

## Article (refereed) - postprint

---

Vanbergen, Adam J., Espíndola, Anahí and Aizen, Marcelo A. 2018. **Risks to pollinators and pollination from invasive alien species**. *Nature Ecology & Evolution*, 2 (1). 16-25. <https://doi.org/10.1038/s41559-017-0412-3>

Copyright © 2017 Macmillan Publishers Limited, part of Springer Nature.

This version available <http://nora.nerc.ac.uk/518743/>

NERC has developed NORA to enable users to access research outputs wholly or partially funded by NERC. Copyright and other rights for material on this site are retained by the rights owners. Users should read the terms and conditions of use of this material at

<http://nora.nerc.ac.uk/policies.html#access>

**This document is the author's final manuscript version of the journal article, incorporating any revisions agreed during the peer review process. There may be differences between this and the publisher's version. You are advised to consult the publisher's version if you wish to cite from this article.**

[www.nature.com/](http://www.nature.com/)

Contact CEH NORA team at  
[noraceh@ceh.ac.uk](mailto:noraceh@ceh.ac.uk)

1 Risks to pollinators and pollination from invasive alien  
2 species

3 *Adam J. Vanbergen<sup>1</sup>, Anahí Espíndola<sup>2</sup> and Marcelo A. Aizen<sup>3</sup>*

4 1. NERC Centre for Ecology and Hydrology, Bush Estate, Penicuik, Edinburgh EH26 0QB, UK

5 2. Department of Biological Sciences, Life Sciences South 252, University of Idaho, 875 Perimeter Dr. MS  
6 3051, Moscow, ID 83844-3051, USA

7 3. Laboratorio Ecotono, INIBIOMA-CONICET and Centro Regional Bariloche, Universidad Nacional del  
8 Comahue, Quintral 1250, 8400 San Carlos de Bariloche, Rio Negro, Argentina

9

10 **Correspondence to [AJV](#).**

11 **e-mail: [ajv@ceh.ac.uk](mailto:ajv@ceh.ac.uk)**

12

13 Abstract

14 Invasive alien species modify pollinator biodiversity and the services they provide that underpin ecosystem  
15 function and human well-being. Building on the IPBES global assessment of pollinators and pollination, we  
16 synthesise current understanding of invasive alien impacts on pollinators and pollination. Invasive alien  
17 species create risks and opportunities for pollinator nutrition, re-organise species interactions to affect native  
18 pollination and community stability, and spread and select for virulent diseases. Risks are complex but  
19 substantial, and depend greatly on the ecological function and evolutionary history of both the invader and  
20 the recipient ecosystem. We highlight evolutionary implications for pollination from invasive alien species,  
21 and identify future research directions, key messages, and options for decision-making.

22

## 23 Introduction

24 Global anthropogenic drivers including land-use change, conventional intensive agriculture, pesticide use or  
25 misuse, pests and pathogens, and climate change threaten pollinators and pollination services<sup>1,2</sup>. Biological  
26 invasions are another major global change driver that can affect this natural capital<sup>1,3</sup>. The Convention on  
27 Biological Diversity ([www.cbd.int/invasive/WhatareIAS.shtml](http://www.cbd.int/invasive/WhatareIAS.shtml)) describes invasive alien species as those  
28 intentionally or accidentally introduced by human actions beyond natural ranges, which subsequently spread  
29 as vigorously growing populations that impact on biota, ecosystems and society. The global growth in  
30 economic wealth, trade, commerce, and transport efficiency facilitates this human-mediated spread of  
31 organisms into novel environments<sup>4-6</sup>, with implications for the benefits that humans derive from nature<sup>1</sup>.

32 Successful invaders have both ecological and evolutionary effects on native species and their  
33 interactions. Invasive alien species can alter the flow of energy and nutrients within an ecosystem<sup>4</sup>, and  
34 disrupt mutualisms including those underpinning crop and wild plant reproduction<sup>7-9</sup>. Strongly interacting  
35 alien invaders can also establish novel selection pressures within a community that can modify evolutionary  
36 trajectories and adversely affect species with low genetic diversity and/or small effective population sizes<sup>10-</sup>  
37 <sup>12</sup>.

38 Scientific and policy concern over various threats to pollinators and pollination led the  
39 Intergovernmental Science-Policy Platform for Biodiversity and Ecosystem Services (IPBES) to carry out a  
40 global evidence-based assessment on their values to humanity, their status and trends and drivers of change,  
41 and to identify policy response options to conserve them for the future<sup>3,13</sup>. In 2016, the Parties to the  
42 Convention on Biological Diversity (CBD-COP13) endorsed the findings of this IPBES assessment.

43 In this review, we build on the peer-reviewed IPBES evaluation<sup>3,13</sup> and earlier review papers<sup>14-17</sup> to  
44 synthesise the current understanding of impacts on pollinators and pollination from invasive alien species  
45 spanning different ecological functions (**Fig. 1**). We evaluate the negative, neutral or positive impacts of: 1)  
46 alien flowering plants on pollinator nutrition, community assembly and native pollination; 2) introduced  
47 alien pollinators on native plant-pollinator systems via competition, genetic exchange and pathogen and  
48 parasite transfer to new hosts; and 3) alien predators that consume pollinators and transform pollination  
49 systems. We outline potential risks to evolutionary dynamics from invasive aliens (**Box 1**) and conclude by  
50 identifying future research directions, key messages, and recommendations for decision-making.

51

## 52 **Invasive alien plants**

53 Global human-mediated dispersal of alien plants has increased, both accidentally (e.g. contamination of  
54 agricultural cargo) and deliberately (e.g. horticultural species)<sup>4-6</sup>. Introduced alien plants may prosper by  
55 escaping biological regulation of population size, by occupying a vacant ecological niche in the recipient  
56 ecosystem, or by possessing or evolving phenotypic traits (e.g. novel defences) that confer competitive  
57 advantage over native plant species<sup>4,8,18</sup>. Insect-pollinated species represent a large proportion of documented  
58 invasive alien plants; however, the capacity for self-pollination often aids initial establishment and spread<sup>19</sup>.  
59 Thereafter, invasive alien plant species that become abundant, and possess copious nectar and pollen rewards  
60 or large and enduring floral displays can lure and co-opt pollinators adapted to exploit such floral resources  
61 (**Fig. 2**)<sup>19-22</sup>. In this manner, invasive alien plant species can dominate species interactions and the diet and  
62 community structure of pollinators<sup>19,23-25</sup>.

### 63 *Impacts on pollinator nutrition*

64 Whilst providing a substantial food resource for pollinators<sup>19,24,26</sup>, a predominance of alien pollen and nectar  
65 in pollinator diets may produce risks for pollinator health. Pollinator species have particular physiological  
66 requirements for energy and a diversity of macronutrients<sup>27-29</sup>, and they forage to balance these needs over  
67 time at both individual and colony levels<sup>26,30-32</sup>. Alien plant domination of floral communities can transform  
68 pollinator diet from a diverse suite of floral species to a largely monotypic diet comprising alien pollen and  
69 nectar (**Fig. 2**)<sup>25</sup>. Pollinating bees are highly sensitive to the specific dietary source and combination of  
70 nutrients, e.g. ratio of different essential amino acids (EAA) to carbohydrates, showing poor growth and  
71 survival when reared on monotypic or nutritionally sub-optimal diets<sup>29,30,33,34</sup>. Consequently, alien plant  
72 invasions may raise the risk of nutritional deficits for pollinators by eroding the ecosystem availability of  
73 combinations of essential nutrients provided by diverse floral resources. Alternatively, invasive alien plants  
74 can adequately supply carbohydrates or essential amino acids exploitable by pollinators with generalized  
75 foraging behaviour and diet<sup>26</sup>. However, the subtle nutrient requirements of pollinators, e.g. protein to lipid  
76 or EAA combinations, and a species' capacity to balance nutrition through flexible foraging<sup>29,30</sup> mean that  
77 the benefits of invasive pollen and nectar for native pollinators remain to be determined. Adverse impacts of  
78 alien pollen or nectar are more likely for relatively specialized pollinator species, either physiologically or

79 morphologically ill adapted to exploit the alien food resource, or dependent on native plants outcompeted by  
80 the invader<sup>34-36</sup>. Secondary compounds in alien pollen and nectar can be differentially toxic to native  
81 pollinator species representing a further risk from plant invasions where they come to dominate diets<sup>34,37,38</sup>.

82 Dominance of plant communities by invasive alien species (**Fig. 2**) could also restrict community-  
83 wide flowering phenology, truncating the period of floral resources' availability. Such curtailment could  
84 cause pollinator population declines and an overall decrease in pollinator diversity, as proposed for  
85 agricultural landscapes<sup>39</sup>. Surprisingly, there are comparatively few recorded examples of alien plant  
86 invasions consistently lowering overall pollinator diversity or abundance<sup>40-42</sup>.

87 Although more research is definitively needed, this scarce evidence implies that pollinators may  
88 either physiologically or behaviourally trade-off or compensate for spatial and temporal changes in nutrient  
89 availability due to invasive alien plants<sup>26</sup>, that effects are subtle, chronic and possibly undetected hitherto, or  
90 that they only adversely affect pollinators in combination with other stressors<sup>2,43</sup>.

#### 91 *Modified interactions and community stability*

92 The dynamic and flexible nature of pollinator foraging behaviour<sup>30,44,45</sup> means interaction networks are  
93 readily penetrated by flowering alien plants<sup>23,46</sup>, where they often assume a key role in community  
94 organisation and function<sup>7,8</sup> (**Fig. 2**). Where the invasive alien plant species is highly abundant or possesses  
95 generalized floral traits that make it highly attractive to pollinators, it can rewire interspecific interactions to  
96 modify network architecture (**Fig. 2**)<sup>44,45,47,48</sup>. For example, they can usurp native interactions and operate as  
97 a hub that increases the size and connectivity of network modules (subsets of highly co-dependent species)<sup>7</sup>,  
98 or weaken the co-dependency of mutualistic relationships in the network<sup>49</sup>. Such changes in modularity and  
99 interaction strength<sup>7,49</sup> can increase community stability by lowering the risk of co-extinction cascades  
100 arising from future environmental changes<sup>50</sup>, unless the invasive alien performing the central role in the  
101 network is itself extirpated. Conversely, as seen with habitat structure, the high dominance of invasive alien  
102 plants could erode the co-phylogenetic structure of native plant-pollinator networks, reflecting poorer  
103 phenotypic matching between interacting partners and less-fitted mutualism, potentially introducing  
104 instability and reduced function of the pollination system<sup>48,51</sup>.

#### 105 *Disrupted native pollination*

106 The influential functional position of invasive alien plants once integrated into pollinator networks may have  
107 ramifications for native plant species reproduction. Invasive alien plants may affect co-flowering native  
108 plants by elevating pollinator activity to facilitate native pollination<sup>22,52,53</sup>. However, if an invasive alien plant  
109 reduces the abundance of native plants that become overly reliant on the invader for facilitation of  
110 pollination services, then there is a potential risk to the native species, should those connections become  
111 eroded or lost due to further environmental changes. Alternatively, invasive alien plants may simply  
112 outcompete native plants for pollinators (**Fig. 2**) and meta-analyses suggest native plant visitation rates do  
113 tend to decrease, indicating that competition prevails<sup>48,54-57</sup>. Whether regional facilitation or local competition  
114 predominates may depend on the spatial scale of the alien plant invasion, and the differing foraging ranges  
115 and ecology of pollinators in the species pool<sup>53,58-60</sup>. Overall, the impact of alien plant invasions on native  
116 plant pollination and reproductive success is greater if, relative to the native flora, the alien produces higher  
117 densities of flowers, they are phylogenetically related, or they possess similar phenology and anatomy of  
118 floral displays<sup>9,19,59,61</sup>. Aside from fundamental competition for pollinators, there may also be native pollen  
119 loss and pick up of foreign pollen during visits to alien flowers. This could either reduce conspecific native  
120 pollen transfer or increase deposition of heterospecific alien pollen that could cause stigma clogging or  
121 chemical inhibition of pollen germination<sup>62</sup>. This improper pollen transfer can translate into reduced native  
122 plant reproduction<sup>55,57,63,64</sup>, yet the extent of this is complicated by plant compensatory mechanisms that can  
123 assure pollination and reproduction, such as the capacity for self-reproduction or recruitment of alternative  
124 pollinators<sup>14,54,56,65</sup>.

## 125 **Invasive alien pollinators**

### 126 *Competitive exclusion and co-existence*

127 Humans have globally translocated many different bee species (e.g. species of *Apis*, *Bombus*, *Osmia*,  
128 *Megachile*) for apiculture and crop pollination services<sup>13,66-68</sup>. The principal managed pollinators, the western  
129 honeybee *Apis mellifera* and the bumblebee *Bombus terrestris*, possess traits such as sociality, generalist  
130 feeding habit and nesting flexibility, that coupled to recurrent introduction of managed colonies and frequent  
131 escape and establishment of feral populations, raise the risk of competition with native species<sup>66,69-72</sup> (**Fig. 1**).  
132 Direct competition from alien honeybees has altered the behaviour and reproductive success of native  
133 pollinators<sup>69,73</sup>. Given their long history of global spread, however, there are surprisingly few accounts of

134 honeybee competition reducing survival or densities of native wild bee species and no reported  
135 extinctions<sup>67,74-76</sup>. One possibility is that the introduced super-generalist honeybee, by occupying a distinct  
136 ecological niche, becomes readily integrated into native pollinator networks, apparently with little  
137 competitive displacement of native pollinators<sup>77,78</sup>. Alternatively, the role of alien honeybees in historic  
138 declines of native pollinators, while noted in certain regions (e.g. decline of congener *Apis cerana* in China)  
139 may have contributed to declines in places like oceanic islands, but gone unrecorded<sup>75,79</sup>. In contrast,  
140 introduced alien bumblebee species, typically *B. terrestris*, often compete with native congeners that occupy  
141 very similar niches for nesting and floral resources, leading to the invader becoming dominant and excluding  
142 natives<sup>66,70,71</sup>. An example is the extirpation of the Patagonian giant bumblebee *Bombus dahlbomii* from most  
143 of its range following the introduction and subsequent establishment of feral populations of managed  
144 European bumblebee species (*B. terrestris* and *B. ruderatus*)<sup>66</sup> (**Fig. 1**).

#### 145 *Genetic effects and mating interference*

146 Another potential risk from anthropogenic introductions of bee species is intra-generic hybridization and  
147 introgression, and reductions of native species fitness through mating interference<sup>80-82</sup>. Despite the history of  
148 global translocation of *A. mellifera*, overall evidence of hybridizations, introgression or mating interference  
149 with endemic sub-species is scant<sup>67,83</sup>. A notable exception was the movement of *A. mellifera capensis* into  
150 the range of *A. m. scutellata* as part of migratory beekeeping in South Africa, where it behaved as a social  
151 parasite, resulting in substantial *A. m. scutellata* colony losses<sup>67,83</sup>. Another example, from South America,  
152 was the introduction (>250 years ago), establishment of feral populations and spread of managed stocks of  
153 European *A. mellifera*, and more recently (1956) an African sub-species (*A. m. scutellata*) regarded as better  
154 suited to tropical environments. Debate continues about the extent that hybridization and introgression of the  
155 European type occurred, nonetheless there seems to be a latitudinal gradient in the extent of hybridization  
156 and the type possessing so-called ‘African’ traits came to dominate bee assemblages across the Neotropics  
157 and Southern USA<sup>67,84</sup>.

#### 158 *Pollination disruption or rescue*

159 Introduced pollinators can influence native pollination processes in complex ways, according to the identity  
160 of the pollinators and the nature of the recipient ecosystem<sup>53</sup>. There is evidence that the introduced  
161 honeybee’s foraging behaviour, i.e. social recruitment of numerous worker bees to a floral resource, can

162 effectively maintain pollination function over great distances, particularly where the ecosystem and  
163 indigenous pollinators have been disrupted by anthropogenic habitat loss and species invasions<sup>77,84,85</sup>.  
164 Interactions between naturalized honeybees and native pollinators have been seen to enhance pollination of  
165 native plants and crops, additively or synergistically<sup>86,87</sup>. However, alien pollinators are efficient pollen  
166 collectors and nectar robbers, so at high densities they can also behave as antagonists rather than mutualists,  
167 adversely affecting plant pollination<sup>72,88</sup>, as seen in South America where frequent visits by abundant  
168 invasive bumblebees reduce crop yields<sup>89</sup>. A preponderance of invasive alien pollinators that either prefer or  
169 are able to exploit alien forage plants, may also produce less effective native mutualisms. To illustrate,  
170 removal of invasive plant species from a Seychelles island ecosystem decreased the domination by invasive  
171 *A. mellifera* of plant-pollinator networks; correspondingly increasing network flower visitation, interaction  
172 diversity and functional redundancy, which resulted in higher fruit production of native plants<sup>48</sup>. Alien  
173 pollinators, by altering mutualistic networks, can raise the likelihood of inbreeding depression via increased  
174 selfing within plant species, or outbreeding depression through hybridization between closely related alien  
175 and native plants<sup>62,66,69,90</sup>. Ultimately, such changes represent a risk to plant fitness, community structure and  
176 function.

### 177 ***Introduction of alien pests and pathogens***

178 An outcome of the trans-continental transport of pollinating bees beyond their native ranges is the greater  
179 likelihood of pathogen and parasite transfer to new hosts, with the potential to elicit population declines of  
180 native pollinators<sup>66,91,92</sup> (**Fig. 1**). Introductions of *A. mellifera* to China in 1896 coincided with a drastic  
181 reduction in the range and population size of the Asian honeybee *A. cerana* with interspecific competition  
182 and pathogen transfer (e.g. Sacbrood viruses) implicated<sup>75,93</sup>. The sustained movement by humans of  
183 managed honeybee (*A. mellifera*) colonies into Asia ultimately resulted in the host shift of the ectoparasitic  
184 *Varroa* mite from sympatric *A. cerana* populations and its subsequent worldwide spread, along with a  
185 complex of viral pathogens (*Picornavirales*) it transmits among bee hosts, as part of trade in managed honey  
186 bees<sup>94,95</sup> (**Fig. 3**). Through vectoring viruses, possibly suppressing bee immune functions, and direct parasitic  
187 feeding the *Varroa* mite is among the major pressures impacting managed and feral honeybee colonies<sup>1,2,96</sup>.  
188 Indeed, the most recent analyses suggest that the *Varroa* host shift may have elicited eco-evolutionary  
189 changes in host-vector-pathogen dynamics resulting in selection for increased virulence of strains of

190 Deformed Wing Virus (DWV) infecting honeybees and implicated in colony losses<sup>94-98</sup> (**Fig. 3**). Moreover,  
191 there are also signs of pathogen transmission between managed bee populations and wild pollinators<sup>91,95,99,100</sup>.  
192 Possibly these pathogens are generalists infecting a broad spectrum of hosts and commonly shared across  
193 flower-visiting insects<sup>100,101</sup>. Alternatively, pathogens introduced along with alien pollinators, managed or  
194 feral, might represent a novel ecological and selective pressure with consequences for pollinator decline and  
195 the epidemiology of pollinator communities (**Fig. 3**).

## 196 **Invasive alien predators**

197 Invasive alien predators, such as cats, rats, and stoats, spread by humans often exert strong top-down  
198 pressure on plant pollination and fitness by consumption of pollinators such as birds, lizards, bats and other  
199 small mammals<sup>13</sup> (**Fig. 1**), especially in the specialised and simpler networks of island ecosystems<sup>6</sup>. A recent  
200 example of a direct threat to already stressed European honey bee populations is the accidental introduction  
201 (2004) of the predatory yellow-legged hornet (*Vespa velutina*) into Europe from Asia<sup>102,103</sup> (**Fig. 1**).

202 Alien predators can also indirectly shift the functioning of native pollination systems through  
203 networks of trophic and competitive interactions. For instance, in Africa, California and Mauritius, invasive  
204 ant species that are more aggressive or competitive than native ants, deter pollinators and seed dispersers  
205 thereby reducing plant fitness<sup>104-106</sup>. Alien insectivorous lizards transformed the pollination system of the  
206 Ogasawara archipelago of Japan by extirpating endemic bee species and leaving the alien honeybee (*A.*  
207 *mellifera*) that prefers flowers of invasive alien plants to dominate, thus completing the shift to an invasive-  
208 dominated pollination ecology<sup>107</sup>.

209 A case that highlights the complex nature of interactions between predators, pollinators and plants is  
210 that of the invasive predatory wasp (*Vespula pensylvanica*) in Hawaii<sup>72,77</sup>. This generalist predator of  
211 arthropods also behaves as a nectar thief, competing with native *Hylaeus* bees and the alien honeybee *A.*  
212 *mellifera* that pollinate the native tree *Metrosideros polymorpha*, thereby lowering pollinator visitation and  
213 resultant fruit production<sup>72,77</sup> (**Fig. 4**). Experimental removal of the wasp revealed the alien *A. mellifera* was  
214 the most effective pollinator in this system, in all likelihood fulfilling a niche previously occupied by extinct  
215 or declining bird pollinators, themselves reduced by introduced vertebrate predators<sup>77</sup> (**Fig. 4**). These  
216 examples serve to illustrate the impact that alien predators can have on the community of interactions

217 affecting pollination, but also how invasive alien pollinator species can maintain pollination in highly  
218 modified ecosystems in the absence of native pollinators.

### 219 **Future research directions**

220 Invasive alien species remain an ongoing threat to pollinator biodiversity and pollination function  
221 worldwide. Nonetheless, our ability to understand and forecast the risk to pollinators and pollination  
222 requires that we fill substantial gaps in knowledge by stimulating future biological, ecological and  
223 evolutionary research.

224         The impact of particular invasive alien species on native pollinators and pollination has been  
225 somewhat overlooked. The impact of introduced solitary bees on the ecology of native pollinators and  
226 pollination is a specific gap in knowledge and risk assessment, warranting further study to help forecast and  
227 prevent future invasions by alien pollinators. For instance, solitary bees such as species of *Osmia* or  
228 *Megachile*, introduced for crop pollination services, sometimes possess similar traits (e.g. dietary  
229 generalism) to the bee species *A. mellifera* and *B. terrestris*, which facilitated the invasion and modification  
230 of native mutualisms by these social bees<sup>13,66-68</sup>. There has also been little investigation of herbivory as an  
231 aspect of pollination invasion ecology, compared to other trophic interactions. Introduced mammalian  
232 herbivores can modify plant communities affecting the floral or nesting resources available to native  
233 pollinators and influencing native plant pollination<sup>108,109</sup>; given the global prevalence of livestock  
234 introductions, this is an understudied research area. Similarly, insect herbivory can influence plant  
235 physiological function and allocations of metabolites to floral displays, pollen and nectar, and emissions of  
236 volatile organic compounds that recruit pollinators<sup>110</sup> and affect pollination<sup>111-113</sup>. Yet, the impact of invasive  
237 insect herbivory on the chemical ecology of native pollination remains a significant knowledge gap with  
238 considerable research potential.

239         Much remains to be discovered about the impact of invasive alien species on the structure, function  
240 and stability of plant-pollinator networks. Henceforward, research should employ recent innovations in  
241 simulation modelling that capture greater biological realism and complexity of species interactions - such as  
242 temporal dynamics, interference competition, variable mutualism dependence - to obtain new insights on  
243 how invasive species re-organise pollinator network structure and affect key mechanisms or properties  
244 underpinning the stability of invaded networks facing future global change<sup>45,48,50,114,115</sup>. Furthermore,

245 research on network structure and stability should be extended beyond impacts from alien plants and alien  
246 pollinators to other invasive groups occupying different trophic or parasitic roles and evaluate the overall  
247 consequences for interconnected mutualistic and antagonistic networks<sup>116</sup>.

248 Research must continue to understand the community dynamics of invasions and their consequences  
249 for pollination processes. We know little about the consequences of massive plant species invasions for  
250 community-wide flowering phenology, and how such temporal changes in distribution of floral resources  
251 link to changes in the temporal dynamics, composition, and diversity of pollinator communities. The extent  
252 that co-flowering native plant species, through their influence on foraging behaviour of different pollinator  
253 groups (e.g. flies, bees, birds), facilitate alien plant establishment is a gap in understanding the dynamics of  
254 alien plant invasions<sup>117</sup>. Similarly, the impact on agricultural crop production of changes in pollinator  
255 foraging due to invasive alien plants has yet to be well studied<sup>60</sup>. Furthermore, by usurping native  
256 interactions<sup>7,49</sup>, alien plant and pollinator species may increase the proportion of ill-matched interactions and,  
257 therefore, decrease pollination function, an untested hypothesis based on a relatively well-established  
258 assumption with important ecological and evolutionary consequences.

259 Evolutionary mechanisms facilitating or hindering invasions by mutualists are largely at a theoretical  
260 stage<sup>10,118</sup>, but recent observations show how rapid adaptation in invading plant populations may aid their  
261 spread and establishment, and also the role of balancing selection at the sex locus of *A. cerana* enabling its  
262 recent establishment in Australia<sup>11,119</sup>. More empirical research is needed to test predictions such as  
263 understanding micro-evolutionary effects, shifting trait structure of plant-pollinator networks, or the role of  
264 genetic diversity in shaping invasion probabilities and dynamics in an ecosystem (see **Box 1**). We need to  
265 understand better the eco-evolutionary constraints to invasion of pollinator communities and their effects on  
266 evolutionary trajectories post- invasion to predict future risk. For instance, community permeability to an  
267 invasive species may be limited by the genetic diversity or the effective population size of the invading  
268 populations, governing their ability to adapt to novel environments. Genetic variability in the native  
269 populations with which the invader will interact may contribute to the success or failure of the invasions,  
270 depending of the type of interaction (e.g. competitive, mutualistic) established with the invasive species.  
271 Once established, an invader has the potential to affect the evolvability of native species, since introductions  
272 can affect the (effective) population sizes, the genetic diversity and the fitness of native populations (**Box 1**).

273           There is considerable scope for increasing our knowledge about the disease risks for native  
274 pollinators and pollination from exposure to invasive alien species. The epidemiology of pollinator  
275 communities is in its infancy with recent detection of pathogen sharing and potential asymmetric  
276 interspecific transmission and virulence<sup>91,95,99,100</sup>. There is an opportunity to unify network theory, evolution,  
277 disease biology and ecology to understand how novel host-vector-pathogen shifts involving alien organisms  
278 affect the evolution of pathogen virulence within hosts; competition and coexistence among assemblages of  
279 ecotoparasites, viral, fungal and bacterial pathogens; and transmission processes and disease frequency  
280 among multiple pollinator hosts<sup>2,96</sup>. Related to this, there is a need to study the underlying mechanisms for  
281 pathogen resistance/tolerance among bee species in their native and invaded ranges, including those living  
282 wild and those reared commercially (e.g. *B. terrestris*)<sup>120</sup>. Furthermore, global trade in agricultural  
283 commodities or the human-mediated translocation of alien plant species increase the risk of spreading alien  
284 plant pathogens<sup>121</sup>. There is some evidence that plant pathogens in native systems may modify plant  
285 physiology and flowering to affect plant-pollinator interactions and plant reproduction<sup>122,123</sup>, but this  
286 possibility during invasion of pollination systems has been hitherto ignored. Moreover, a single study  
287 provides some evidence that a plant pathogenic RNA virus (TRSV) due to its evolutionary history may  
288 infect bees via *Varroa* mite vectors, albeit without apparent effects on bee colony health, intriguingly  
289 pointing to the potential for viruses to transcend kingdoms<sup>124</sup>. Overall, the biological and evolutionary  
290 complexity and phylogenetic breadth of potential plant-pollinator-pathogen epidemiology arising from  
291 species invasions is considerable and warrants investigation.

## 292 **Conclusions and policy responses**

293 The effects of invasive alien species on pollinators and pollination are complex and substantial, particularly  
294 under the biogeographical circumstances of oceanic islands<sup>6,13</sup>, but depend greatly on the functional ecology  
295 and phylogenetic history of the invader and the recipient ecosystem. For example, invasive alien species  
296 possessing generalised ecological traits or evolutionarily close to natives are readily incorporated into species  
297 networks and ecosystems, and when attaining great abundance, they substantially modify structure and  
298 function of pollination systems, often negatively for native species. Alien predators exert considerable top-  
299 down pressure on native pollination systems through direct and more subtle indirect trophic interactions that  
300 can transform the pollination ecology into a state dominated by alien interactions. Global trade in managed

301 bees and horticultural or agricultural plants increases disease risks through the interspecific spread and  
302 selection of novel pathogens with the potential to impact pollinators and pollination in unforeseen ways.  
303 Invasive alien species thus tend to represent a significant biological risk to pollinators and pollination, albeit  
304 one that varies with species identity, abundance and environmental context.

305 In the globalised economy, there is considerable scope for interactions among drivers of biodiversity  
306 change, thus the impact of invasive alien species on pollinators and pollination is exacerbated or complicated  
307 when it occurs in combination with other threats such as diseases, climate or land-use change<sup>2,6,43</sup>. Policies  
308 that minimize impacts on pollinators from stresses such as conventional intensive agricultural management  
309 and climate change, for example by diversifying agricultural landscapes and building ecological  
310 infrastructure<sup>1,2,13,125</sup>, are likely to relieve some of this overall multifactorial pressure on pollinators. In  
311 principal, this could increase the resilience of native plant-pollinator communities to alien species invasions.  
312 Current and future research focused on the interplay between invasive species and other global change  
313 drivers affecting pollinator biodiversity in different ecosystems will enable subsequent refinement of  
314 intergovernmental policy (e.g. CBD) tackling invasive alien species.

315 Eradication or control of established invasive aliens is often prohibitively expensive and rarely  
316 successful beyond oceanic islands and vertebrate species. Consequently, the most effective policy response is  
317 a tiered approach to mitigate the risk. Crucial to forestalling invasions is horizon scanning for emerging  
318 threats and forecasting likely impacts, which allows for timely scientific, technical and policy  
319 responses<sup>3,103,126,127</sup>. Thereafter, actions leading to improving regulation, e.g. of trade in managed pollinators  
320 or horticultural plants, maintaining surveillance and establishing rigorous monitoring<sup>3,126,128</sup>, and once  
321 detected, rapid assertive management to avoid establishment by the alien species are expected to prevent new  
322 invasions or limit their impacts<sup>3,126</sup>. If invasive alien species go unchecked, the risk to pollinators and  
323 pollination is elevated, ultimately with unpredictable but mostly negative consequences for ecosystem health  
324 and human well-being<sup>1</sup>.

325

## 326 **Competing interests**

327 There are no competing interests.

## 328 **Author Contributions**

329 In the cited IPBES report AJV and AE authored the section in Chapter 2 on invasive species impacts on  
330 pollinators and pollination. AJV conceived and led this article, AE & MAA provided insight, co-wrote the  
331 review and all authors performed revisions following peer review.

### 332 **Acknowledgements**

333 We thank for their leadership, support, input and comments Simon G. Potts and Vera Imperatriz-Fonseca  
334 (co-chairs), Hien Ngo (secretary) and all the expert authors and reviewers of the IPBES Assessment Report  
335 on Pollinators, Pollination and Food Production<sup>3,13</sup>. We also thank Heather Lowther for creating the figures  
336 in this review. AJV was supported by NERC-CEH National Capability Funding (NEC05106). AE was  
337 supported by the Swiss National Science Foundation (grants P300P3\_151141 and PBNEP3\_140192). MAA  
338 was partially supported by FONCYT (PICT 2015-2333).

### 339 **References**

340

- 341 1 Potts, S. G. *et al.* Safeguarding pollinators and their values to human well-being. *Nature*.  
342 **540**, 220–229, doi:10.1038/nature20588 (2016).
- 343 2 Vanbergen, A. J. & the Insect Pollinators Initiative. Threats to an ecosystem service:  
344 pressures on pollinators. *Front. Ecol. Environ.* **11**, 251–259, doi:10.1890/120126 (2013).
- 345 3 IPBES. Summary for Policymakers of the Assessment Report of the Intergovernmental  
346 Science-Policy Platform on Biodiversity and Ecosystem Services on Pollinators, Pollination  
347 and Food Production. S.G. Potts *et al.* (eds.) Secretariat of the Intergovernmental Science-  
348 Policy Platform on Biodiversity and Ecosystem Services, Bonn, Germany. 36 (2016).
- 349 4 Mack, R. N. *et al.* Biotic invasions: causes, epidemiology, global consequences, and control.  
350 *Ecol. Appl.* **10**, 689–710, doi:10.1890/1051-0761(2000)010[0689:bicegc]2.0.co;2 (2000).
- 351 5 Hulme, P. E. Trade, transport and trouble: managing invasive species pathways in an era of  
352 globalization. *J. Appl. Ecol.* **46**, 10–18, doi:10.1111/j.1365-2664.2008.01600.x (2009).
- 353 6 Dawson, W. *et al.* Global hotspots and correlates of alien species richness across taxonomic  
354 groups. *Nature Ecology & Evolution.* **1**, 0186, doi:10.1038/s41559-017-0186 (2017).

- 355 7 Albrecht, M., Padrón, B., Bartomeus, I. & Traveset, A. Consequences of plant invasions on  
356 compartmentalization and species' roles in plant–pollinator networks. *Proc. Roy. Soc. B.*  
357 **281**, doi:10.1098/rspb.2014.0773 (2014).
- 358 8 Goodell, K. & Parker, I. M. Invasion of a dominant floral resource: effects on the floral  
359 community and pollination of native plants. *Ecology*. **98**, 57-69, doi:10.1002/ecy.1639  
360 (2017).
- 361 9 Herron-Sweet, C. R., Lehnhoff, E. A., Burkle, L. A., Littlefield, J. L. & Mangold, J. M.  
362 Temporal- and density-dependent impacts of an invasive plant on pollinators and pollination  
363 services to a native plant. *Ecosphere*. **7**, e01233, doi:10.1002/ecs2.1233 (2016).
- 364 10 Jones, E. I. & Gomulkiewicz, R. Biotic interactions, rapid evolution, and the establishment  
365 of introduced species. *Am. Nat.* **179**, E28-36, doi:10.1086/663678 (2012).
- 366 11 Vandepitte, K. *et al.* Rapid genetic adaptation precedes the spread of an exotic plant species.  
367 *Mol. Ecol.* **23**, 2157-2164, doi:10.1111/mec.12683 (2014).
- 368 12 Bossdorf, O. *et al.* Phenotypic and genetic differentiation between native and introduced  
369 plant populations. *Oecologia*. **144**, 1-11, doi:10.1007/s00442-005-0070-z (2005).
- 370 13 IPBES. The assessment report of the Intergovernmental Science-Policy Platform on  
371 Biodiversity and Ecosystem Services on pollinators, pollination and food production. S.G.  
372 Potts, V. L. Imperatriz-Fonseca, & H. T. Ngo, (eds). Secretariat of the Intergovernmental  
373 Science-Policy Platform on Biodiversity and Ecosystem Services, Bonn, Germany. (2016).
- 374 14 Traveset, A. & Richardson, D. M. Mutualistic interactions and biological invasions. *Ann.*  
375 *Rev. Ecol. Evol. Sys.* **45**, 89-113, doi:10.1146/annurev-ecolsys-120213-091857 (2014).
- 376 15 Traveset, A. & Richardson, D. M. Biological invasions as disruptors of plant reproductive  
377 mutualisms. *Trends Ecol. Evol.* **21**, 208-216, doi:10.1016/j.tree.2006.01.006 (2006).
- 378 16 Bjercknes, A. L., Totland, O., Hegland, S. J. & Nielsen, A. Do alien plant invasions really  
379 affect pollination success in native plant species? *Biol. Conserv.* **138**, 1-12,  
380 doi:10.1016/j.biocon.2007.04.015 (2007).

- 381 17 Stout, J. C. & Morales, C. L. Ecological impacts of invasive alien species on bees.  
382 *Apidologie*. **40**, 388-409, doi:10.1051/apido/2009023 (2009).
- 383 18 Uesugi, A. & Kessler, A. Herbivore exclusion drives the evolution of plant competitiveness  
384 via increased allelopathy. *New Phytol.* **198**, 916-924, doi:10.1111/nph.12172 (2013).
- 385 19 Pysek, P. *et al.* Successful invaders co-opt pollinators of native flora and accumulate insect  
386 pollinators with increasing residence time. *Ecol. Monogr.* **81**, 277-293, doi:10.1890/10-  
387 0630.1 (2011).
- 388 20 Morales, C. L. & Traveset, A. A meta-analysis of impacts of alien vs. native plants on  
389 pollinator visitation and reproductive success of co-flowering native plants. *Ecol. Lett.* **12**,  
390 716-728, doi:10.1111/j.1461-0248.2009.01319.x (2009).
- 391 21 Chrobock, T. *et al.* Effects of native pollinator specialization, self-compatibility and  
392 flowering duration of European plant species on their invasiveness elsewhere. *J. Ecol.* **101**,  
393 916-923, doi:10.1111/1365-2745.12107 (2013).
- 394 22 Masters, J. A. & Emery, S. M. The showy invasive plant *Ranunculus ficaria* facilitates  
395 pollinator activity, pollen deposition, but not always seed production for two native spring  
396 ephemeral plants. *Biol. Invasions.* **17**, 2329-2337, doi:10.1007/s10530-015-0878-3 (2015).
- 397 23 Traveset, A. *et al.* Invaders of pollination networks in the Galápagos Islands: emergence of  
398 novel communities. *Proc. Roy. Soc. B.* **280**, doi:10.1098/rspb.2012.3040 (2013).
- 399 24 Vilà, M. *et al.* Invasive plant integration into native plant-pollinator networks across Europe.  
400 *Proc. Roy. Soc. B.* **276**, 3887-3893, doi:10.1098/rspb.2009.1076 (2009).
- 401 25 Kleijn, D. & Raemakers, I. A retrospective analysis of pollen host plant use by stable and  
402 declining bumble bee species. *Ecology.* **89**, 1811-1823 (2008).
- 403 26 Harmon-Threatt, A. N. & Kremen, C. Bumble bees selectively use native and exotic species  
404 to maintain nutritional intake across highly variable and invaded local floral resource pools.  
405 *Ecol. Entomol.* **40**, 471-478, doi:10.1111/een.12211 (2015).

- 406 27 Sedivy, C., Muller, A. & Dorn, S. Closely related pollen generalist bees differ in their ability  
407 to develop on the same pollen diet: evidence for physiological adaptations to digest pollen.  
408 *Funct. Ecol.* **25**, 718-725, doi:10.1111/j.1365-2435.2010.01828.x (2011).
- 409 28 Paoli, P. *et al.* Nutritional balance of essential amino acids and carbohydrates of the adult  
410 worker honeybee depends on age. *Amino Acids.* **46**, 1449-1458, doi:10.1007/s00726-014-  
411 1706-2 (2014).
- 412 29 Stabler, D., Paoli, P. P., Nicolson, S. W. & Wright, G. A. Nutrient balancing of the adult  
413 worker bumblebee (*Bombus terrestris*) depends on the dietary source of essential amino  
414 acids. *J. Exp. Biol.* **218**, 793-+, doi:10.1242/jeb.114249 (2015).
- 415 30 Vaudo, A. D., Patch, H. M., Mortensen, D. A., Tooker, J. F. & Grozinger, C. M.  
416 Macronutrient ratios in pollen shape bumble bee (*Bombus impatiens*) foraging strategies and  
417 floral preferences. *PNAS.* **113**, E4035-E4042, doi:10.1073/pnas.1606101113 (2016).
- 418 31 Requier, F. *et al.* Honey bee diet in intensive farmland habitats reveals an unexpectedly high  
419 flower richness and a major role of weeds. *Ecol. Appl.* **25**, 881-890, doi:10.1890/14-1011.1  
420 (2014).
- 421 32 Jha, S. & Kremen, C. Resource diversity and landscape-level homogeneity drive native bee  
422 foraging. *PNAS.* **110**, 555-558, doi:10.1073/pnas.1208682110 (2013).
- 423 33 Tasei, J.-N. & Aupinel, P. Nutritive value of 15 single pollens and pollen mixes tested on  
424 larvae produced by bumblebee workers (*Bombus terrestris*, Hymenoptera: Apidae).  
425 *Apidologie.* **39**, 397-409, doi:10.1051/apido:2008017 (2008).
- 426 34 Praz, C. J., Müller, A. & Dorn, S. Specialized bees fail to develop on non-host pollen: do  
427 plants chemically protect their pollen? *Ecology.* **89**, 795-804, doi:10.1890/07-0751.1 (2008).
- 428 35 Palladini, J. & Maron, J. Reproduction and survival of a solitary bee along native and exotic  
429 floral resource gradients. *Oecologia.* **176**, 789-798, doi:10.1007/s00442-014-3028-1 (2014).
- 430 36 Vanderplanck, M. *et al.* The importance of pollen chemistry in evolutionary host shifts of  
431 bees. *Sci. Rep.* **7**, 43058, doi:10.1038/srep43058 (2017).

- 432 37 Tiedeken, E. J. *et al.* Nectar chemistry modulates the impact of an invasive plant on native  
433 pollinators. *Funct. Ecol.* **30**, 885-893, doi:10.1111/1365-2435.12588 (2016).
- 434 38 Arnold, S. E. J., Peralta Idrovo, M. E., Lomas Arias, L. J., Belmain, S. R. & Stevenson, P.  
435 C. Herbivore defence compounds occur in pollen and reduce bumblebee colony fitness. *J.*  
436 *Chem. Ecol.* **40**, 878-881, doi:10.1007/s10886-014-0467-4 (2014).
- 437 39 Mandelik, Y., Winfree, R., Neeson, T. & Kremen, C. Complementary habitat use by wild  
438 bees in agro-natural landscapes. *Ecol. Appl.* **22**, 1535-1546, doi:10.1890/11-1299.1 (2012).
- 439 40 Moron, D. *et al.* Wild pollinator communities are negatively affected by invasion of alien  
440 goldenrods in grassland landscapes. *Biol. Conserv.* **142**, 1322-1332,  
441 doi:10.1016/j.biocon.2008.12.036 (2009).
- 442 41 Nienhuis, C. M., Dietzsch, A. C. & Stout, J. C. The impacts of an invasive alien plant and its  
443 removal on native bees. *Apidologie.* **40**, 450-463, doi:10.1051/apido/2009005 (2009).
- 444 42 Lopezaraiza-Mikel, M. E., Hayes, R. B., Whalley, M. R. & Memmott, J. The impact of an  
445 alien plant on a native plant-pollinator network: an experimental approach. *Ecol. Lett.* **10**,  
446 539-550, doi:10.1111/j.1461-0248.2007.01055.x (2007).
- 447 43 Schweiger, O. *et al.* Multiple stressors on biotic interactions: how climate change and alien  
448 species interact to affect pollination. *Biological Reviews.* **85**, 777-795, doi:10.1111/j.1469-  
449 185X.2010.00125.x (2010).
- 450 44 Ramos-Jiliberto, R., Valdovinos, F. S., Moisset de Espanés, P. & Flores, J. D. Topological  
451 plasticity increases robustness of mutualistic networks. *J. Anim. Ecol.* **81**, 896-904,  
452 doi:10.1111/j.1365-2656.2012.01960.x (2012).
- 453 45 CaraDonna, P. J. *et al.* Interaction rewiring and the rapid turnover of plant-pollinator  
454 networks. *Ecol. Lett.* **20**, 385-394, doi:10.1111/ele.12740 (2017).
- 455 46 Montero- Castaño, A. & Vilà, M. Influence of the honeybee and trait similarity on the effect  
456 of a non-native plant on pollination and network rewiring. *Funct. Ecol.* **31**, 142-152,  
457 doi:10.1111/1365-2435.12712 (2017).

- 458 47 Bartomeus, I., Vilà, M. & Santamaria, L. Contrasting effects of invasive plants in plant-  
459 pollinator networks. *Oecologia* **155**, 761-770, doi:10.1007/s00442-007-0946-1 (2008).
- 460 48 Kaiser-Bunbury, C. N. *et al.* Ecosystem restoration strengthens pollination network  
461 resilience and function. *Nature*. **542**, 223-227, doi:10.1038/nature21071 (2017).
- 462 49 Aizen, M. A., Morales, C. L. & Morales, J. M. Invasive mutualists erode native pollination  
463 webs. *PLoS Biol.* **6**, e31, doi:10.1371/journal.pbio.0060031 (2008).
- 464 50 Vanbergen, A. J., Woodcock, B. A., Heard, M. S. & Chapman, D. S. Network size, structure  
465 and mutualism dependence affect the propensity for plant–pollinator extinction cascades.  
466 *Funct. Ecol.* **31**, 1285-1293, doi:10.1111/1365-2435.12823 (2017).
- 467 51 Aizen, M. A. *et al.* The phylogenetic structure of plant–pollinator networks increases with  
468 habitat size and isolation. *Ecol. Lett.* **19**, 29-36, doi:10.1111/ele.12539 (2016).
- 469 52 McKinney, A. M. & Goodell, K. Plant-pollinator interactions between an invasive and  
470 native plant vary between sites with different flowering phenology. *Plant Ecol.* **212**, 1025-  
471 1035, doi:10.1007/s11258-010-9882-y (2011).
- 472 53 Albrecht, M., Ramis, M. R. & Traveset, A. Pollinator-mediated impacts of alien invasive  
473 plants on the pollination of native plants: the role of spatial scale and distinct behaviour  
474 among pollinator guilds. *Biol. Invasions* **18**, 1801-1812, doi:10.1007/s10530-016-1121-6  
475 (2016).
- 476 54 Dietzsch, A., Stanley, D. & Stout, J. Relative abundance of an invasive alien plant affects  
477 native pollination processes. *Oecologia* **167**, 469-479, doi:10.1007/s00442-011-1987-z  
478 (2011).
- 479 55 Thijs, K., Brys, R., Verboven, H. F. & Hermy, M. The influence of an invasive plant species  
480 on the pollination success and reproductive output of three riparian plant species. *Biol*  
481 *Invasions* **14**, 355-365, doi:10.1007/s10530-011-0067-y (2012).

- 482 56 Morales, C. L. & Traveset, A. A meta-analysis of impacts of alien vs. native plants on  
483 pollinator visitation and reproductive success of co-flowering native plants. *Ecol. Lett.* **12**,  
484 716-728, doi:10.1111/j.1461-0248.2009.01319.x (2009).
- 485 57 Brown, B. J., Mitchell, R. J. & Graham, S. A. Competition for pollination between an  
486 invasive species (purple loosestrife) and a native congener. *Ecology.* **83**, 2328-2336,  
487 doi:10.1890/0012-9658(2002)083[2328:cfpbai]2.0.co;2 (2002).
- 488 58 Montero- Castaño, A. & Vilà, M. Impact of landscape alteration and invasions on  
489 pollinators: a meta-analysis. *J. Ecol.* **100**, 884-893, doi:10.1111/j.1365-2745.2012.01968.x  
490 (2012).
- 491 59 Carvalheiro, L. G. *et al.* The potential for indirect effects between co-flowering plants via  
492 shared pollinators depends on resource abundance, accessibility and relatedness. *Ecol. Lett.*  
493 **17**, 1389-1399, doi:10.1111/ele.12342 (2014).
- 494 60 Russo, L., Nichol, C. & Shea, K. Pollinator floral provisioning by a plant invader:  
495 quantifying beneficial effects of detrimental species. *Divers. Distrib.* **22**, 189-198,  
496 doi:10.1111/ddi.12397 (2016).
- 497 61 Bruckman, D. & Campbell, D. R. Pollination of a native plant changes with distance and  
498 density of invasive plants in a simulated biological invasion. *Am. J. Bot.* **103**, 1458-1465,  
499 doi:10.3732/ajb.1600153 (2016).
- 500 62 Morales, C. L. & Traveset, A. Interspecific pollen transfer: Magnitude, prevalence and  
501 consequences for plant fitness. *Crit. Rev. Plant Sci.* **27**, 221-238,  
502 doi:10.1080/07352680802205631 (2008).
- 503 63 Chittka, L. & Schurkens, S. Successful invasion of a floral market - An exotic Asian plant  
504 has moved in on Europe's river-banks by bribing pollinators. *Nature.* **411**, 653-653 (2001).
- 505 64 Bruckman, D. & Campbell, D. R. Timing of invasive pollen deposition influences pollen  
506 tube growth and seed set in a native plant. *Biol. Invasions.* **18**, 1701-1711,  
507 doi:10.1007/s10530-016-1113-6 (2016).

- 508 65 Emer, C., Vaughan, I. P., Hiscock, S. & Memmott, J. The impact of the invasive alien plant,  
509 *Impatiens glandulifera*, on pollen transfer networks. *PLoS ONE*. **10**, e0143532,  
510 doi:10.1371/journal.pone.0143532 (2015).
- 511 66 Morales, C. L., Arbetman, M. P., Cameron, S. A. & Aizen, M. A. Rapid ecological  
512 replacement of a native bumble bee by invasive species. *Front. Ecol. Environ.* **11**, 529-534,  
513 doi:10.1890/120321 (2013).
- 514 67 Moritz, R. F. A., Hartel, S. & Neumann, P. Global invasions of the western honeybee (*Apis*  
515 *mellifera*) and the consequences for biodiversity. *Ecoscience*. **12**, 289-301 (2005).
- 516 68 Goulson, D. Effects of introduced bees on native ecosystems. *Annu. Rev. Ecol. Evol. Syst.*  
517 **34**, 1-26, doi:10.1146/annurev.ecolsys.34.011802.132355 (2003).
- 518 69 Dohzono, I. & Yokoyama, J. Impacts of alien bees on native plant-pollinator relationships: a  
519 review with special emphasis on plant reproduction. *Appl. Entomol. Zool.* **45**, 37-47,  
520 doi:10.1303/aez.2010.37 (2010).
- 521 70 Dafni, A., Kevan, P., Gross, C. L. & Goka, K. *Bombus terrestris*, pollinator, invasive and  
522 pest: an assessment of problems associated with its widespread introductions for commercial  
523 purposes. *Appl. Entomol. Zool.* **45**, 101-113, doi:10.1303/aez.2010.101 (2010).
- 524 71 Ings, T. C., Ward, N. L. & Chittka, L. Can commercially imported bumble bees out-compete  
525 their native conspecifics? *J. Appl. Ecol.* **43**, 940-948, doi:10.1111/j.1365-2664.2006.01199.x  
526 (2006).
- 527 72 Hanna, C., Foote, D. & Kremen, C. Competitive impacts of an invasive nectar thief on  
528 plant-pollinator mutualisms. *Ecology*. **95**, 1622-1632, doi:10.1890/13-1276.1 (2014).
- 529 73 Thomson, D. Competitive interactions between the invasive European honey bee and native  
530 bumble bees. *Ecology* **85**, 458-470 (2004).
- 531 74 Roubik, D. W. & Wolda, H. Do competing honey bees matter? Dynamics and abundance of  
532 native bees before and after honey bee invasion. *Popul. Ecol.* **43**, 53-62,  
533 doi:10.1007/pl00012016 (2001).

- 534 75 Yang, G. Harm of introducing the western honeybee *Apis mellifera* L. to the Chinese  
535 honeybee *Apis cerana* F. and its ecological impact. *Acta Entomology Sinica*. **3**, 015 (2005).
- 536 76 Paini, D. R. Impact of the introduced honey bee (*Apis mellifera*) (Hymenoptera: Apidae) on  
537 native bees: a review. *Austral Ecology*. **29**, 399-407, doi:10.1111/j.1442-9993.2004.01376.x  
538 (2004).
- 539 77 Hanna, C., Foote, D. & Kremen, C. Invasive species management restores a plant–pollinator  
540 mutualism in Hawaii. *J. Appl. Ecol.* **50**, 147-155, doi:10.1111/1365-2664.12027 (2013).
- 541 78 Montero-Castaño, A. & Vilà, M. Influence of the honeybee and trait similarity on the effect  
542 of a non-native plant on pollination and network rewiring. *Funct. Ecol.* **31**, 142-152,  
543 doi:10.1111/1365-2435.12712 (2017).
- 544 79 Kato, M. & Kawakita, A. Plant-pollinator interactions in New Caledonia influenced by  
545 introduced honey bees. *Am. J. Bot.* **91**, 1814-1827, doi:10.3732/ajb.91.11.1814 (2004).
- 546 80 Remnant, E. J. *et al.* Reproductive interference between honeybee species in artificial  
547 sympatry. *Mol. Ecol.* **23**, 1096-1107, doi:10.1111/mec.12669 (2014).
- 548 81 Kondo, N. I. *et al.* Reproductive disturbance of Japanese bumblebees by the introduced  
549 European bumblebee *Bombus terrestris*. *Naturwissenschaften*. **96**, 467-475,  
550 doi:10.1007/s00114-008-0495-4 (2009).
- 551 82 Brito, R. M., O. Francisco, F., Ho, S. Y. W. & Oldroyd, B. P. Genetic architecture of the  
552 *Tetragonula carbonaria* species complex of Australian stingless bees (Hymenoptera: Apidae:  
553 Meliponini). *Biol. J. Linn. Soc.* **113**, 149-161, doi:10.1111/bij.12292 (2014).
- 554 83 Byatt, M. A., Chapman, N. C., Latty, T. & Oldroyd, B. P. The genetic consequences of the  
555 anthropogenic movement of social bees. *Insectes Soc.* **63**, 15-24, doi:10.1007/s00040-015-  
556 0441-3 (2016).
- 557 84 Aizen, M. A. & Feinsinger, P. Habitat fragmentation, native insect pollinators, and feral  
558 honey bees in Argentine chaco serrano *Ecol. Appl.* **4**, 378-392, doi:10.2307/1941941 (1994).

- 559 85 Dick, C. W. Genetic rescue of remnant tropical trees by an alien pollinator. *Proc. Roy. Soc. B.* **268**, 2391-2396, doi:10.1098/rspb.2001.1781 (2001).
- 560
- 561 86 Brittain, C., Williams, N., Kremen, C. & Klein, A.-M. Synergistic effects of non-*Apis* bees  
562 and honey bees for pollination services. *Proc. Roy. Soc. B.* **280**, doi:10.1098/rspb.2012.2767  
563 (2013).
- 564 87 Greenleaf, S. S. & Kremen, C. Wild bees enhance honey bees' pollination of hybrid  
565 sunflower. *PNAS.* **103**, 13890-13895, doi:10.1073/pnas.0600929103 (2006).
- 566 88 Aizen, M. A. *et al.* When mutualism goes bad: density-dependent impacts of introduced  
567 bees on plant reproduction. *New Phytol.* **204**, 322-328, doi:10.1111/nph.12924 (2014).
- 568 89 Sáez, A., Morales, C. L., Ramos, L. Y. & Aizen, M. A. Extremely frequent bee visits  
569 increase pollen deposition but reduce drupelet set in raspberry. *J. Appl. Ecol.* **51**, 1603-1612,  
570 doi:10.1111/1365-2664.12325 (2014).
- 571 90 Kenta, T., Inari, N., Nagamitsu, T., Goka, K. & Hiura, T. Commercialized European  
572 bumblebee can cause pollination disturbance: an experiment on seven native plant species in  
573 Japan. *Biol. Cons.* **134**, 298-309, doi:10.1016/j.biocon.2006.07.023 (2007).
- 574 91 Arbetman, M. P., Meeus, I., Morales, C. L., Aizen, M. A. & Smagghe, G. Alien parasite  
575 hitchhikes to Patagonia on invasive bumblebee. *Biol. Invasions.* **15**, 489-494,  
576 doi:10.1007/s10530-012-0311-0 (2013).
- 577 92 Schmid-Hempel, R. *et al.* The invasion of southern South America by imported bumblebees  
578 and associated parasites. *J. Anim. Ecol.* **83**, 823-837, doi:10.1111/1365-2656.12185 (2014).
- 579 93 Ji, R., Xie, B., Yang, G. & Li, D. From introduced species to invasive species--a case study  
580 on the Italian bee *Apis mellifera* L. *Chinese Journal of Ecology.* **22**, 70-73 (2002).
- 581 94 Martin, S. J. *et al.* Global honey bee viral landscape altered by a parasitic mite. *Science.* **336**,  
582 1304-1306, doi:10.1126/science.1220941 (2012).
- 583 95 Wilfert, L. *et al.* Deformed wing virus is a recent global epidemic in honeybees driven by  
584 *Varroa* mites. *Science.* **351**, 594-597, doi:10.1126/science.aac9976 (2016).

- 585 96 Brosi, B. J., Delaplane, K. S., Boots, M. & de Roode, J. C. Ecological and evolutionary  
586 approaches to managing honeybee disease. *Nature Ecology & Evolution* **1**, 1250-1262,  
587 doi:10.1038/s41559-017-0246-z (2017).
- 588 97 McMahon, D. P. *et al.* Elevated virulence of an emerging viral genotype as a driver of  
589 honeybee loss. *Proc. Roy. Soc. B.* **283**, doi:10.1098/rspb.2016.0811 (2016).
- 590 98 Ryabov, E. V. *et al.* A virulent strain of deformed wing virus (DWV) of honeybees (*Apis*  
591 *mellifera*) prevails after *Varroa destructor*-mediated, or *in vitro*, transmission. *PLoS Pathog.*  
592 **10**, e1004230, doi:10.1371/journal.ppat.1004230 (2014).
- 593 99 Furst, M. A., McMahon, D. P., Osborne, J. L., Paxton, R. J. & Brown, M. J. F. Disease  
594 associations between honeybees and bumblebees as a threat to wild pollinators. *Nature.* **506**,  
595 364-366, doi:10.1038/nature12977 (2014).
- 596 100 McMahon, D. P. *et al.* A sting in the spit: widespread cross-infection of multiple RNA  
597 viruses across wild and managed bees. *J. Anim. Ecol.* **84**, 615-624, doi:10.1111/1365-  
598 2656.12345 (2015).
- 599 101 Singh, R. *et al.* RNA viruses in hymenopteran pollinators: evidence of inter-taxa virus  
600 transmission via pollen and potential impact on non-*Apis* hymenopteran species. *PLoS ONE.*  
601 **5**, e14357, doi:10.1371/journal.pone.0014357 (2010).
- 602 102 Monceau, K., Bonnard, O. & Thiery, D. *Vespa velutina*: a new invasive predator of  
603 honeybees in Europe. *J. Pest Sci.* **87**, 1-16, doi:10.1007/s10340-013-0537-3 (2014).
- 604 103 Keeling, M. J., Franklin, D. N., Datta, S., Brown, M. A. & Budge, G. E. Predicting the  
605 spread of the Asian hornet (*Vespa velutina*) following its incursion into Great Britain.  
606 *Scientific Reports.* **7**, 6240, doi:10.1038/s41598-017-06212-0 (2017).
- 607 104 Hansen, D. M. & Müller, C. B. Invasive ants disrupt gecko pollination and seed dispersal of  
608 the endangered plant *Roussea simplex* in Mauritius. *Biotropica.* **41**, 202-208,  
609 doi:10.1111/j.1744-7429.2008.00473.x (2009).

610 105 Lach, L. A mutualism with a native membracid facilitates pollinator displacement by  
611 Argentine ants. *Ecology*. **88**, 1994-2004, doi:10.1890/06-1767.1 (2007).

612 106 Hanna, C. *et al.* Floral visitation by the Argentine ant reduces bee visitation and plant seed  
613 set. *Ecology*. **96**, 222-230, doi:10.1890/14-0542.1 (2015).

614 107 Abe, T., Wada, K., Kato, Y., Makino, S. & Okochi, I. Alien pollinator promotes invasive  
615 mutualism in an insular pollination system. *Biol. Invasions*. **13**, 957-967,  
616 doi:10.1007/s10530-010-9882-9 (2011).

617 108 Vazquez, D. P. & Simberloff, D. Changes in interaction biodiversity induced by an  
618 introduced ungulate. *Ecol. Lett.* **6**, 1077-1083 (2003).

619 109 Vazquez, D. P. & Simberloff, D. Indirect effects of an introduced ungulate on pollination  
620 and plant reproduction. *Ecol. Monogr.* **74**, 281-308 (2004).

621 110 Stokl, J., Brodmann, J., Dafni, A., Ayasse, M. & Hansson, B. S. Smells like aphids: orchid  
622 flowers mimic aphid alarm pheromones to attract hoverflies for pollination. *Proc. Roy. Soc.*  
623 *B.* **278**, 1216-1222, doi:10.1098/rspb.2010.1770 (2011).

624 111 Kessler, A., Halitschke, R. & Poveda, K. Herbivory-mediated pollinator limitation: negative  
625 impacts of induced volatiles on plant–pollinator interactions. *Ecology*. **92**, 1769-1780,  
626 doi:10.1890/10-1945.1 (2011).

627 112 Barber, N. A., Adler, L. S., Theis, N., Hazzard, R. V. & Kiers, E. T. Herbivory reduces plant  
628 interactions with above- and belowground antagonists and mutualists. *Ecology*. **93**, 1560-  
629 1570, doi:10.1890/11-1691.1 (2012).

630 113 Desurmont, G. A. *et al.* Alien interference: disruption of infochemical networks by invasive  
631 insect herbivores. *Plant, Cell Environ.* **37**, 1854-1865, doi:10.1111/pce.12333 (2014).

632 114 Vieira, M. C. & Almeida-Neto, M. A simple stochastic model for complex coextinctions in  
633 mutualistic networks: robustness decreases with connectance. *Ecol. Lett.* **18**, 144-152,  
634 doi:10.1111/ele.12394 (2015).

635 115 Valdovinos, F. S. *et al.* Niche partitioning due to adaptive foraging reverses effects of  
636 nestedness and connectance on pollination network stability. *Ecol. Lett.* **19**, 1277-1286,  
637 doi:10.1111/ele.12664 (2016).

638 116 Sauve, A. M. C., Fontaine, C. & Thébault, E. Structure–stability relationships in networks  
639 combining mutualistic and antagonistic interactions. *Oikos* **123**, 378-384,  
640 doi:10.1111/j.1600-0706.2013.00743.x (2014).

641 117 Carvalho, G. O., Medel, R. & Navarro, L. Assessing the effects of native plants on the  
642 pollination of an exotic herb, the blueweed *Echium vulgare* (Boraginaceae). *Arthropod-  
643 Plant Interactions.* **7**, 475-484, doi:10.1007/s11829-013-9265-0 (2013).

644 118 Guimaraes, P. R., Jr., Jordano, P. & Thompson, J. N. Evolution and coevolution in  
645 mutualistic networks. *Ecol. Lett.* **14**, 877-885, doi:10.1111/j.1461-0248.2011.01649.x  
646 (2011).

647 119 Gloag, R. *et al.* An invasive social insect overcomes genetic load at the sex locus. *Nature  
648 Ecology & Evolution* **1**, 0011, doi:10.1038/s41559-016-0011 (2016).

649 120 Arbetman, M. P., Gleiser, G., Morales, C. L., Williams, P. & Aizen, M. A. Global decline of  
650 bumblebees is phylogenetically structured and inversely related to species range size and  
651 pathogen incidence. *Proc. Roy. Soc. B.* **284**, doi:10.1098/rspb.2017.0204 (2017).

652 121 Santini, A. *et al.* Biogeographical patterns and determinants of invasion by forest pathogens  
653 in Europe. *New Phytol.* **197**, 238-250, doi:10.1111/j.1469-8137.2012.04364.x (2013).

654 122 Shykoff, J. A. & Bucheli, E. Pollinator visitation patterns, floral rewards and the probability  
655 of transmission of *Microbotryum violaceum*, a venereal disease of plants. *J. Ecol.* **83**, 189-  
656 198, doi:10.2307/2261557 (1995).

657 123 Roy, B. A. The effects of pathogen-induced pseudoflowers and buttercups on each others  
658 visitation. *Ecology.* **75**, 352-358, doi:10.2307/1939539 (1994).

659 124 Li, J. L. *et al.* Systemic spread and propagation of a plant-pathogenic virus in European  
660 honeybees, *Apis mellifera*. *Mbio.* **5**, doi:10.1128/mBio.00898-13 (2014).

661 125 Kovács-Hostyánszki, A. *et al.* Ecological intensification to mitigate impacts of conventional  
662 intensive land use on pollinators and pollination. *Ecol. Lett.* **20**, 673-689,  
663 doi:10.1111/ele.12762 (2017).

664 126 Brown, M. J. F. *et al.* A horizon scan of future threats and opportunities for pollinators and  
665 pollination. *PeerJ.* **4**, e2249, doi:10.7717/peerj.2249 (2016).

666 127 Roy, H. E. *et al.* Horizon scanning for invasive alien species with the potential to threaten  
667 biodiversity in Great Britain. *Global Change Biology.* **20**, 3859-3871,  
668 doi:10.1111/gcb.12603 (2014).

669 128 Keeling, M. J. *et al.* Efficient use of sentinel sites: detection of invasive honeybee pests and  
670 diseases in the UK. *J R Soc Interface.* **14**, doi:10.1098/rsif.2016.0908 (2017).

671 129 Prior, K. M., Robinson, J. M., Meadley Dunphy, S. A. & Frederickson, M. E. Mutualism  
672 between co-introduced species facilitates invasion and alters plant community structure.  
673 *Proc. Roy. Soc. B.* **282**, doi:10.1098/rspb.2014.2846 (2015).

674 130 Sakai, A. K. *et al.* The population biology of invasive species. *Annu. Rev. Ecol. Syst.* **32**,  
675 305-332 (2001).

676 131 Blackburn, T. M., Lockwood, J. L. & Cassey, P. The influence of numbers on invasion  
677 success. *Mol. Ecol.* **24**, 1942-1953, doi:10.1111/mec.13075 (2015).

678 132 Willi, Y., Van Buskirk, J. & Hoffmann, A. A. Limits to the adaptive potential of small  
679 populations. *Annu. Rev. Ecol. Evol. Syst.* **37**, 433-458 (2006).

680 133 Gomulkiewicz, R., Nuismer, S. L. & Thompson, J. N. Coevolution in variable mutualisms.  
681 *Am. Nat.* **162**, S80-S93 (2003).

682 134 Stotz, G. C., Gianoli, E. & Cahill, J. F. Spatial pattern of invasion and the evolutionary  
683 responses of native plant species. *Evol. Appl.* **9**, 939-951, doi:10.1111/eva.12398 (2016).

684

685 **Box 1. Evolutionary perspectives on impacts of invasive alien species on pollinators.** Evolution is driven by  
686 four processes: mutation, gene flow, drift and selection. Anthropogenic changes to a pollination system that  
687 modify these processes have the capacity to affect the evolutionary outcomes for species, co-evolutionary  
688 dynamics and community structure and function. Several eco-evolutionary characteristics of the interacting  
689 communities can modulate these evolutionary processes, such as the extent of mutual dependence between  
690 the interacting species, the probability of encounter, the demography of the invasion, and the phylogenetic  
691 histories of the plants and pollinators<sup>10,129-131</sup>.

692 Species invading a pollination community create and are exposed to novel selective pressures have  
693 the ability to modify ongoing evolutionary trajectories<sup>10-12</sup>. Indeed, the newly interacting species compete for  
694 resources (e.g. floral rewards, pollination service), and asymmetric interactions will allow some to dominate  
695 the community. This is one of the reasons why mathematical models predicted that the widespread  
696 introduction of the super-generalist and very competitive honeybee *A. mellifera* is expected to select for  
697 convergence in flower traits across many wild plant species, affecting plant-pollinator community function  
698 and structure in the longer term<sup>118</sup>. The relative changes of both the census and effective population sizes of  
699 the invasive and native species<sup>131</sup> can also have a direct impact on the evolutionary paths of the interacting  
700 species. Because invasive species usually reach large population sizes, they can affect the populations of co-  
701 occurring natives negatively through either interference or exploitative competition. Ultimately, this can, on  
702 the one hand, decrease the native population's chances of demographic recovery, and on the other hand,  
703 reduce the native's effective population size increasing the effects of genetic drift. Likewise, the effects of  
704 genetic drift are also expected to be amplified in species that already have low effective population sizes,  
705 such as is usually the case in endangered or rare species<sup>132</sup>. Further, organisms with small effective  
706 population sizes are less responsive to selection, which negatively affects the ability of natives to adapt to the  
707 new conditions created by the arrival and establishment of the invasive species.

708 Through its effect on the population sizes of co-occurring native species, invasive species can also  
709 affect connectivity among native populations. Loss of connectivity decreases gene flow and in some cases  
710 genetic diversity and evolvability, rendering native species less able to adapt to new conditions or to recover  
711 from the effects of drift<sup>132-134</sup>. Impoverished genetic diversity may affect adaptive processes contributing to  
712 the success or failure of invasions, depending of the type of interaction the native has with the invasive  
713 species. On this point, modelling approaches indicated that an alien species with high genetic diversity

714 (usually associated with a higher ability to adapt) is expected to establish in the community. Further, higher  
715 genetic diversity in the resident (native) species than in the invasive species can lead to exclusion of the  
716 invasive in predator-prey interactions, and may allow adaptation to the invasive and survival of both species  
717 in other types of interactions (e.g., mutualistic, competition)<sup>10</sup>.  
718

719 **Figure 1. Conceptual synthesis of the direct and indirect impacts on (A) native pollinators and (B)**  
720 **native plant pollination from invasive alien species of (C) plants, (D) predators, (E) introduced**  
721 **pollinators and their (F) pests and pathogens.** Images are representative examples of native and invasive  
722 alien species and do not portray a particular ecological system: (A) native Patagonian giant bumblebee  
723 *Bombus dahlbomii* (source Carolina Morales); (B) native British wildflowers (source Claire Carvell); (C)  
724 Himalayan balsam *Impatiens glandulifera* invasive in Europe (source Dan Chapman); (D) Asian hornet  
725 *Vespa velutina* invasive in Europe (source Gilles San Martin) (E) managed pollinators translocated  
726 worldwide include the western honeybee *Apis mellifera* (source Eugene Ryabov) and *Bombus terrestris*  
727 (source Adam Vanbergen), which has spread (F) pests and pathogens e.g. *Varroa* mite (source USDA);  
728 Deformed Wing Virus (source Pavel Plevka).

729

730 **Figure 2. Invasive alien plant impact on pollinator visitation and network structure.** An example of an  
731 alien plant species (A) Himalayan balsam, *Impatiens glandulifera* native to Asia and invasive in Europe.  
732 This plant attains high densities, produces copious nectar and pollen and possesses a large, enduring floral  
733 display, all of which enables it to readily penetrate and dominate plant-pollinator networks by co-opting  
734 pollinators, such as (B) the honeybee and (C) syrphid hoverflies. In turn, alien plant invasions can alter the  
735 composition and structure of native plant-pollinator networks from (D) to (E). This raises the risk of (E)  
736 pollinator nutritional deficits due to reductions in availability of essential nutrients from diverse floral  
737 resources, poorly matched mutualisms and impaired native plant pollination (but see<sup>59</sup> for an exception).  
738 Source of images: Dan Chapman, Claire Carvell and Adam Vanbergen.

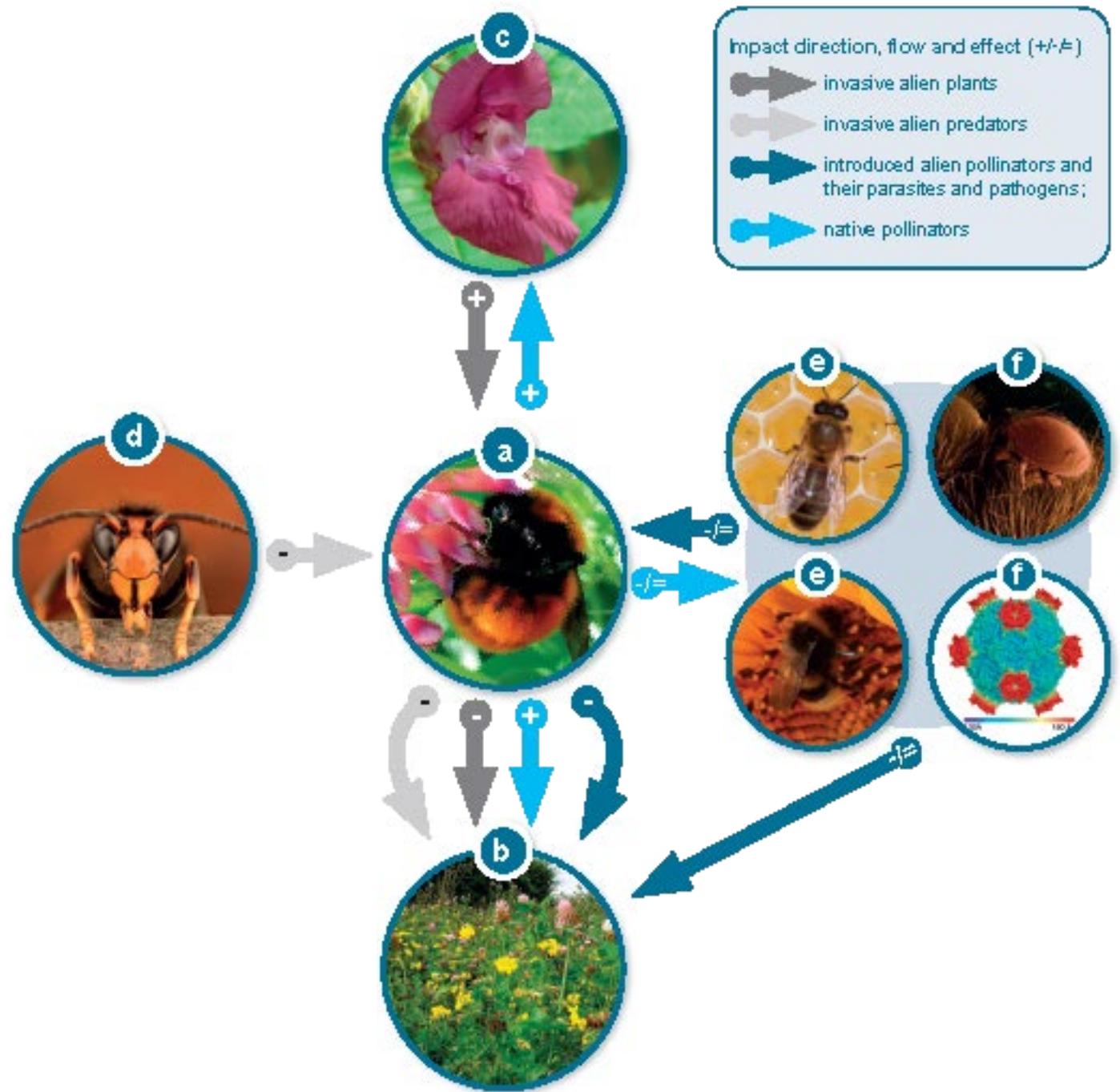
739

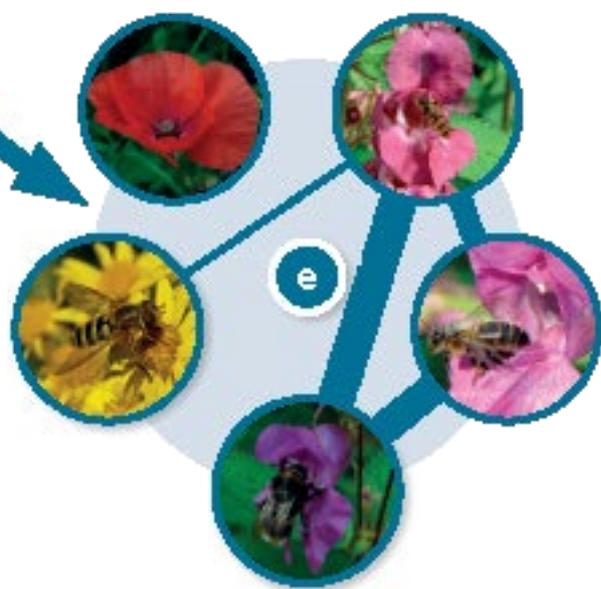
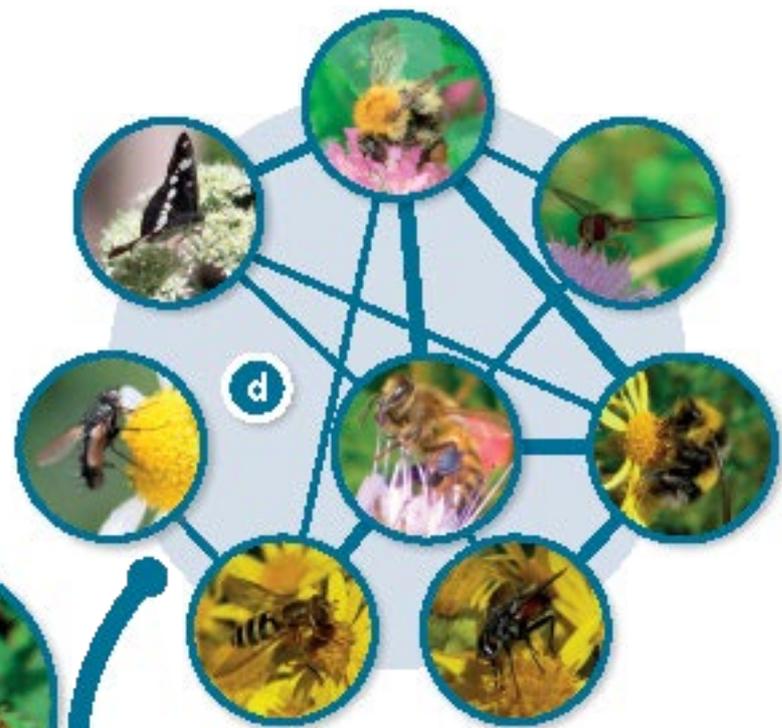
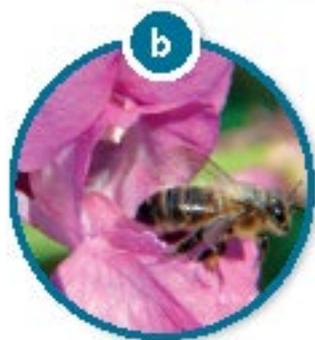
740 **Figure 3. Global movement of managed pollinators and risk of altered host-vector-pathogen dynamics.**  
741 The historic and current human-assisted translocation of (A) the western honey bee *Apis mellifera* for  
742 apiculture and pollination services led to its range extending from its native range (vertical lines) to a near  
743 global distribution (shaded green area) that overlapped with other *Apis* species including the Asian honey bee  
744 *A. cerana* (horizontal lines). This led to (B) the *Varroa* mite, a parasite of *A. cerana*, infecting sympatric  
745 colonies of *A. mellifera* and subsequently spreading worldwide in association with the new host bee. *Varroa*  
746 is now the major worldwide pest of managed honeybees between which it transmits many viruses<sup>2,13</sup>. Recent

747 evidence suggests that (C) the novel eco-evolutionary interaction between *Varroa*, *A. mellifera* and the  
748 Deformed Wing Virus (DWV) has increased viral virulence and that DWV (D) co-infects bumblebee species  
749 with (E) unknown implications for pollinator community epidemiology. Image sources: *Apis mellifera*  
750 (Eugene Ryabov); *Apis cerana* (Dino Martins); *Varroa* mite (USDA); Deformed Wing Virus (Pavel Plevka).

751

752 **Figure 4. Complex interactions between alien predators, alien and native pollinators and native plants**  
753 **transform and maintain pollination in highly modified ecosystems.** Within the Hawaiian archipelago  
754 (map outline), historic introductions of (A) mammalian predators (e.g. cats and rats) led to (B) extinctions  
755 and declines of birds, particularly of the charismatic Hawaiian honeycreepers, that (C) pollinated the tree  
756 *Metrosideros polymorpha* among many other native plant species. More recently, the invasion by (D)  
757 *Vespula pensylvanica* the predatory wasp and nectar thief has increased competition for floral resources,  
758 deterred flower visitation by (E) native *Hylaeus* bees and the (F) alien honeybee *A. mellifera* and thereby (C)  
759 reduced *M. polymorpha* pollination and fruit production. Experimental exclusion of the wasp showed the  
760 alien honeybee (F) is now the most effective pollinator in this system with the decline or loss of bird  
761 pollinators. Double-headed arrows indicate mutualisms. Single headed arrows show impacts. Grey arrows =  
762 alien interactions; Blue arrows = native interactions. Dashed arrow = declining or extinct interactions. Image  
763 sources: *V. pensylvanica* (J. Gallacher CC-BY-2.0); *Hylaeus* spp. Forrest & Kim Starr; feral cat (Batty CC-  
764 BY-2.0); rat (US-NPS).





Altered network structure and stability  
Reduced availability of balanced pollinator diets  
Greater interspecific competition  
Increased pollination interference

