficial insects (Kishinevsky et al., 2017; Mei et al., 2021).

(2) Wind-pollinated flowers (cover and diversity) can promote beneficial insects, as a source of pollen or potential habitat (Dong et al., 2021; Saunders, 2018).

(3) Honeybees abundance negatively impacts other beneficial insect groups (Jeavons et al., 2020).

(4) Hoverflies have a negative effect on rosy apple aphid infestation (Moerkens et al., 2021; Dib et al., 2011).

(5) Parasitoids have a negative effect on rosy apple aphid infestation (Rodriguez-Gasol et al., 2019).

(6) Parasitoids observed in spring have a negative impact on the first generation of codling moth (Martinez-Sastre et al., 2021).

(7) RAA infestation conducts to distorted apples (i.e. sign of aphid damage) (Blommers, 2004).

(8) A high density for the first generation of codling moth conducts to a high number of larvae for the third generation.

(9) We observed a correlation between I-P flower cover and diversity as well as between W-P flower cover and diversity.

Thus, we included correlated errors.

Figure A.1. Initial pSEMs and associated hypotheses. A: Initial pSEM related to interactions between hoverflies, parasitoid wasps, wild bees, honeybees and spontaneous flowering vegetation; B: Initial pSEM related to cascading effects on pest regulation.

Figure A.2. Temporal dynamics from March to June of the percentage of flower cover (A), specific plant richness (B) and Shannon plant diversity (C).

Figure A.3. Comparison of flower cover between organic and conventional orchards and between transect locations within the orchard. Bars indicate standard errors. W-P flower: Wind-pollinated flower and I-P flower: Insect pollinated flower

Figure A.4. Comparison of flower diversity (Shannon index) between organic and conventional orchards and between transect location within the orchard. Bars indicate standard errors. W-P flower: Wind-pollinated flower and I-P flower: Insect pollinated flower.

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