

Environmental factors affecting insect pollinators and pests at urban farmlands: a case study with honey bees and cabbage white butterflies

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Rapid urbanization results in a significantly increased urban population, but also the loss of agricultural lands, thus raising a concern for food security. Urban agriculture has received increasing attention as a way of improving city citizens' food accessibility and local people's livelihood. Although vegetable-dominant small urban farmlands are relatively common in China, little is known about environmental factors associated with insects that could affect ecosystem services at these urban farmlands, which in turn influence agricultural productivity. Using Asian honey bee (*Apis cerana*) and cabbage white butterfly (*Pieris rapae*) as examples, I investigated how environmental features within and surrounding urban farmlands affected insect pollinator (bee) and pest (butterfly) abundance in a megacity of China during winters. I considered environmental features at three spatial scales: fine (5m-radius area), local (50m-radius area), and landscape (500m-radius and 1km-radius areas). While the abundance of cabbage white butterfly increased with local crop diversity, it was strongly negatively associated with landscape-scale crop and weed covers. Asian honey bees responded positively to flower cover at the fine scale. Their abundance also increased with local-scale weed cover but decreased with increasing landscape-scale weed cover. Although farmland fragmentation, i.e., patch density of farmland within a landscape did not affect cabbage white butterfly, the abundance of Asian honey bees tended to decrease with increasing farmland fragmentation. These results suggest that cultivating too diverse crops at urban farmlands can increase crop damage; however, the damage may be alleviated at farmlands embedded in a landscape with more crop cover. Retaining small amount of un-harvested flowering crops and weedy vegetation within a farmland, especially less fragmented farmland can benefit honey bees when natural resources are scarce.

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Abstract

Rapid urbanization results in a significantly increased urban population, but also the loss of agricultural lands, thus raising a concern for food security. Urban agriculture has received increasing attention as a way of improving city citizens' food accessibility and local people's livelihood. Although vegetable-dominant small urban farmlands are relatively common in China, little is known about environmental factors associated with insects that could affect ecosystem services at these urban farmlands, which in turn influence agricultural productivity. Using Asian honey bee (*Apis cerana*) and cabbage white butterfly (*Pieris rapae*) as examples, I investigated how environmental features within and surrounding urban farmlands affected insect pollinator (bee) and pest (butterfly) abundance in a megacity of China during winters. I considered environmental features at three spatial scales: fine (5m-radius area), local (50m-radius area), and landscape (500m-radius and 1km-radius areas). While the abundance of cabbage white butterfly increased with local crop diversity, it was strongly negatively associated with landscape-scale crop and weed covers. Asian honey bees responded positively to flower cover at the fine scale. Their abundance also increased with local-scale weed cover but decreased with increasing landscape-scale weed cover. Although farmland fragmentation, i.e., patch density of farmland within a landscape did not affect cabbage white butterfly, the abundance of Asian honey bees tended to decrease with increasing farmland fragmentation. These results suggest that cultivating too diverse crops at urban farmlands can increase crop damage; however, the damage may be alleviated at farmlands embedded in a landscape with more crop cover. Retaining small amount of un-harvested flowering crops and weedy vegetation within a farmland, especially less fragmented farmland can benefit honey bees when natural resources are scarce.

Introduction

Urban expansion in many countries is often accompanied by the loss of agricultural lands (Bren d'Amour et al., 2017). The world's built-up areas are a quarter of total agricultural lands; however, the expansion of built-up areas is faster than urban population growth and takes place on productive agricultural lands, leading to reduction in global crop production (Seto & Ramankutty, 2016; Bren d'Amour et al., 2017). Moreover, the proportion of global population in urban areas is forecast to increase from 55% in 2018 to 68% by 2050 (United Nations, 2019). Approximately 90% of the increase would occur in Asia and Africa, which are also the hotspot of future urban expansion. These changes, i.e., growing urban population and agricultural land loss, raise concerns for not only food security but also for the livelihoods of smallholders, especially in developing countries (IFPRI, 2017; Huang et al., 2020).

Urban agriculture has received increasing attention as one of the practices that may improve the accessibility of food, the livelihood of local people such as farmers and rural migrants, and human well-being (Deelstra & Girardet, 2000; De Bon et al., 2010; Mok et al., 2013; Martellozzo et al., 2014). One recent study also shows that small-scale urban agriculture can bring high yields (McDougall et al., 2018). Urban agriculture can be broadly defined as agricultural production, such as vegetables, fruits, ornamental plants, and other dietary products, occurring in urban environments that include both inner city and city fringe areas (Mougeot, 2000). In China, small-scale vegetable farming is common in cities, especially at vacant lots and city fringe areas. Urban agriculture, also called peri-urban agriculture in some cases, has been supported by the Chinese government policy "Vegetable Basket" for decades (Zong et al., 2021). Like other countries, urban expansion has been a major driver causing farmland loss in China (Tu et al., 2020). Although farmland protection policy started in mid-1990s has alleviated the loss to some extent, urban development frequently occurs on highly productive farmlands near urban edge areas, often leaves reclaimed but less productive farmlands as an offset for the development, and increases farmland fragmentation (Liang et al., 2015; Huang et al., 2019; Tu et al., 2021). In this situation, understanding ecological factors that could affect agricultural production at urban farmlands is crucial to promote the sustainability of these farmlands.

Insects are well known for their role in ecosystem service (e.g., pollination and pest control) and disservice (e.g., crop damage) in agricultural landscapes (Losey, 2006; Kremen & Chaplin-Kramer, 2007; Oliveira et al., 2014; Omark, 2018). Their effects on crop yields have been widely studied. Over 70% of main global crops benefit from animal-mediated pollination (Klein et al., 2007) and 9-25% of staple food crops are estimated to be lost due to animal pests (Oerke, 2006), in which insects are the major group in both cases. Bees are considered the most important pollinators. For example, in USA, 11% of the agricultural gross domestic product in 2009 depended on pollination, mainly contributed by bees (Lautenbach et al., 2012). Bee abundance affects yields of certain crops such as oilseed rape more than pesticides do (Catarino et al., 2019 and references therein). Diverse wild bees support pollination of multiple crops in urban gardens (Lowenstein et al., 2015). A number of factors, e.g., quantity and quality of floral resources, semi-natural vegetation, crop diversity such as crop compositional or configurational

heterogeneity and landscape complexity, influence the abundance and diversity of bees (Klein et al., 2012; Scheper et al., 2013; Potts et al., 2003; Priyadarshana et al., 2021; Raderschall et al., 2021).

In contrast to bees, cabbage white butterflies (e.g., *Pieris rapae* and *P. brassicae*) are recognized as agricultural pests because their larvae, cabbageworms, feed on the family Brassicaceae crops (cruciferous crops) and can severely damage these crops. For example, cabbage white butterflies caused about 70% yield loss in cruciferous crop in Meghalaya, the northeastern state of India (Singh et al., 2014). While presence or abundance of Brassicaceae crops is positively associated with cabbageworm density, this effect may be amplified if the host plants are congregated as a large pure stand (resource concentration hypothesis; Root, 1973) or sparsely distributed (resource diffusion hypothesis; Yamamura, 1999). The type of surrounding crop has an impact on cabbageworm density (Maguire, 1984) and distance between cruciferous patches influences egg density of cabbage white butterfly (Fahrig & Paloheimo, 1988). Floral resource availability can also be important to the adult butterflies (Curtis et al., 2015). Here, using Asian honey bee (*Apis cerana*) and cabbage white butterfly (*Pieris rapae*), I investigated how environmental features within and surrounding urban farmlands in a megacity of China influenced the abundance of these two species, during winter in which Brassicaceae crops are of particular prevalent. I focused on farmlands largely cultivating vegetables, which are dominant agricultural products of urban farming. I expected that the percentage of flowering plants and Brassicaceae crops would positively affect honey bees and cabbage white butterflies, respectively. Given that these farmlands are in urban areas mostly filled with built-up structures, landscape weedy vegetation (spontaneous vegetation in vacant lots or parks) as well as available farmlands could also be critical to both species. Crop diversity could negatively affect cabbage white butterflies as found in the effect of crop diversification on insect pests in general (Hooks & Johnson 2003; Beillouin et al., 2021). Alternatively, the effect may not be significant because urban farmlands in this study region often contain more diverse crops than in rural agricultural lands, and predator/parasitoid abundance is low in winter.

Materials & Methods

Study area

Guangzhou is the capital city of Guangdong Province, China (Fig. 1) and home to ~18 million peoples. It also lies within the Indo-Burma Biodiversity hotspot region, the most urbanized area among biodiversity hotspot regions in China (Güneralp & Seto, 2013). Guangzhou has a subtropical monsoon climate: warm and dry winter and hot and humid summer, with a mean annual temperature of 22.2°C (<https://en.climate-data.org>). During the past two decades, farmlands located in city center area and between the center and boundary areas have drastically decreased due to urban expansion and economic development (Shi & Shi, 2020).

I selected 33 farmlands in 2020 and 57 farmlands (23 from previous year) in 2021 (Fig.1). Several farmlands chosen in 2020 could not be surveyed in 2021 due to development, abandonment, and too-early growth stage of crops (very small seedlings). The size of all

farmlands was $\geq 7,850\text{m}^2$. Within a farmland, I established a 50m-radius area (site, hereafter) that was also used for bird surveys in another study (Lee et al., 2022). All samplings were performed within the 50m-radius area during January each year. Cabbage white butterfly was surveyed at 90 sites (32 in 2020 + 58 in 2021) across two winters. Bee sampling was conducted at 58 sites (25 sites in 2020 and 33 sites in 2021). Nearest distance between the centers of sites surveyed for cabbage white butterfly was $1.87 \pm 1.09\text{km}$ (mean \pm standard deviation) in 2020 and $0.82 \pm 1.18\text{km}$ in 2021.

I did not distinguish between urban and peri-urban agriculture, including farmlands located in the inner city as well as close to the city boundary (Mougeot, 2000). However, I emphasize that all sites were in urban areas; built-up cover within a 500m-radius area of sample sites averaged $61 \pm 13.8\%$, consistent with the criterion of urban areas containing $> 50\%$ build-up (Marzluff, 2000).

Cabbage white butterfly

Cabbage white butterfly (CWB) was surveyed twice (≥ 12 days apart) using a transect method: one survey in the morning (between 9-11:30am) and one in the afternoon (between 1:00-4:00pm). During each survey, I placed randomly two 50m-line transects along a walk path within a site. However, if Brassicaceae and non-Brassicaceae crop areas were clearly divided, I placed one transect close to more Brassicaceae crops and the other near non-Brassicaceae crops. I also changed the locations of transects between surveys. I walked the transect at a rate of 5m per minute, and counted CWB individuals detected within a 10m x 5m x 5m imaginary window (5m to each side of the transect, 5m from the ground, and 5m in front). Although a 5m width is commonly used, I expanded the width of the imaginary window because CWBs were often active slightly farther from the edge of path and moved along the row crops, which were not easy to access because of irrigated water and farmer's attendance at some sites. Special care was taken not to count the same individuals twice while walking along the transect. During the first winter, I also carried out another survey using an area search method to verify the transect method. I quickly walked around 50% area of a site, e.g., north-east-south side for 1 minute, and counted CWB. This scanning process was repeated three times at different sides of the site, with 3 minutes between surveys. Data collected from transect method and area search methods were highly correlated: Pearson's correlation (r) between maximum counts of each data was 0.94 ($P < 0.001$). Thus, I assumed that the transect survey data adequately represented CWB abundance in this study.

Bee sampling

To capture bees, I used three-colored (blue, white, and yellow) pan traps (Campbell & Hanula, 2007). Three sample stations were established $> 30\text{m}$ apart from each other within a site. Sample stations were placed randomly from the center of site. However, most locations were restricted by presence/absence of flowering plants (crops and weeds), farmer's permission, and accessibility (avoiding deep furrows filled with water). I installed a set of three-colored pan traps

at each sample station. I attached pan traps to a pole at ~ 0.8m high, which was tall enough to allow bees to see them, and placed them near flowering crops or weeds if they were present. Pan traps were 1/2-2/3 filled with soapy water (10 drops of dish soap per liter) to remove the surface tension. All traps were retrieved 2 days after installation. While I brought all bees captured to a lab, washed, and preserved in 70% ethyl alcohol for identification, I counted only Asian honey bees (*Apis* hereafter) because they were the main bee species captured and abundant enough to conduct analysis.

Environmental data at fine and local scales

Local scale environmental data were collected within a site, that is, “site” represents the local scale of this study. At each site, crop and non-crop features such as built-up structures (storage house and path), herbaceous weeds, trees/shrubs, and open water (small pond and water channel) were identified and marked on a printed satellite image downloaded from Google Earth (<https://www.google.com/earth/>). The images were taken between 2018 and 2020. Many farmlands were composed of highly diverse crop species, but each crop was cultivated in similar-sized rows. Thus, I divided the site into several blocks. I identified crops and counted the number of rows of each crop within a block. I summed all counts by block. In a block, I calculated the relative frequency of each crop from the sum and converted that to “area” by multiplying it by the size of the block. Sum of the crop’s area across all blocks represented total area of the crop within the site. All non-crop features and blocks were delineated in ArcGIS using georeferenced Google Earth images as a base map.

I calculated the percentage of weedy vegetation (weed50) and cruciferous crops (Brassicaceae; the mustard family). Crop diversity (cropdiv50) was calculated as a Shannon-Wiener diversity index. While over 120 crops were found across sites, several crops were very minor or variants of the same crop, sharing similar biological and ecological characteristics. I also found cases of misidentifications when crops look similar, especially when they belong to the same genus or family. Thus, I used the family of crop to calculate cropdiv50 (Table S1 for the list of crop genera). At the family level, all sites had ≥ 2 crops and approximately 95% of sites contained at least 6 crops, averaging 13.9 ± 5.4 , and ranging from 2 to 24.

For *Apis*, crop surveys were also performed at a fine scale, a 5m-radius area surrounding a sample station. Within the 5m-radius area, crops were identified and the percentage of each crop was visually estimated. Similar to cropdiv50, I calculated fine-scale crop diversity based on crop species (cropdiv5). In 2021, the percentage of flowering plants (flower) including crops and weeds was also estimated at the fine scale.

Landscape data

I choose two landscape-scale sizes by considering flight distances of CWB and *Apis* (Jones et al., 1980; Dyer & Seeley, 1991), the logistics of creating a land cover map, and the matrix context of study sites, selecting 500m-radius and 1km-radius areas surrounding the center of site. The 500m-radius area was used to examine compositional aspect of the landscape matrix in

which a farmland was embedded, and the 1km-radius area was used to characterize the spatial configuration and fragmentation of farmlands within a landscape.

To generate the land cover map, I downloaded satellite images from Google Earth and georeferenced them. Using the georeferenced images as a base map, I delineated 3 land cover types within the 500m-radius area in ArcGIS: vegetable-dominant farmland (crop cover), weedy vegetation (mostly spontaneous herbaceous vegetation in vacant lots and construction sites), and built-up structure (building, road, any impervious surface). I then calculated the percentage of farmland or crop cover (crop500) and weedy vegetation (weed500). I also delineated farmlands within the 1km-radius area and calculated patch density (pd1000) and edge density of farmlands (edge1000) using Fragstats v 4.2.1 (McGarigal et al., 2012). Although these indices may be the simplest measure of the spatial configuration of habitat patch (McGarigal & Marks, 1995), they can affect diversity and abundance of pollinators and butterflies by increasing edge habitats or facilitating movements between patches (Flick et al., 2012; Hass et al., 2018; Martin et al., 2019; among others). Edge density is the same as the total edge length in this study because landscape sizes were identical across all sites.

Statistical analysis

I pooled 2 years of data together for CWB; even if the same site was surveyed both years, each year's data were considered independent because local scale features such as crop diversity and amount of weedy vegetation differed between two winters at the same site. Any potential bias associated with this approach, i.e., year effect was also examined before final analysis (see below). Among environmental variables, edge1000 was highly correlated with pd1000 and crop500 ($r = 0.83$ and 0.68 , respectively, $P < 0.001$ in CWB data), which increased the variance inflation factor of these variables. Thus, edge1000 was not included in analyses.

I selected the maximum count of CWB between two transects and summed it over two visits at each site. The sum was used as the "abundance" of CWB. I $\log(x+1)$ -transformed the abundance of CWB to minimize potential bias caused by high abundance values and overdispersion that can affect type I error. I first tested whether survey year (2020 or 2021) had an effect on CWB abundance. I compared two generalized linear models (GLMs) with Gamma distribution: intercept-only model vs model with a year variable. The AICc (Akaike Information Criterion adjusted for small sample size) value of the intercept-only model was lower (< 2) than the other model, suggesting CWB abundance was not significantly affected by year-related variations. Thus, I constructed 4 GLMs without "year" variable: null model (intercept-only model), local model (3 local variables — Brassicaceae, cropdiv50, weed50), landscape model (3 landscape variables — crop500, weed500, pd1000), and full model (3 local and 3 landscape variables). These four models were compared based on their AICc values. If there were no competing models to a top model (model with a lowest AICc), that is, if other models had Δ AICc (AICc difference from top model) ≥ 4 , the top model was used to make inferences. Otherwise, all models with Δ AICc < 4 were considered as plausible models (Burnham &

Anderson, 2002) and their parameter estimates were model-averaged (“MuMIn” package; Bartoń, 2022).

I summed Apis individuals caught in a set of pant traps at each sample station, which was considered as a unit for the analysis. The sum represented the abundance of Apis per sample station. Environmental data for Apis included fine-scale data. However, flower covers at the fine scale was collected in 2021 only and thus I performed two separate analyses: the one with the data of two years (2020 and 2021) and the other with 2021 data only. I built a total of 8 GLMs with negative binomial distribution for each analysis using a combination of scales (TableS2). The survey year had a significant effect on Apis abundance and thus year was incorporated into all models that used both 2020 and 2021 data. The abundance data of Apis had zero values; while zero-inflated Poisson and negative binomial distributions can be used in this case, I chose negative binomial distribution because of overdispersion in the GLMs with zero-inflated Poisson. Model selection and averaging were carried out as for CWB.

Spatial dependence of bee and butterfly counts was also examined by conducting Moran’s I test on the residuals of the full models of CWB and Apis (“ape” package; Paradis & Schliep, 2019): $P > 0.1$ in all cases, indicating that spatial dependence was negligible. I checked other regression model assumptions such as homoscedasticity and overdispersion (“DHARMA” package; Hartig, 2022) and did not find any cases of significant violation. The values of variance inflation factor were < 2.5 , suggesting little issue of multicollinearity.

Results

A total of 1189 CWB individuals was observed across all sites. Mean abundance was 13.21 butterflies per site with a SD of 10.16, ranging from 1 to 39. The abundance of CWB was strongly associated with a combination of local and landscape variables: the full model showed the lowest AICc, and $\Delta AICc$ of other models was > 9 (Table 1).

Crop diversity had a positive effect on the abundance of CWB, whereas percent cover of farmland and weedy vegetation within the landscape affected negatively (Fig. 2 and Table S3). The fragmentation of farmlands, i.e., patch density, was not associated with CWB abundance given wide 95% confidence intervals across 0 and very low estimate (TableS3). Although percent cover of cruciferous crops tended to have a positive impact on CWB abundance, the effect was not as strong as other significant variables (TableS3).

For two winters, 265 Apis individuals were captured. While more individuals were caught in 2021 than 2020, Apis abundance varied by sample stations: 1.15 ± 2.11 (mean abundance \pm SD), ranging from 0 to 14 in 2020, and 1.82 ± 2.03 , ranging from 0 to 11 in 2021. Of 8 GLM models, 4 models were selected as plausible models in 2021 data analysis and 2 models in combined years data analysis (Table 2).

Effects of weedy vegetation on Apis abundance were consistent across two winters given that the 95% confidence intervals did not include 0 or slightly overlapped with 0. The effects also depended on scale: the abundance of Apis was affected by weedy vegetation cover negatively at the landscape scale but positively at the local scale (Fig. 3 and TableS3). In

combined years data analysis, increasing percent cover of farmland at the landscape scale was positively associated with increasing Apis abundance. Apis abundance was also low in a landscape with more fragmented farmlands. The percentage of flowering plants had a strong positive effect on Apis abundance in 2021: its coefficient was ≥ 1.5 higher than other variables associated with Apis, suggesting the amount of floral resource at the fine scale could be critical to Apis (Fig. 3).

Discussion

The results of this study reveal variation in environmental factors at relevant spatial scales associated with a host-specific pest, i.e., cabbage white butterflies, and a pollinator, i.e., Asian honey bees. Cabbage white butterfly was more related to crop cover at the landscape scale (negatively), whereas Asian honey bee was strongly related to flower cover at the fine scale and weedy vegetation at the local scale (positively). Both species also responded negatively to weedy vegetation cover at the landscape scale. These patterns can provide insight into how small urban farmlands can be managed to improve ecosystem service, i.e., pollination by bees, while minimizing ecosystem disservice, i.e., crop damage by cabbage white butterflies.

The resource concentration hypothesis predicts high abundance of host-specific herbivorous insects at their host plant rich patches, i.e., dense, large, or pure stands of host plants (Root, 1973). I expected a positive relationship between cabbage white butterfly abundance and percent cover of Brassicaceae, and I found a tendency for this relationship. However, landscape-scale crop cover was the environmental variable most strongly associated with the abundance of cabbage white butterfly. Certainly the presence or absence of Brassicaceae in surrounding farmlands could influence CWB distribution. If Brassicaceae cover is low in a landscape with otherwise high crop cover, we might expect a negative association. Although I lack data on Brassicaceae cover at the landscape scale, local Brassicaceae cover was not correlated with landscape crop cover. Also, almost all farmlands surveyed contained Brassicaceae: of 90 sites, 44 included $> 20\%$ Brassicaceae at the local scale and only 7 had $< 5\%$ Brassicaceae. Thus, the negative response of cabbage white butterfly is unlikely related to the amount of landscape Brassicaceae cover. Matteson and Langellotto (2012) found that cabbage white butterfly spends less time in urban gardens in a landscape with more green spaces, which could lead to low detection of cabbage butterfly in these gardens. I often observed more cabbage white butterflies and their mating, landing, and oviposition behaviors at farmlands isolated in a landscape dominated by built-up structure. Considering that crop and weedy vegetation covers are part of green spaces, the responses of cabbage white butterfly to both covers parallels the previous finding. Increasing crop covers in a landscape likely diffuses cabbage white butterflies, lowering abundance per farmland. It may also interrupt visual cues and consequently have a negative impact on their ability to search host and floral resources because cabbage white butterflies depend on vision to locate these resources (Hern et al., 1996).

The positive effect of crop diversity is somewhat unexpected. Polyculture stands often show lower density or higher mortality of pest insects compared to monoculture stands (Altieri et

al., 1978; Russell, 1989; Letourneau et al., 2011; Iverson et al., 2014; among others). The diversification of cruciferous crops through mixing or intercropping can effectively control pest insects due to higher abundance and richness of parasitoids/predators, visual camouflage effect, and masking effect of host plant odors, among other mechanisms (Finch & Collier 2000; Hooks & Johnson, 2003). The positive effect I observed may be affected by the season of survey. Although winter is relatively warm and dry in southern China, the abundance and richness of predator and parasitoid insects are lower than other seasons such as spring (unpublished data). Lowenstein and Minor (2018) reported different trends in abundance between herbivores including cabbage white butterflies and predator/parasitoid insects at urban gardens and farms in Chicago, USA: areas with high predators and parasitoids tended to have low herbivores. Compared to large-scale rural agricultural lands, urban farmlands have also more diverse crops despite their small size. Brassicaceae crops are cultivated at most of these farmlands in the study area. This creates a spatially and temporally heterogeneous environment due to variations in harvest time and growth stage between crops, even between Brassicaceae crops. These conditions may benefit cabbage white butterflies by providing host and floral resources, shelters, or mating chances for a longer period during the winter season. I also noticed seasonal variations in crop composition, although crop diversity at each study site did not change significantly between seasons. For example, in spring, the proportion of Brassicaceae crops declines but other crops (especially vine crops, e.g., bean and gourd family crops) increase and flowering weeds are abundant. Cabbage white butterflies may respond differently to crop diversity in spring, but the extent of possible seasonal variations in the responses of cabbage white butterflies requires further research.

Bee populations are strongly regulated by food resource availability (Roulston & Goodell, 2011). The establishment of semi-natural vegetation, especially planting flowering herbaceous plants at field edges (i.e., flower strips) is often recommended to promote crop pollination and yield by insect pollinators (Blaauw & Isaacs, 2014; Williams et al. 2015; Shutter et al., 2018; but see Nicholson et al., 2018). One recent meta-analysis shows that crop pollination decreases with increasing distance to floral resources, indicating the importance of floral resources to pollinators (Albrecht et al., 2020). Crop diversity and semi-natural vegetation at landscape scales also increase bumble bee density in wheat-dominant agricultural areas (Raderschall et al., 2021). The result of the current study, i.e., the strong effect of percent cover of flowering plants on bee abundance, partly supports the general trend. While I do not have data on the diversity and abundance of flowering plants at local and landscape scales, the result highlights the benefit of even small amount of flowering crop and weed to honey bees in urban environments, especially during the winter in which natural floral resources may be scarce. This could also explain the different effects of weedy vegetation cover at local and landscape scales. The subtropical climate in southern China enables farmers to cultivate vegetable crops year-round. While some weedy vegetation showed withering in winter, the condition was less severe at urban farmlands than some vacant lots (e.g., construction sites and abandoned lands), which is likely affected by

differences in water availability. There were also more weed plants blooming at urban farmlands based on my observation.

One potential concern is the quality of floral resources in these urban farmlands because most common weed plants flowering in the study area are non-native plants, particularly *Bidens* species. There are few studies comparing nutritional values of nectars to bees between crops and weed plants in this region. However, honey bees are generalists, visiting a wide range of flowering plants, and require different diets to maintain colony health (Requier et al., 2015). During times of food shortage, weed plants and non-native plants can be an important component of the diets of *A. mellifera* and *A. cerana*, respectively (Requier et al., 2015; Koyama et al., 2018). *Apis mellifera* also forages on flowers of *Bidens* species (Kajobe, 2006). Floral resource use and preference by bees are significantly associated with dominant plants and affected by season as floral resource availability changes throughout the year (Lowe et al., 2021). Most flowering plants at sample sites in winter are weed plants and several crops that belong to the genera *Brassica*, *Chrysanthemum*, and *Allium*. Weed plants can be one of major floral resources to Asian honey bees in winter regardless of their nutritional values. However, with seasonal variations in the diversity and abundance of floral resources, Asian honey bees may prefer certain crops or weed plants in other seasons such as spring, when trees and shrubs as well as more crops bloom. The degree of effects of flower strips on local bee abundance and diversity can also depend on the interaction between characteristics of flower strips and floral resources available in surrounding landscape (Scheper et al., 2015). Similarly, it is possible that the strong response of Asian honey bees to fine-scale flower cover could be influenced by the total amount of floral resources within farmland and landscape, which will vary by season.

Conclusions

Overall, the findings of current study provide a scientific basis for urban planners, policy makers, and farmers to consider in the management of urban farmlands across scales to enhance the sustainability of these farmlands. At the farmland (local) scale, cultivating too diverse crops would not be recommended as it can increase the risk of crop damage by cabbage white butterfly. However, the risk can be alleviated at farmlands embedded in a landscape with more overall crop cover, which may partly benefit Asian honey bees as well. Retaining non-crop area such as weedy vegetation patches or small portion of flowering plants including crops within a farmland can be an effective practice to maintain Asian honey bee population and promote pollination. It also benefits birds as local weedy vegetation has a positive effect on winter bird diversity at urban farmlands (Lee et al., 2022). With growing urban expansion, farmlands in China have been significantly converted into built-up structure, leading to a decline in net primary productivity of cropland (He et al., 2017) despite farmland protection policy restrictions on development. Farmlands in a city are not exceptional. In major cities in China, new development largely spreads from the edge of city or occurs at old villages within a city (Tu et al., 2021), lowering crop cover in surrounding landscapes and isolating farmlands (Liang et al., 2015). Policy makers and urban planners must consider preserving farmlands, especially at the

city fringe or suburb for permanent agricultural activities. Priority should be given to farmlands embedded in landscapes with relatively high crop covers. It is also critical that city government monitors potential environmental problems associated with urban farmlands. For example, most of these farmlands, especially ones next to main districts where development and population are concentrated, use waste water. The average contents of heavy metals in agricultural soil in Guangzhou and Foshan cities are higher than background values of Guangdong province (Xiao et al., 2020). While the levels of metals are still lower than national standard values, the ecological risk of heavy metal pollution in these cities is not negligible. Combined with further research on seasonal variations in the patterns found in current study, the assessment of environmental quality of urban farmlands can contribute to improve the sustainability of urban agroecosystem.

Acknowledgements

I would like to thank Y. Zhang and D. Cheng for their assistance in collecting local environmental data as well as honey bee data, D. Chen for handling logistic issues, and J. Rotenberry for edits and comments on the draft of this manuscript.

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Table 1(on next page)

Model selection results of cabbage white butterfly based on Akaike information criterion value adjusted for small sample size (AICc).

Note a big difference in AICc values (Δ AICc) between top model and other models. Null model is an intercept-only model, whereas full model includes both local- and landscape-scale variables. Abbreviation: DF, degree of freedom; LogLik, log-likelihood.

1

Model	DF	LogLik	AICc	Δ AICc	AICc weight
Full	8	-81.506	180.8	0.00	0.988
Landscape	5	-89.493	189.7	8.91	0.011
Local	5	-95.622	202.0	21.17	0.000
Null	2	-113.276	230.7	49.90	0.000

2

3

Table 2 (on next page)

Summary of model comparisons of Asian honey bee.

Null model is an intercept-only model, whereas full model includes variables of all three scales. A combination of two spatial scales is denoted with “+”: for example, Fine+Local model contains variables at fine and local scales. Note that fine-scale flower cover was considered in 2021 only. Models in “Both” year are constructed using two years of data, i.e., 2020 and 2021. Abbreviation: DF, degree of freedom; LogLik, log-likelihood; Δ AICc, AICc difference from top model.

1

Year	Model	DF	LogLik	AICc	Δ AICc	AICc weight
2021	Full	9	-163.529	347.1	0.00	0.444
	Fine+Local	6	-167.580	348.1	0.99	0.270
	Fine+Local	4	-170.216	348.9	1.78	0.182
	Fine+Landscape	7	-167.387	350.0	2.92	0.103
	Local+Landscape	7	-172.985	361.2	14.12	0.000
	Local	4	-176.948	362.3	15.24	0.000
	Landscape	5	-176.131	362.9	15.83	0.000
	Null	2	-179.720	363.6	16.49	0.000
Both	Local+Landscape	8	-280.330	577.5	0.00	0.582
	Full	9	-279.815	578.7	1.20	0.320
	Landscape	6	-284.734	582.0	4.44	0.063
	Fine+Landscape	7	-284.620	583.9	6.38	0.024
	Local	5	-288.486	587.3	9.79	0.004
	Null	3	-291.062	588.3	10.73	0.003
	Fine+Local	6	-288.123	588.8	11.21	0.002
	Fine	4	-291.054	590.3	12.81	0.001

2

3

Figure 1

Study sites established in the city of Guangzhou, Guangdong Province, China.

All sites are located in the Indo-Burma Biodiversity Hotspot (Indo-Burma) areas in Guangzhou. Number of sites used for each taxon and year are variable.

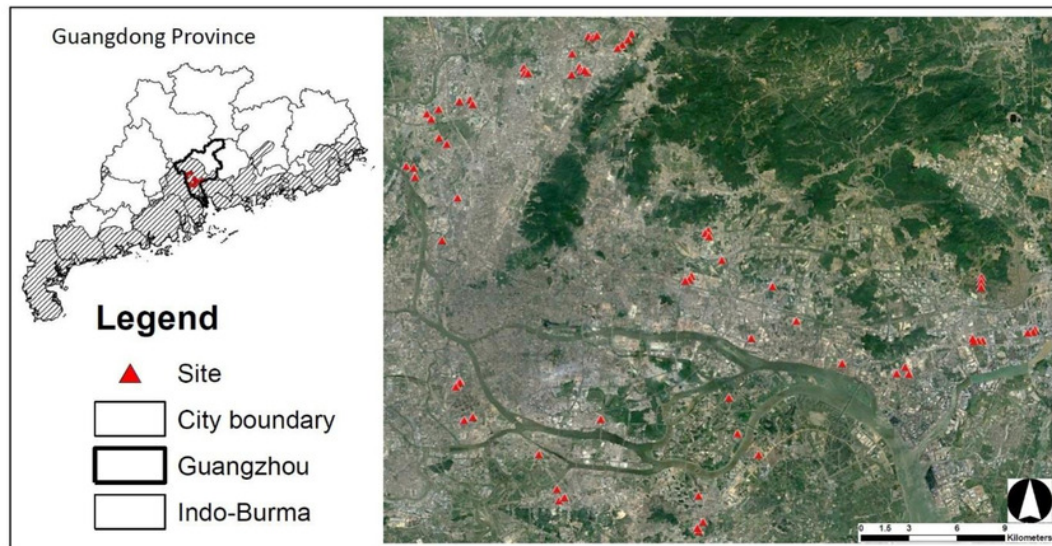


Figure 2

Significant relationship between the abundance of cabbage white butterfly and environmental variables: local crop diversity (cropdiv50, A), landscape crop cover (crop500, B), and landscape weedy vegetation cover (weed500, C).

Y-axis represents predicted abundance that is $\log(x+1)$ transformed. X-axis shows standardized values of each environmental variable.

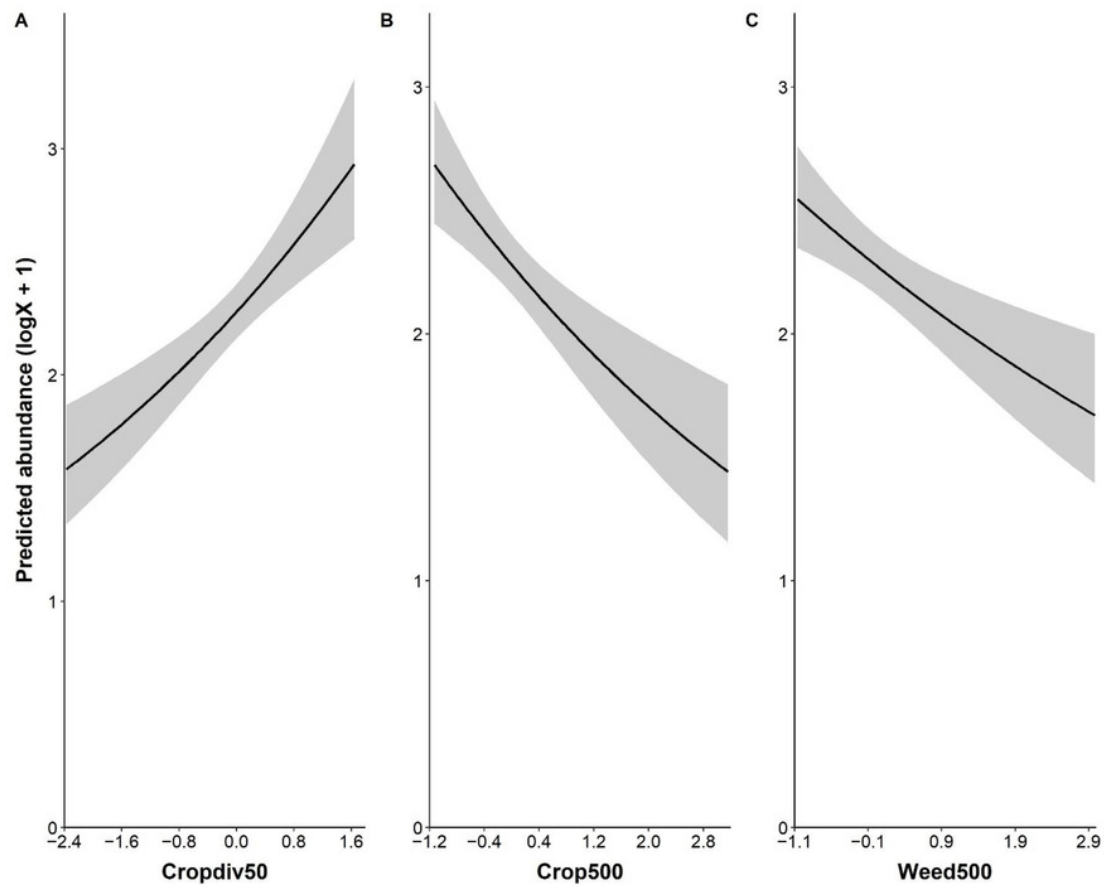


Figure 3

Model averaged parameter estimates and their 95% confidence intervals (error bars on the graph) from the analysis of Asian honey bee using 2021 data (A) and two years of data (B).

X-axis shows all environmental variables at each of three spatial scales: fine scale variables, flower (flower cover) and cropdiv5 (crop diversity); local scale variables, cropdiv50 (crop diversity) and weed50 (weedy vegetation cover); landscape scale variables, crop500 (crop cover), weed500 (weedy vegetation cover), and pd1000 (patch density).

