

Working title, The impact of heterospecific pollen on plant reproductive success is mediated by phylogenetic distance and floral reproductive traits

Jose B. Lanuza, Ignasi Bartomeus, Tia-Lynn Ashman, Romina Rader * ^{1,2,3}

¹US Environmental Protection Agency, Office of Research and Development, National Health and Environmental Effects Research Laboratory, Atlantic Ecology Division, 27 Tarzwell Drive Narragansett, RI, 02882, USA

²Big Name University, Department of R, City, BN, 01020, USA

³Estacion Biologica de Donana (EBD-CSIC), E-41092 Sevilla, Spain

* corresponding author: barragansljose@gmail.com

Pollinator sharing can have negative consequences for species fitness with the arrival of foreign pollen. However, the costs of heterospecific pollen are not yet well understood. For this reason, we have conducted a glasshouse experiment where we try to understand how phylogenetic relatedness and the different traits of these species are involved in this process. We experimentally crossed 10 species belonging to three different families: Brassicaceae, Solanaceae and Convolvulaceae. Overall, more than 4000 crosses were done and seed set and pollen tubes were considered as proxy of effect. We found that for all species foreign pollen (50% or less) reduced seed set. Moreover, the seed set reduction is not dependent on the degree of relatedness of the pollen donor. However, the effect is governed by the degree of relatedness and the traits of the species recipient. Our results show that the outcome of heterospecific pollen deposition is determined in greater degree by the traits of the pollen recipient than the pollen donor and that certain traits such as compatibility system are crucial to understand the costs of heterospecific pollen.

Keywords: heterospecific pollen, plant reproduction, fitness, interspecific competition, phylogenetic distance.

INTRODUCTION

Paragraph 1

In natural systems plant species normally coexist and share their floral visitors with other species Bascompte et al. (2003). This pollinator sharing from the plant perspective at the pre-pollination stage can be negative due to competition Pauw (2013) or positive due to facilitation Carvalheiro et al. (2014). Once the floral visitor has arrived to the flower, pollen deposition on the stigma can take place and

34 hence ovule fertilization. An increasing number of visits generally correlates with higher chances of
35 fertilization Engel and Irwin (2003). However this is not always the case, among these possible flower
36 visitors we find also nectar robbers and pollen thieves Inouye (1980) and the quality of pollen that is
37 deposited on the stigma is also highly relevant to the pollination succes Aizen and Harder (2007).
38 Moreover, other less study issues in the pollination process are conspecific pollen loss and the arrival of
39 foreign pollen which can have important detrimental effects for the species fitness Morales and Traveset
40 (2008) Ashman and Arceo-Gómez (2013).

41 **Paragraph 2**

42 Recent studies have advanced in the ecological importance of heterospecific pollen effect Morales and
43 Traveset (2008) (???) Arceo-Gómez and Ashman (2016). A general overview of heterospecific pollen
44 effect is that it can play an important role on species fitness but seems to be context dependent and not
45 always fitness decreased is produced Morales and Traveset (2008). There are several traits that are
46 postulated to be highly relevant to this process but the correlation with heterospecific pollen effect
47 remains to be tested. Moreover, the different methods used make difficult to compare among different
48 experiments. For instance many articles do not describe the foreign pollen quantity used to test for
49 effect or how this pollen was applied (Refs). The small size of pollen, great environmental variability
50 and population outcrossing rates among others make of this field of study a daunting task and highly
51 unpredictable.

52 **Paragraph 3**

53 can reduce species fitness (REFS) but seems to be highly contex-dependent. There are hypothesized
54 that some traits can play a crucial role in this species interaction such as stigma type, pollen size,

55 Mention invasive species in this paragraph

56 Few studies have tried to understand how relatedness is involved in the hp effect but generally

57 Until our knowledge

58 Rescue from here the useful things:

59 Invasive species are supposed to have greater negative effect than native ones Arceo-Gómez and
60 Ashman (2016). Although when non-natives species don't have greater negative effect we still don't
61 know why. For this reason, this ecological question is non a native non native one is a trait based issue
62 that is still to be solved. Moreover, the quantity of pollen that integrates in the network can be quite
63 variable ranging from low quantities Bartomeus et al. (2008) to intermediate (ref) to high (ref).
64 Moreover, closely related species are supposed to reduce fitness in greater effect but the evidence is
65 scarce and based on independent studies with different methodologies (Arceo-gomez & Ashman 2016)
66 or studies that just check it with a pair of species that are highly related with the aim to understand
67 hybridization costs (refs). There is a need to deepen into how relatedness is involved in the costs of
68 heterospecific pollen effect. Furthermore, following the conceptual trait framework of Ashman and
69 Arceo-Gomez on heterospecific pollen there are good theoretical basis for trait effect. Notwithstanding,
70 non empirical work has tested how really these traits are involved in heterospecific pollen effect.

71 Explain traits. Put examples

72 what is closely related? same genus? Just that right, the rest is far related?

73 I would like to add that the experiments focus on two proxies of effect prezygotic and postzygotic. Why
74 focus on postzygotic? Is the final stage where we can see the effect. Further studies should also study
75 germination rates.

76 **Paragraph 3**

77 Traditionally heterospecific pollen effect has focused its attention on different pollen donors as a main
78 driver of different effect. However in this article we want to emphasize that this is true for the cases
79 that the species are highly close related where pollen recognition can take place (eg hybridization) but
80 not when this pollen is from less closely related species which the main driver of effect is determined by
81 the reproductive biology of the female part of the plant (compatibility system, stigma type, stigma area
82 and number of ovules).

83 **Paragraph 4**

84 Sell well our work: We are the first empirical experiment testing the effect of heterospecific pollen with

85 phylogenetic distance

86 The great difficulty of working with pollen in a coflowering community make the understanding of
87 heterospecific pollen effect a real challenge. For this reason we have created an artificial co-flowering
88 community in a glasshouse to test the effect with all the possible combinations among them. Where we
89 test the following hypothesis: 1) Does heterospecific pollen reduce seed set, if so, 2) Does heterospecific
90 pollen effect depend on the relatedness of the species, 3) Does heterospecific pollen effect depend on any
91 floral trait?

92 Maybe another possible hypothesis to test is the reciprocity of the effect of heterospecific pollen????

93 Use the sterile species as a proof of the mechanical interference. Was a mistake but seems cool proof!!!

94 METHODS

95 comment starts Glasshouse trial • Species selected and why – how you made them co-flower • Give
96 details of sources and planting seeds, growth medium in pots, temperature and light details • Hand
97 crosses and how you did them, how you measured seed set over time. • Analyses of data –
98 standardization, means, matrices etc.

99 • Analyses and technical difficulties: We calculated effect size by subtracting the mean of the cross
100 pollinated seed set by the mean effect of the HP pollen (explain exactly what figures you used to
101 calculate this) – check with liam about potentially using missing values analyses for the species we
102 don't have?

103 Check that the method is working well to prove that your crosses were close to 50% results in SI i.e not
104 all mixes were 50/50% and we have now counted all the pollen to make this a quantitative variable. We
105 also need to factor in the point that we have different total abundances of pollen across our treatments,
106 irrespective of ratios. To what extent are differences in the ratios of pollen applied by hand across
107 different plant families influenced by plant traits such as pollen size, morphology and stigma surface
108 type?

109 Results – may need to include amount of pollen in models as random factor- prefill matrix with missing
110 value analyse for the species you don't have.

111 Question 1: how do different pollination treatments (100% HP, 50% HP, self and cross) impact HP
112 pollen across different plant families? Even with 100% HP one (or more species?) still produced seed
113 set.

114 Result Effect size of Seed set ~ phylogenetic distance relationship We found that the variation ?/ mean
115 effect size of seed set is positively related to phylogenetic distance. This means the more unrelated the
116 species are, the greater the negative impact of heterospecific pollen (give stats effect size i.e. Procrustes,
117 $X = 0.35$; $P = 0.03$)

118 Question 2 : what are the main traits impacting HP impacts? (compatibility system, pollen size,
119 stigma surface, wet/dry stigma, length of style etc.

120 Effect size of seed set ~ floral traits/ reproductive plant traits We found that the three best terms to
121 explain the variation in seed set is pollen/ovule ratio, stigma width and style length (Stats effect size
122 i.e. $X = 0.39$, $P = 0.02$).

123 Need to provide correlation matrix for all traits just for 10 species Show both ways to present this.
124 Which particular traits do you find significant effects for? Show this and give stats. Present plot for
125 each trait and effect size

126 comment finishes

127 The study was conducted in a glasshouse at University of New England (Armidale, Australia) from
128 November 2017 to March 2018. Rooms were temperature controlled depending on the requirements of
129 the species with day and night temperature differences. The species selected (Table 1) belonged to
130 three different families, Solanaceae, Brassicaceae and Convolvulaceae. The criteria of species/family
131 selection was based on close/distant related species (see phylogenetic tree for relatedness fig 1),
132 heterogeneous traits, low structural flower complexity and fast life cycle. For the purpose of the
133 experiment all the species were considered as pollen recipient and as pollen donor (see interaction
134 matrix, fig 2). Species were watered once or twice per day and fertilized weekly (NPK 23: 3.95: 14).

135 Brown and Mitchell 2001 could be a good paper to explain why we pick seed set as a proxy and not

136 fruit set. We cannot see changes on it, losing information with it.

137 **Hand-pollination**

138 Foreign pollen effect was studied through two different treatments, one with 50% conspecific pollen and
139 50% heterospecific pollen and a second one with 100% foreign pollen (N=10). Seed set was the proxy of
140 effect (see Brown and Mitchell 2001, for differences in effect between seed set and fruit set) and “pollen
141 tubes”. Moreover, hand cross pollination, hand self pollination, apomixis (bagged emasculated flowers)
142 and natural selfing were tested (N=10). Flowers were emasculated the day prior anthesis and hand
143 pollinated next day with a toothpick. Hand-pollination was realized with 3-4 gentle touches on the
144 surface of the stigma. The mixes of pollen were performed on an eppendorf based on the pollen counts
145 made with Neubauer chamber (each anther was counted 4 times for 20 different anthers per species).

146 **Evolutionary distance**

147 Two types of evolutionary distances were calculated with MEGA7 two kinds of markers: 1) Internal
148 transcribed spacer (ITS) and 2) ribulose-bisphosphate carboxylase (RBCL)

149 **Traits**

150 Several traits of the ten species were measured. Pollen per anther was counted, number of ovules,
151 stigma width and length and stigmatic area, style width and length, ovary width and length. Moreover
152 stigma type was tested. Self-incompatibility was

153 We used the statistical language R (R Core Team 2018) for all our analyses. These were implemented in
154 dynamic markdown documents using `knitr` (Xie 2014, 2015, 2018) and `rmarkdown` (Allaire et al.
155 2018) packages. All the multilevel models were fitted with `lme4` (Bates et al. 2015).

156 RESULTS

157 DISCUSSION

158 Discussion

- 159 1. What are the implications of the findings?

160 CONCLUSIONS

161 ACKNOWLEDGEMENTS

162 REFERENCES

- 163 Aizen, M. A., and L. D. Harder. 2007. Expanding the limits of the pollen-limitation concept: Effects of
164 pollen quantity and quality. *Ecology* 88:271–281.
- 165 Allaire, J., Y. Xie, J. McPherson, J. Luraschi, K. Ushey, A. Atkins, H. Wickham, J. Cheng, and W.
166 Chang. 2018. Rmarkdown: Dynamic documents for r.
- 167 Arceo-Gómez, G., and T.-L. Ashman. 2016. Invasion status and phylogenetic relatedness predict cost
168 of heterospecific pollen receipt: Implications for native biodiversity decline. *Journal of Ecology*
169 104:1003–1008.
- 170 Ashman, T.-L., and G. Arceo-Gómez. 2013. Toward a predictive understanding of the fitness costs of
171 heterospecific pollen receipt and its importance in co-flowering communities. *American Journal of*
172 *Botany* 100:1061–1070.
- 173 Bartomeus, I., J. Bosch, and M. Vilà. 2008. High invasive pollen transfer, yet low deposition on native

174 stigmas in a carpobrotus-invaded community. *Annals of Botany* 102:417–424.

175 Bascompte, J., P. Jordano, C. J. Melián, and J. M. Olesen. 2003. The nested assembly of plant–animal
176 mutualistic networks. *Proceedings of the National Academy of Sciences* 100:9383–9387.

177 Bates, D., M. Mächler, B. Bolker, and S. Walker. 2015. Fitting linear mixed-effects models using lme4.
178 *Journal of Statistical Software* 67:1–48.

179 Carvalheiro, L. G., J. C. Biesmeijer, G. Benadi, J. Fründ, M. Stang, I. Bartomeus, C. N.
180 Kaiser-Bunbury, M. Baude, S. I. Gomes, V. Merckx, and others. 2014. The potential for indirect effects
181 between co-flowering plants via shared pollinators depends on resource abundance, accessibility and
182 relatedness. *Ecology letters* 17:1389–1399.

183 Engel, E. C., and R. E. Irwin. 2003. Linking pollinator visitation rate and pollen receipt. *American*
184 *Journal of Botany* 90:1612–1618.

185 Inouye, D. W. 1980. The terminology of floral larceny. *Ecology* 61:1251–1253.

186 Morales, C. L., and A. Traveset. 2008. Interspecific pollen transfer: Magnitude, prevalence and
187 consequences for plant fitness. *Critical Reviews in Plant Sciences* 27:221–238.

188 Pauw, A. 2013. Can pollination niches facilitate plant coexistence? *Trends in ecology & evolution*
189 28:30–37.

190 R Core Team. 2018. R: A language and environment for statistical computing. R Foundation for
191 Statistical Computing, Vienna, Austria.

192 Xie, Y. 2014. Knitr: A comprehensive tool for reproducible research in R. *in* V. Stodden, F. Leisch,
193 and R. D. Peng, editors. *Implementing reproducible computational research*. Chapman; Hall/CRC.

194 Xie, Y. 2015. *Dynamic documents with R and knitr*. 2nd editions. Chapman; Hall/CRC, Boca Raton,
195 Florida.

196 Xie, Y. 2018. Knitr: A general-purpose package for dynamic report generation in r.

