

# UC Riverside

## UC Riverside Previously Published Works

### Title

Effects of global change on insect pollinators: multiple drivers lead to novel communities

### Permalink

<https://escholarship.org/uc/item/3412b1h0>

### Author

Rafferty, Nicole E

### Publication Date

2017-10-01

### DOI

10.1016/j.cois.2017.06.009

### Copyright Information

This work is made available under the terms of a Creative Commons Attribution-NonCommercial License, available at <https://creativecommons.org/licenses/by-nc/4.0/>

Peer reviewed

Dear author,

Please note that changes made in the online proofing system will be added to the article before publication but are not reflected in this PDF.

We also ask that this file not be used for submitting corrections.



ELSEVIER

ScienceDirect

Current Opinion in  
Insect Science

# Effects of global change on insect pollinators: multiple drivers lead to novel communities

Nicole E Rafferty<sup>1,2</sup>

Global change drivers, in particular climate change, exotic species introduction, and habitat alteration, affect insect pollinators in numerous ways. In response, insect pollinators show shifts in range and phenology, interactions with plants and other taxa are altered, and in some cases pollination services have diminished. Recent studies show some pollinators are tracking climate change by moving latitudinally and elevationally, while others are not. Shifts in insect pollinator phenology generally keep pace with advances in flowering, although there are exceptions. Recent data demonstrate competition between exotic and native bees, along with rapid positive effects of exotic plant removal on pollinator richness. Genetic analyses tie bee fitness to habitat quality. Across drivers, novel communities are a common outcome that deserves more study.

## Addresses

<sup>1</sup> Department of Biology, University of California, 900 University Ave., Riverside, CA 92521, USA

<sup>2</sup> Rocky Mountain Biological Lab, P.O. Box 519, Crested Butte, CO 81224, USA

Corresponding author: Rafferty, Nicole E ([nicole.rafferty@ucr.edu](mailto:nicole.rafferty@ucr.edu))

**Current Opinion in Insect Science** 2017, **23**:xx–yy

This review comes from a themed issue on **Global change biology**

Edited by **Brandon Barton** and **Jason Harmon**

<http://dx.doi.org/10.1016/j.cois.2017.06.009>

2214-5745/© 2017 Elsevier Inc. All rights reserved.

## Introduction

Global change is affecting insect pollinators in profound ways. Climate change, exotic species introduction, and habitat loss are affecting all major aspects of the biology of insects that pollinate plants in both natural and agricultural communities, altering their distribution, phenology, abundance, physiology, and morphology [1–5]. The consequences of these effects are complex, perturbing plant–pollinator interactions in subtle but important ways and in some cases resulting in local extinction [2]. Despite the complexity, understanding these consequences is critical: just as the vast majority of flowering plants depend on insects for pollination [6], we rely in large part on insects to pollinate our crops, a valuable ecosystem service [7].

Among the many insect taxa that serve as pollinators, bees, flies, butterflies, and moths have received the most study in the context of global change. Within these taxa, bees are key pollinators of both crop plants and wild plants [8], and studies on bees have dominated the literature on plant–pollinator interactions under global change. Because bees rely heavily on floral resources both for their own sustenance and to provision their offspring, their fitness is strongly determined not only by the direct effects of global change but also by the influence of global change drivers on flowering plants.

Here, I consider the effects of several global change drivers on insect pollinators, with an emphasis on what we know about the effects on native bees. First, I discuss how climate change is affecting insect pollinators, as this is a topic of active research that illustrates a suite of responses. Second, I review the effects of exotic species, both insect and plant taxa, on insect pollinators. Third, I consider another global change factor, habitat alteration and loss, and its effects on insect pollinators. Throughout, I consider both direct effects on pollinators and effects that are mediated via plants and other interspecific interactions. Given biotic pollination is by definition a multitrophic interaction, greater consideration of how global change alters species interactions is needed to improve conservation and management of pollination services.

## Effects of climate change

The responses of insect pollinators to climate change have been relatively well-studied, although much remains to be resolved. For the most part, experimental studies of climate change factors on insect pollinators have focused on temperature [9–12], an important determinant of developmental rate [13]. Manipulations of other factors, such as carbon dioxide [14] or precipitation [15], have been applied to plants with subsequent measures of pollinator responses to altered floral traits. Complementing experimental approaches are long-term data, historical observations, and museum specimen records that can be correlated with ambient temperatures and other climate variables to describe insect responses [1,16].

Among the most striking consequences of climate change have been shifts in the spatial distributions of insect pollinators. Given the rapid life cycles and high mobility of most insect pollinators, are they able to keep pace with anthropogenic climate change by tracking environmental

## 2 Global change biology

90 conditions over space? Evidence is mixed. On the one  
 91 hand, Kerr *et al.* [4\*\*] discovered bumble bees (*Bombus*  
 92 spp.) across two continents have not tracked warming  
 93 temperatures, as evidenced by a failure to expand their  
 94 northern latitudinal range limits. On the other hand,  
 95 several studies have shown that bumble bees have moved  
 96 upward in elevation in montane ecosystems [4\*\*,17,18],  
 97 and some butterflies have shifted up in altitude [19]. Both  
 98 a nymphalid butterfly (*Polygonia c-album*) and a lycaenid  
 99 butterfly (*Aricia agestis*) in Britain have greatly expanded  
 100 their ranges northward in association with warming  
 101 [20,21]. A key question that has been not been considered  
 102 for most taxa is how these spatial shifts affect interactions  
 103 with floral resources and thereby influence both pollinator  
 104 fitness and patterns of pollen flow and reproductive  
 105 output of plants. Differential shifts among taxa will  
 106 almost certainly translate into modified communities,  
 107 especially as perennial plants are likely to lag behind  
 108 their pollinators. In addition, it remains largely unknown  
 109 whether traits or phylogenetic relationships can explain  
 110 variable spatial responses among taxa (but see [4\*\*,22]).  
 111 To understand constraints on the distributions of insect  
 112 pollinator populations and predict how distributions will  
 113 be affected by climate change directly and via effects on  
 114 host plants and other species with which pollinators  
 115 interact, species distribution models can be a useful tool  
 116 [23,24].

117 Shifts in the phenologies of insect pollinators are another  
 118 conspicuous signal of climate change. Multiple species of  
 119 bees have significantly advanced their phenologies [1], as  
 120 have many butterflies and moths [25,26]. Among lepi-  
 121 dopterans, variable responses can be partially explained  
 122 by traits such as diet breadth [26]. In contrast to spatial  
 123 shifts, the consequences of climate change-induced tem-  
 124 poral shifts for plant–pollinator interactions have received  
 125 much attention. Community-level analyses indicate bees  
 126 and the plants they pollinate are advancing at similar rates  
 127 [1], whereas butterflies and their nectar sources show  
 128 different sensitivities to temperature [27\*]. In general,  
 129 experimental studies suggest phenological mismatches  
 130 are unlikely to lead to complete decoupling of interac-  
 131 tions among insect pollinators and plants [28,29]. In part  
 132 this outcome is not surprising: plant–pollinator interac-  
 133 tions tend to be generalized [30] and nested, with  
 134 specialists interacting with generalists [31], and high  
 135 rates of interaction turnover [32]. However, there are  
 136 examples of specialized plant–bee interactions that are  
 137 likely becoming disrupted as phenologies shift [33,34].  
 138 Even subtle phenological mismatches are likely to have  
 139 consequences for interaction strengths, fitness, and the  
 140 evolution of life histories [35]. Whereas the conse-  
 141 quences of mismatches for plants have been commonly  
 142 measured in terms of seed production [29,36], the conse-  
 143 quences for pollinators have gone unquantified [37].  
 144 Also in contrast to the situation for insect pollinator  
 145 phenology, where few studies have linked responses

146 to traits or phylogenies, flowering phenology responses  
 147 to climate change have been associated with traits such  
 148 as flowering season, life history, and pollination mode  
 149 [38,39] and exhibit phylogenetic signal across continents  
 150 [40]. Together, these gaps in understanding point to a  
 151 need for more studies at the community level; a com-  
 152 munity approach should simultaneously create opportu-  
 153 nities for trait-based analyses and enable the conse-  
 154 quences of phenological mismatches from the  
 155 pollinator perspective to be quantified.

156 Other aspects of climate change that have been demon-  
 157 strated to affect insect pollinators via flowering plants  
 158 include elevated carbon dioxide and decreased precipi-  
 159 tation. Plants grown under elevated carbon dioxide can  
 160 have altered floral traits, such as nectar composition [14]  
 161 and pollen protein concentration [41]. In turn, these  
 162 altered traits can influence the fitness of insect pollinators;  
 163 Hoover *et al.* [14] found that *Bombus terrestris* workers  
 164 exhibited reduced longevity when fed synthetic nectar  
 165 mimicking that of flowers produced under elevated car-  
 166 bon dioxide, and Ziska *et al.* [41] posit that reduced  
 167 protein in goldenrod pollen could negatively affect bees.  
 168 Experimental drought had variable effects on floral vola-  
 169 tiles but consistently reduced flower size and floral display  
 170 across four species, resulting in different communities of  
 171 bees, flies, and butterflies visiting the flowers in the  
 172 drought treatment [15]. In general, a tight link between  
 173 the direct effects of climate change on floral resources and  
 174 the consequent effects on insect pollinators has yet to be  
 175 made. In part, this is because it is difficult to isolate the  
 176 effects of complex floral responses on mobile insects,  
 177 particularly in the field and at the population and com-  
 178 munity levels. As molecular genetic techniques and tech-  
 179 nologies that allow automated identification of individual  
 180 bees, for example as they pass over radio frequency  
 181 identification readers, are refined, larger-scale field-based  
 182 studies of pollinator fitness and foraging responses should  
 183 become more feasible.

### Effects of exotic species

184 Human-aided transport and introduction of exotic species  
 185 is a major driver of global change, reshaping fundamental  
 186 ecological relationships [42]. Focusing in on exotic insect  
 187 pollinators, we know the most about the impacts of non-  
 188 native bees on native bees [43]. Non-native bees include  
 189 long-established domesticated honey bees (*Apis melli-  
 190 fera*), more recently-introduced commercial pollinators,  
 191 such as *Bombus terrestris* [44], and accidental introductions  
 192 of species such as *Hylaeus communis* [45]. Alien pollinators  
 193 can compete with native pollinators for resources, poten-  
 194 tially reducing their fitness, altering patterns of pollen  
 195 flow, and ultimately changing community structure to the  
 196 disruption of ecosystem services [46,47]. Not surprisingly,  
 197 the best-studied interactions between exotic and native  
 198 bees involve honey bees. Building on prior experimental  
 199 work that demonstrated competition for floral resources

200 between honey bees and a native bumble bee (*B. occi-*  
201 *dentalis*; [48]), Thomson [49\*\*] used a 15-year-long data  
202 set to show a negative relationship between feral *A.*  
203 *mellifera* densities and *Bombus* spp. densities. Similarly,  
204 after honey bees invaded a tropical reserve, solitary bees  
205 were observed to visit different plant species because of  
206 competition, but declines in the native bees were not  
207 detected [50]. Thus, the effects of exotic insect pollina-  
208 tors on native pollinators likely depend on factors that  
209 modify the strength of competition, such as niche overlap  
210 and flexibility, as well as interacting effects of other  
211 stressors, such as drought, that modulate floral resource  
212 availability.

211 Turning briefly to non-native plants, several studies  
212 have investigated how exotic plants influence plant–  
213 pollinator interactions [46,51–53]. Recently, a large  
214 experiment by Kaiser-Bunbury *et al.* [54\*\*] showed  
215 exotic plant removal resulted in about 20% more polli-  
216 nator species in restored sites, with more generalized  
217 plant–pollinator networks and higher fruit set of com-  
218 mon species. These results suggest removal of non-  
219 native species can rapidly enhance pollinator richness  
220 but may, as the authors note, hinge on nearby popula-  
221 tions of pollinators to colonize restored sites [54\*\*]. More  
222 broadly, no real consensus on the effects of exotic plants  
223 on insect pollinators has emerged, with both positive and  
224 negative effects reported [46,51]. Moving forward,  
225 greater integration of the study of exotic species with  
226 the study of phenological and range shifts, which can  
227 similarly modify interaction strengths and create novel  
228 communities, would be productive.

### 229 Effects of habitat alteration and loss

230 Habitat alteration and loss is widely recognized as a  
231 contributor to declines of insect pollinators [55].  
232 Changes in land use are associated with changes in  
233 pollinator community composition and richness; in par-  
234 ticular, conversion to arable land is associated with  
235 declines in bee and wasp species richness over 80 years  
236 in Britain [56]. Agricultural intensification carries its own  
237 suite of effects on insect pollinators, including the direct  
238 effects of pesticides such as neonicotinoids, which can  
239 have multiple debilitating effects on bees [57–59], weak-  
240 ening pollination services [60]. Using genetic analyses, a  
241 recent study by Carvell *et al.* [61\*] showed that lineage  
242 survival of three bumble bee species increased as a  
243 function of nearby high-quality foraging habitat, quanti-  
244 fied as semi-natural vegetation, spring floral resources for  
245 queens, and overall flower cover in spring and summer.  
246 Bumble bee nesting density also can be negatively  
247 related to the percent of paved surface and positively  
248 related to the amount of natural oak woodland-chaparral  
249 habitat [62].

250 Some traits serve as predictors of the severity of effects of  
251 habitat alteration and loss on insect pollinators. Generally,

252 specialized pollinators are more sensitive to land use  
253 impacts [63,64]. Within bees, a global analysis indicated  
254 stronger negative effects of overall agricultural intensifi-  
255 cation and isolation from natural habitat for species that  
256 nest above ground, whereas species that nest below-  
257 ground were adversely affected by land tilling [65].  
258 The abundance of social bees was also more negatively  
259 affected by isolation than was the case for solitary bees  
260 [65]. Some pollinators may be able to adjust their foraging  
261 distances in response to landscape-scale variables, as seen  
262 with bumble bees capable of foraging farther to find  
263 patches of greater floral diversity in landscapes that are  
264 relatively homogeneous [62]. Altogether, multiple studies  
265 indicate that ecological intensification practices, such as  
266 increasing floral resource availability and diversity across  
267 landscapes, have positive effects on insect pollinator  
268 persistence in the face of habitat alteration [66]. Never-  
269 theless, with changing land use, pollinator behavior and  
270 species composition are likely to change, modifying inter-  
271 actions and pollination services.

### 272 Conclusions

273 As we become increasingly aware that species interactions  
274 shape species distributions in time and space and modu-  
275 late the direct effects of global change, considering insect  
276 pollinators in a community context should be a priority.  
277 For example, Forrest and Chisholm [67\*] demonstrated  
278 that warmer temperatures led simultaneously to higher  
279 rates of activity and nest provisioning by a solitary bee  
280 (*Osmia iridis*) and to increased rates of brood parasitism by  
281 a wasp (*Sapyga* sp.). Thus, positive effects of warming are  
282 likely to be negated by altered interaction frequencies  
283 with a natural enemy [67\*], a result that would not be  
284 predicted in isolation of community context. Community-  
285 level analyses also detect broader trends before pairwise  
286 interactions are disrupted or individual species decline.  
287 For example, a study of phenological overlap in  
288 Greenland over 18 years points to disrupted plant–polli-  
289 nator interactions as the flowering season shrinks, poten-  
290 tially leaving pollinators without floral resources late in  
291 the season [68].

292 Much progress has been made in understanding the  
293 effects of individual global change drivers on insect  
294 pollinators. Moving forward, further progress in under-  
295 standing and mitigating anthropogenic disturbances  
296 could be made by searching for common outcomes across  
297 drivers. All three of the global change drivers highlighted  
298 here are likely to result in novel interactions and commu-  
299 nities. Climate change, for example, alters overlap among  
300 species via spatial and temporal shifts, among other  
301 mechanisms. Introduced exotic species interact with  
302 native species in novel ways. And habitat alteration  
303 and loss can result in novel species composition and cause  
304 species to modify behavior, altering interactions. By tying  
305 these common outcomes to resulting eco-evolutionary  
306 dynamics, we can begin to anticipate how global change

## 4 Global change biology

307 will reshape insect pollinator communities and pollina-  
308 tion services.

309 **Conflict of interest statement**

310 Nothing declared.

311 **Acknowledgements**

312 I thank Brandon Barton and Jason Harmon for organizing this issue. I am  
313 grateful to Paul Nabity and an anonymous reviewer for comments that  
314 improved this manuscript. This research did not receive any specific grant  
315 from funding agencies in the public, commercial, or not-for-profit sectors.

316 **References and recommended reading**

317 Papers of particular interest, published within the period of review,  
318 have been highlighted as:

- of special interest
- of outstanding interest

- 319 1. Bartomeus I, Ascher JS, Wagner D, Danforth BN, Colla S,  
320 Kornbluth S, Winfree R: **Climate-associated phenological**  
321 **advances in bee pollinators and bee-pollinated plants.** *Proc*  
*Natl Acad Sci U S A* 2011, **108**:20645–20649.
- 322 2. Burkle LA, Martin JC, Knight TM: **Plant–pollinator interactions**  
323 **over 120 years: loss of species, co-occurrence, and function.**  
324 *Science* 2013, **339**:1611–1615.
- 325 3. Scaven VL, Rafferty NE: **Physiological effects of climate**  
326 **warming on flowering plants and insect pollinators and**  
327 **potential consequences for their interactions.** *Curr Zool* 2013,  
328 **59**:418–426.
- 329 4. Kerr JT, Pindar A, Galpern P, Packer L, Potts SG, Colla S: **Climate**  
330 **change impacts on bumblebees converge across continents.**  
331 *Science* 2015, **349**:177–180.  
332 Using a spatially and temporally extensive dataset, the authors show that  
333 bumble bees across two continents have experienced range contractions  
334 at their southern limits and have failed to track climatic changes by  
335 expanding their northern range limits, although species have shifted  
336 up in elevation in some regions. Importantly, this study shows bumble  
337 bees are showing range responses opposite to that of most species, and  
338 these responses cannot be explained by land or pesticide use.
- 339 5. Miller-Struttman NE, Geib JC, Franklin JD, Kevan PG, Holdo RM,  
340 Ebert-May D, Lynn AM, Kettenbach JA, Hedrick E, Galen C:  
**Functional mismatch in a bumble bee pollination mutualism**  
**under climate change.** *Science* 2015, **349**:1541–1544.
- 341 6. Ollerton J, Winfree R, Tarrant S: **How many flowering plants are**  
342 **pollinated by animals?** *Oikos* 2011, **120**:321–326.
- 343 7. Klein AM, Vaissiere BE, Cane JH, Steffan-Dewenter I,  
344 Cunningham SA, Kremen C, Tscharntke T: **Importance of**  
345 **pollinators in changing landscapes for world crops.** *Proc R Soc*  
*B* 2007, **274**:303–313.
- 346 8. Potts SG, Biesmeijer JC, Kremen C, Neumann P, Schweiger O,  
347 Kunin WE: **Global pollinator declines: trends, impacts and**  
**drivers.** *Trends Ecol Evol* 2010, **25**:345–353.
- 348 9. Williams CM, Hellmann J, Sinclair BJ: **Lepidopteran species**  
349 **differ in susceptibility to winter warming.** *Clim Res* 2012,  
350 **53**:119–130.
- 351 10. Frund J, Zieger SL, Tscharntke T: **Response diversity of wild**  
352 **bees to overwintering temperatures.** *Oecologia* 2013,  
353 **173**:1639–1648.
- 354 11. Jevanandam N, Goh AGR, Corlett RT: **Climate warming and the**  
355 **potential extinction of fig wasps, the obligate pollinators of**  
356 **figs.** *Biol Lett* 2013, **9**:20130041.
- 357 12. Bennett MM, Cook KM, Rinehart JP, Yocum GD, Kemp WP,  
358 Greenlee KJ: **Exposure to suboptimal temperatures during**  
359 **metamorphosis reveals a critical developmental window in the**  
360 **solitary bee, *Megachile rotundata*.** *Physiol Biochem Zool* 2015,  
**88**:508–520.

13. Kingsolver JG, Huey RB: **Size, temperature, and fitness: three**  
361 **rules.** *Evol Ecol Res* 2008, **10**:251–268. 362
14. Hoover SER, Ladley JJ, Shchepetkina AA, Tisch M, Gieseg SP,  
363 Tylianakis JM: **Warming, CO<sub>2</sub>, and nitrogen deposition**  
**interactively affect a plant–pollinator mutualism.** *Ecol Lett*  
2012, **15**:227–234. 364 365
15. Burkle LA, Runyon JB: **Drought and leaf herbivory influence**  
366 **floral volatiles and pollinator attraction.** *Glob Change Biol* 2016,  
22:1644–1654. 367 368
16. Ovaskainen O, Skorokhodova S, Yakovleva M, Sukhov A,  
369 Kutenkov A, Kutenkova N, Shcherbakov A, Meyke E, del Mar  
Delgado M: **Community-level phenological response to**  
**climate change.** *Proc Natl Acad Sci U S A* 2013, **110**:13434–  
13439. 370 371
17. Ploquin EF, Herrera JM, Obeso JR: **Bumblebee community**  
372 **homogenization after uphill shifts in montane areas of**  
**northern Spain.** *Oecologia* 2013, **173**:1649–1660. 373 374
18. Pyke GH, Thomson JD, Inouye DW, Miller TJ: **Effects of climate**  
375 **change on phenologies and distributions of bumble bees and**  
**the plants they visit.** *Ecosphere* 2016, **7**:e01267. 376 377
19. Konvicka M, Maradova M, Benes J: **Uphill shifts in distribution of**  
378 **butterflies in the Czech Republic: effects of changing climate**  
**detected on a regional scale.** *Glob Ecol Biogeogr* 2003, **12**:403–  
410. 379 380 381
20. Braschler B, Hill JK: **Role of larval host plants in the climate-**  
382 **driven range expansion of the butterfly *Polygonia c-album*.** *J*  
*Anim Ecol* 2007, **76**:415–423. 383 384
21. Pateman RM, Hill JK, Roy DB, Fox R, Thomas CD: **Temperature-**  
385 **dependent alterations in host use drive rapid range expansion**  
**in a butterfly.** *Science* 2012, **336**:1028–1030. 386 387
22. Pöyry J, Luoto M, Heikkinen RK, Kuussaari M, Saarinen K:  
388 **Species traits explain recent range shifts of Finnish butterflies.**  
389 *Glob Change Biol* 2009, **15**:732–743. 390
23. Van der Putten WH, Macel M, Visser ME: **Predicting species**  
391 **distribution and abundance responses to climate change: why**  
**it is essential to include biotic interactions across trophic**  
**levels.** *Philos Trans R Soc Lond B Biol Sci* 2010, **365**:2025–2034. 392 393 394
24. Giannini TC, Chapman DS, Saraiva AM, Alves-dos-Santos I,  
395 Biesmeijer JC: **Improving species distribution models using**  
**biotic interactions: a case study of parasites, pollinators and**  
**plants.** *Ecography* 2013, **36**:649–656. 396 397
25. Forister ML, Shapiro AM: **Climatic trends and advancing spring**  
398 **flight of butterflies in lowland California.** *Glob Change Biol*  
2003, **9**:1130–1135. 399 400
26. Altermatt F: **Tell me what you eat and I'll tell you when you fly:**  
401 **diet can predict phenological changes in response to climate**  
**change.** *Ecol Lett* 2010, **13**:1475–1484. 402 403
27. Kharouba HM, Vellend M: **Flowering time of butterfly nectar**  
404 **food plants is more sensitive to temperature than the timing of**  
**butterfly adult flight.** *J Anim Ecol* 2015, **84**:1311–1321. 405 406  
This study uses butterfly collection and herbarium records across British  
407 Columbia, Canada, to demonstrate that plant phenology is more sensitive  
408 than butterfly phenology to changing temperatures. The authors also look  
409 at interacting butterfly–plant species pairs to explore how phenological  
410 overlap may change with temperature. This work represents an important  
411 step toward understanding whether climate change will differentially  
412 affect insect pollinator and flowering plant phenologies, with insight  
413 gained from analyses at both the community-wide and species-pair  
414 levels.
28. Rafferty NE, Ives AR: **Effects of experimental shifts in flowering**  
415 **phenology on plant–pollinator interactions.** *Ecol Lett* 2011,  
14:69–74. 416 417
29. Gezon ZJ, Inouye DW, Irwin RE: **Phenological change in a spring**  
418 **ephemeral: implications for pollination and plant**  
**reproduction.** *Glob Change Biol* 2016, **22**:1779–1793. 419 420
30. Waser NM, Chittka L, Price MV, Williams NM, Ollerton J:  
421 **Generalization in pollination systems, and why it matters.**  
422 *Ecology* 1996, **77**:1043–1060. 423

- 424 31. Bascombe J, Jordano P, Melián CJ, Olesen JM: **The nested**  
425 **assembly of plant-animal mutualistic networks.** *Proc Natl Acad Sci U S A* 2003, **100**:9383-9387. 486
- 427 32. CaraDonna PJ, Petry WK, Brennan RM, Cunningham JL,  
428 Bronstein JL, Waser NM, Sanders NJ: **Interaction rewiring and**  
429 **the rapid turnover of plant-pollinator networks.** *Ecol Lett* 2017, **20**:385-394. 489
- 430 33. Kudo G, Ida TY: **Early onset of spring increases the**  
431 **phenological mismatch between plants and pollinators.**  
432 *Ecology* 2013, **94**:2311-2320. 490
- 433 34. Robbirt KM, Roberts DL, Hutchings MJ, Davy AJ: **Potential**  
434 **disruption of pollination in a sexually deceptive orchid by**  
435 **climatic change.** *Curr Biol* 2014, **24**:1-5. 491
- 436 35. Gienapp P, Reed TE, Visser ME: **Why climate change will**  
437 **invariably alter selection pressures on phenology.** *Proc Biol Sci*  
438 2014, **281**:20141611. 492
- 439 36. Rafferty NE, Ives AR: **Pollinator effectiveness varies with**  
440 **experimental shifts in flowering time.** *Ecology* 2012, **93**:803-  
441 814. 493
- 442 37. Forrest JRK: **Plant-pollinator interactions and phenological**  
443 **change: what can we learn about climate impacts from**  
444 **experiments and observations?** *Oikos* 2015, **124**:4-13. 494
- 445 38. Fitter AH, Fitter R: **Rapid changes in flowering time in British**  
446 **plants.** *Science* 2002, **296**:1689-1691. 495
- 447 39. Calinger KM, Queenborough S, Curtis PS: **Herbarium specimens**  
448 **reveal the footprint of climate change on flowering trends**  
449 **across north-central North America.** *Ecol Lett* 2013, **16**:1037-  
450 1044. 496
- 451 40. Rafferty NE, Nability PD: **A global test for phylogenetic signal in**  
452 **shifts in flowering time under climate change.** *J Ecol* 2016,  
453 **105**:627-633. 497
- 454 41. Ziska LH, Pettis JS, Edwards J, Hancock JE, Tomecek MB,  
455 Clark A, Dukas JS, Loladze I, Polley HW: **Rising atmospheric CO<sub>2</sub>**  
456 **is reducing the protein concentration of a floral pollen source**  
457 **essential for North American bees.** *Proc R Soc B* 2016,  
458 **283**:20160414. 498
- 459 42. Helmus MR, Mahler DL, Losos JB: **Island biogeography of the**  
460 **Anthropocene.** *Nature* 2014, **513**:543-546. 499
- 461 43. Russo L: **Positive and negative impacts of non-native bee**  
462 **species around the world.** *Insects* 2016, **7**:69. 500
- 463 44. Dafni A, Kevan P, Gross CL, Goka K: ***Bombus terrestris*,**  
464 **pollinator, invasive and pest: an assessment of problems**  
465 **associated with its widespread introductions for commercial**  
466 **purposes.** *Appl Entomol Zool* 2010, **45**:101-113. 501
- 467 45. Martins KT, Normandin É, Ascher JS: ***Hylaeus communis***  
468 **(Hymenoptera: Colletidae), a new exotic bee for North America**  
469 **with generalist foraging and habitat preferences.** *Can Entomol*  
470 2017 <http://dx.doi.org/10.4039/tce.2016.62>. 502
- 471 46. Stout JC, Morales CL: **Ecological impacts of invasive alien**  
472 **species on bees.** *Apidologie* 2009, **40**:388-409. 503
- 473 47. Schweiger O, Biesmeijer JC, Bommarco R, Hickler T, Hulme PE,  
474 Klotz S, Kühn I, Moora M, Nielsen A, Ohlemüller R *et al.*: **Multiple**  
475 **stressors on biotic interactions: how climate change and alien**  
476 **species interact to affect pollination.** *Biol Rev* 2010, **85**:777-795. 504
- 477 48. Thomson D: **Competitive interactions between the invasive**  
478 **European honey bee and native bumble bees.** *Ecology* 2004,  
479 **85**:458-470. 505
- 480 49. Thomson DM: **Local bumble bee decline linked to recovery of**  
481 **honey bees, drought effects on floral resources.** *Ecol Lett* 2016,  
482 **19**:1247-1255. 506
- 483 The author uses a 15-year-long dataset on correlate increases in honey  
484 bee abundance to bumble bee declines and reduced niche overlap, with  
485 bumble bees shifting away from visiting a plant frequented by honey  
486 bees. Bumble bee abundance was also linked to key floral resources,  
487 which diminished with drought during the study period in California, USA.  
488 This paper demonstrates the important insights to be gained by long time  
489 series and the natural experiments that those time series can capture. 507
50. Roubik DW, Villanueva-Gutierrez R: **Invasive Africanized honey**  
511 **bee impact on native solitary bees: a pollen resource and trap**  
512 **nest analysis.** *Biol J Linn Soc* 2009, **98**:152-160. 487
51. Bjerknes A-L, Totland Ø, Hegland SJ, Nielsen A: **Do alien plant**  
513 **invasions really affect pollination success in native plant**  
514 **species?** *Biol Conserv* 2007, **138**:1-12. 488
52. Lopezarazola Mikel ME, Hayes RB, Whalley MR, Memmott J: **The**  
515 **impact of an alien plant on a native plant-pollinator network:**  
516 **an experimental approach.** *Ecol Lett* 2007, **10**:539-550. 489
53. Baskett CA, Emery SM, Rudgers JA: **Pollinator visits to**  
517 **threatened species are restored following invasive plant**  
518 **removal.** *Int J Plant Sci* 2011, **172**:411-422. 490
54. Kaiser-Bunbury CN, Mougil J, Whittington AE, Valentin T,  
519 • Gabriel R, Olesen JM, Blüthgen N: **Ecosystem restoration**  
520 **strengthens pollination network resilience and function.**  
521 *Nature* 2017, **542**:223-227. 491
- 522 After experimentally removing exotic plants from four sites, the authors  
523 compared plant-pollinator interactions and network metrics with control  
524 sites, finding increased pollinator species richness, visitation rates, and fruit  
525 set in the restored sites. This study is important for tying large-scale  
526 manipulation of exotic species presence to community function and pollina-  
527 tion services. 492
55. Goulson D, Nicholls E, Botias C, Rotheray EL: **Bee declines driven**  
528 **by combined stress from parasites, pesticides, and lack of**  
529 **flowers.** *Science* 2015, **347**:1255957. 493
56. Senapathi D, Carvalheiro LG, Biesmeijer JC, Dodson CA,  
530 Evans RL, McKechar M, Morton RD, Moss ED, Roberts SPM,  
531 Kunin WE *et al.*: **The impact of over 80 years of land cover**  
532 **changes on bee and wasp pollinator communities in England.**  
533 *Proc R Soc B* 2015, **282**:20150294. 494
57. Whitehorn PR, O'Connor S, Wackers FL, Goulson D: **Neonicotinoid**  
534 **pesticide reduces bumble bee colony growth**  
535 **and queen production.** *Science* 2012, **336**:351-352. 495
58. Gill RJ, Raine NE: **Chronic impairment of bumblebee natural**  
536 **foraging behaviour induced by sublethal pesticide exposure.**  
537 *Funct Ecol* 2014, **28**:1459-1471. 496
59. Rundlöf M, Andersson GKS, Bommarco R, Fries I, Hederström V,  
538 Herbertsson L, Jonsson O, Klatt BK, Pedersen TR, Yourstone J  
539 *et al.*: **Seed coating with a neonicotinoid insecticide negatively**  
540 **affects wild bees.** *Nature* 2015, **521**:77-80. 497
60. Stanley DA, Garratt MPD, Wickens JB, Wickens VJ, Potts SG,  
541 Raine NE: **Neonicotinoid pesticide exposure impairs crop**  
542 **pollination services provided by bumblebees.** *Nature* 2015,  
543 **528**:548-550. 498
61. Carvell C, Bourke AFG, Dreier S, Freeman SN, Hulmes S,  
544 • Jordan WC, Redhead JW, Sumner S, Wang J, Heard MS:  
545 **Bumblebee family lineage survival is enhanced in high-quality**  
546 **landscapes.** *Nature* 2017, **543**:547-549. 499
- 547 Combining a variety of methods, most notably molecular genetic ana-  
548 lyses, the authors connect the availability of floral resources in the spring  
549 and summer to the survival of bumble bee colony lineages from one year  
550 to the next. The results of this paper make a convincing case that  
551 measures to improve foraging habitat for native bees can positively affect  
552 their fitness and could help stem pollinator declines driven by land  
553 conversion for intensive agriculture. 500
62. Jha S, Kremen C: **Resource diversity and landscape-level**  
554 **homogeneity drive native bee foraging.** *Proc Natl Acad Sci U S A*  
555 2013, **110**:555-558. 501
63. Winfree R, Bartomeus I, Cariveau DP: **Native pollinators in**  
556 **anthropogenic habitats.** *Annu Rev Ecol Syst* 2011, **42**:1-22. 502
64. Weiner CN, Werner M, Linsenmair KE, Blüthgen N: **Land-use**  
557 **impacts on plant-pollinator networks: interaction strength**  
558 **and specialization predict pollinator declines.** *Ecology* 2014,  
559 **95**:466-474. 503
65. Williams NM, Crone EE, Roulston TH, Minckley RL, Packer L,  
560 Potts SG: **Ecological and life-history traits predict bee species**  
561 **responses to environmental disturbances.** *Biol Conserv* 2010,  
562 **143**:2280-2291. 504
66. Kovács-Hostyánszki A, Espíndola A, Vanbergen AJ, Settele J,  
563 Kremen C, Dicks LV: **Ecological intensification to mitigate**  
564 505

## 6 Global change biology

- 545 **impacts of conventional intensive land use on pollinators and** 552  
546 **pollination.** *Ecol Lett* 2017, **20**:673-689. 553
- 547 67. Forrest JRK, Chisholm SPM: **Direct benefits and indirect costs** 554  
548 **of warm temperatures for high-elevation populations of a**  
549 **solitary bee.** *Ecology* 2017, **98**:359-369. 555  
550 The authors examine the relationships among temperature, nest provision-  
551 cell production increased with temperature and with floral resource density,  
but because brood parasitism rate also increased with temperature, repro-  
ductive output was not elevated under warmer temperatures. This work  
illustrates the importance of top-down forces and community context for  
predicting the effects of climate change on insect pollinators.
68. Schmidt NM, Mosbacher JB, Nielsen PS, Rasmussen C, Høye TT, 556  
Roslin T: **An ecological function in crisis? The temporal overlap**  
**between plant flowering and pollinator function shrinks as the** 557  
**Arctic warms.** *Ecography* 2016, **39**:1250-1252. 558

UNCORRECTED PROOF