working title Compatibility system and stygma size of pollen recipient as main predictors of heterospecific pollen effect

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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.

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

28 INTRODUCTION

- Paragraph 1 General idea to our concept
- 30 In natural systems plant species normally coexist and share their floral visitors with other species
- 31 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

- fertilization Engel and Irwin (2003). However this is not always the case, among these possible flower visitors we find also nectar robbers and pollen thiefs Inouye (1980) and the quality of pollen that is deposit on the stigma is also highly relevant to the pollination succes Aizen and Harder (2007).

 Moreover, other less study issues in the pollination process are conspecific pollen loss and the arrival of foreign pollen which can have important detrimental effects on species fitness Morales and Traveset (2008) Ashman and Arceo-Gómez (2013).
- Paragraph 2 Introducing topic and knowledge gap
- Recent studies have advanced in the ecological understanding of heterospecific pollen effect Morales and 42 Traveset (2008) Ashman and Arceo-Gómez (2013) Arceo-Gómez and Ashman (2016). A general 43 overview of foreign pollen arrival is that it can play an important role on species fitness but seems to be context dependent and not always produce a decrease in fitness Morales and Traveset (2008). Part of this unpredictability is due to the enormous variability of foreing pollen transferred in nature, where levels between 0 and 75 percent are seen, but most commonly values ranges between 0 and 20 percent of the total pollen load Bartomeus et al. (2008) Montgomery and Rathcke (2012) Ashman and Arceo-Gómez (2013) Fang and Huang (2013), being the generalist species the ones that receive greater loads of heterospecific pollen Fang and Huang (2013). Surprisingly, this low ranges of heterospecific pollen have been shown to decrease fitness greatly Thomson et al. (1982). Although heterospecific pollen quantity is fundamental to understand the outcome of the interaction so is the different traits of both pollen donor and recipient. Ashman and Arceo-Gómez (2013) postulated the first predictive framework for traits of heterospecific pollen effect, where different traits such as compatibility system and pollen size among others seems to be crucial to understand foreing pollen effect. Moreover, in Tong and Huang (2016) an assymetric effect was shown in a crossing experiment between 6 species of the genus Pedicularis where the pollen of long styled species was able to grow the full length of the style on short styled species but not viceversa. Despite these recent caveats, we still lack empirical evidence to affirm what are the main traits that drive heterospecific pollen effect for both pollen donor and recipient at seed production level. Interestingly, to comprehend how these traits interact is also crucial to look at the phylogenetic relatedness of the species. There is a considerable amount of literature of crosses between close related species Brown and Mitchell (2001) Arceo-Gómez et al. (2016) Tong and Huang (2016) but few works focused on heterospecific pollen of far related species Thomson et al. (1982)

Galen and Gregory (1989) Neiland and Wilcock (1999) which also show a noteworthy fitness decrease.

Although the effect of close related species is predicted to be greater Ashman and Arceo-Gómez (2013)

the presence of pollen of non related species on multiple species Arceo-Gómez and Ashman (2016) and

the higher chances to coexist with a species that has less niche overlap (Ref) make foreign pollen from

far related species also an important subject of study in order to understand the importance of

heterospecific pollen in natural systems. Notwithstanding, the effect of heterospecific pollen of far and

close related species at community level remains to be explored beyond single pairwise interactions.

71 Paragraph 3 Expanding ideas with examples

Interestingly, incompatibility system seems to play an important role in foreign pollen effect where species that are self incompatible would have stronger barriers towards heterospecific pollen than self compatible species Ashman and Arceo-Gómez (2013). The type of incompatibility, sporophytic or gametophytic is related with the place of pollen recognition where the former take place at the sitgma 75 level and the latter occurs within the style, this last late acting pollen recognition mechanism is associated with greater negative effect Barrett (1988). Remarkably, there is a great variability in 77 mating systems across populations Whitehead et al. (2018) and therefore predict an effect of foreign pollen is a bit obscured by the variability within species, however species that are strong selfers or strong outcrossers have less variablity in mating systems and predictions of effect could be more realistic (see figure 1 from Whitehead et al. (2018)). Moreover, other traits such as number of pollen 81 grains per flower and number of ovules have been tradittionally associated with the type of 82 incompatibility system where species with higher pollen ovule ratios are predicted to be xenogamous and species with low pollen ovule ratios autogamous (REF). Selfer species would have a reduction of herkogamy (REF), less pollen production per ovule (REF) and therefore less likely to contribute in exporting pollen in the community. Other morphological traits, like stigma size can be determinant for the total pollen quatity that a stigma can receive and therefore related to do that pollen size would also play an important role. Example with pollen here.

Paragraph 4 Introducing our experiment

The great environmental variability in natural systems and complexity of floral structures make
heterospecific pollination studies a daunting task. Moreover, variation in sampling effort have been

shown to be determinant to characterize pollen transfer interactions Arceo-Gómez et al. (2018).

Although plant-pollinator network and pollen network studies can give a first picture of the importance of foreign pollen is necessary to address how its effect is shaped with both traits and relatedness of the species. For this reason, in this study we have created an artificial co-flowering community with 10 species belonging to three different families where we try to test the following questions: 1) Does heterospecific pollen reduce seed set, if so, 2) Does heterospecific pollen effect depend on the relatedness of the species, 3) Does heterospecific pollen effect depend on any floral trait?

99 METHODS

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

Table 1

Family	Genus	Species	
Brassicaceae	Brassica	Brassica rapa	
Brassicaceae	Brassica	Brassica oleracea	
Brassicaceae	Eruca	Eruca versicaria	
Brassicaceae Sinapis		Sinapis alba	
Convolvulaceae	Ipomoea	Ipomoea aquatica	
Convolvulaceae	Ipomoea	Ipomoea purpurea	
Solanaceae	Capsicum	Capsicum annuum	
Solanaceae	Petunia	Petunia integrifolia	

Family	Genus	Species
Solanaceae	Solanum	Solanum lycopersicum
Solanaceae	Solanum	Solanum melongena

109 Hand-pollination

Foreign pollen effect was studied through two different treatments, one with 50% conspecific pollen and 110 50% heterospecific pollen and a second one with 100% foreign pollen (N=10). Therefore, 180 different 111 combinations were perform with N=10. Seed set was the proxy of effect for all our teratments. Moreover, hand cross pollination, hand self pollination, apomixis (bagged emasculated flowers) and natural selfing were tested (N=10). Flowers were emasculated the day prior anthesis and hand 114 pollinated next day with a toothpick. HaNd-pollination was realized with 3-4 gentle touches on the 115 stigma surface. The mixes of pollen were performed on an eppendorf based on the pollen counts maded 116 with Neubaeur chamber (each anther was counted 4 times for 20 different anthers per species). In order 117 to confirm that the treatments applied were 50-50% pollen we counted for each species the total pollen 118 stigmatic load from a donor of each family (N=3). 119

120 Traits and evolutive distance

The traits measured for each species were pollen per anther, number of ovules, stigma width and length and stigmatic area, style width and length, ovary width and length. Moreover stigma type was tested. Pollen was counted for 20 anthers of each species with 4 replicates per sample with an hemocytometer. 123 Previously anthers were squashed on a known solution with the pippete tip and homogeneize with a 124 vortex for 30 seconds. Ovule number was counted with the help of an stereomicroscope and a small 125 grid over a petri dish from 15 randomly selected flowers. The different morphometrical traits were 126 measured with XXXX. Levels of self incompatibility were estimated by dividing the the fruit set of hand self pollination by hand cross pollination. Moreover, two types of evolutive distances were 128 calculated with MEGA7 for two kinds of markers: 1) Internal transcribed spacer (ITS) and 2) 129 ribulose-bisphosphate carboxylase (RBCL)

31 Analysis

We used the statistical language R (R Core Team 2018) for all our analyses. These were implemented in dynamic rmarkdown documents using knitr (Xie 2014, 2015, 2018) and rmarkdown (Allaire et al. 2018) packages. To test the effect of heterospecific pollen, we substracted the to seed set of hand cross pollination the seed set of the heterospecific pollen treatments. Therefore, small values mean low effect and viceversa. To be able to compare among species seed set was previously scaled for all species. In order to see relations between heterospecific pollen effect and traits we performed Mantel test between the matrix of effect and the distance matrix of each trait (euclidean distances). Moreover, Mantel test was also conducted between heterospecific pollen effect and phylogenetic distance.

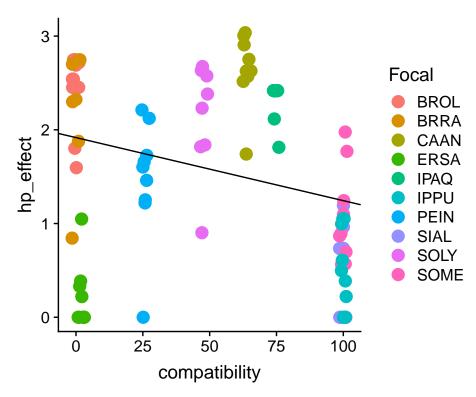


Figure 1: The effect of heterospecific pollen (scaled see set) is represented in function of the compatibility system (self/cross*100) for the the different species. Each coulored dot represents the interaction of a focal species with a different pollen donor.

140 RESULTS

Heterospecific pollen produced fitness for xxx cases.

142 DISCUSSION

143 Discussion

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1. What are the implications of the findings?

145 CONCLUSIONS

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147 REFERENCES

- Aizen, M. A., and L. D. Harder. 2007. Expanding the limits of the pollen-limitation concept: Effects of
- pollen quantity and quality. Ecology 88:271–281.
- Allaire, J., Y. Xie, J. McPherson, J. Luraschi, K. Ushey, A. Atkins, H. Wickham, J. Cheng, and W.
- 151 Chang. 2018. Rmarkdown: Dynamic documents for r.
- Arceo-Gómez, G., C. Alonso, T.-L. Ashman, and V. Parra-Tabla. 2018. Variation in sampling effort
- affects the observed richness of plant-plant interactions via heterospecific pollen transfer: Implications
- for interpretation of pollen transfer networks. American journal of botany 105:1601–1608.
- Arceo-Gómez, G., and T.-L. Ashman. 2016. Invasion status and phylogenetic relatedness predict cost
- of heterospecific pollen receipt: Implications for native biodiversity decline. Journal of Ecology
- 157 104:1003-1008.
- ¹⁵⁸ Arceo-Gómez, G., R. A. Raguso, and M. A. Geber. 2016. Can plants evolve tolerance mechanisms to
- heterospecific pollen effects? An experimental test of the adaptive potential in clarkia species. Oikos
- 160 125:718-725.
- Ashman, T.-L., and G. Arceo-Gómez. 2013. Toward a predictive understanding of the fitness costs of
- 162 heterospecific pollen receipt and its importance in co-flowering communities. American Journal of
- 163 Botany 100:1061-1070.
- Barrett, S. C. 1988. The evolution, maintenance, and loss of self-incompatibility systems. Plant
- reproductive ecology: patterns and strategies:98–124.
- Bartomeus, I., J. Bosch, and M. Vilà. 2008. High invasive pollen transfer, yet low deposition on native

- stigmas in a carpobrotus-invaded community. Annals of Botany 102:417–424.
- Bascompte, J., P. Jordano, C. J. Melián, and J. M. Olesen. 2003. The nested assembly of plant–animal
- mutualistic networks. Proceedings of the National Academy of Sciences 100:9383–9387.
- Brown, B. J., and R. J. Mitchell. 2001. Competition for pollination: Effects of pollen of an invasive
- plant on seed set of a native congener. Oecologia 129:43–49.
- Carvalheiro, L. G., J. C. Biesmeijer, G. Benadi, J. Fründ, M. Stang, I. Bartomeus, C. N.
- Kaiser-Bunbury, M. Baude, S. I. Gomes, V. Merckx, and others. 2014. The potential for indirect effects
- between co-flowering plants via shared pollinators depends on resource abundance, accessibility and
- relatedness. Ecology letters 17:1389–1399.
- Engel, E. C., and R. E. Irwin. 2003. Linking pollinator visitation rate and pollen receipt. American
- 177 Journal of Botany 90:1612–1618.
- Fang, Q., and S.-Q. Huang. 2013. A directed network analysis of heterospecific pollen transfer in a
- biodiverse community. Ecology 94:1176–1185.
- Galen, C., and T. Gregory. 1989. Interspecific pollen transfer as a mechanism of competition:
- 181 Consequences of foreign pollen contamination for seed set in the alpine wildflower, polemonium
- viscosum. Oecologia 81:120–123.
- Inouye, D. W. 1980. The terminology of floral larceny. Ecology 61:1251–1253.
- Montgomery, B. R., and B. J. Rathcke. 2012. Effects of floral restrictiveness and stigma size on
- heterospecific pollen receipt in a prairie community. Oecologia 168:449–458.
- Morales, C. L., and A. Traveset. 2008. Interspecific pollen transfer: Magnitude, prevalence and
- consequences for plant fitness. Critical Reviews in Plant Sciences 27:221–238.
- Neiland, M., and C. Wilcock. 1999. The presence of heterospecific pollen on stigmas of nectariferous
- and nectarless orchids and its consequences for their reproductive success. Protoplasma 208:65–75.
- Pauw, A. 2013. Can pollination niches facilitate plant coexistence? Trends in ecology & evolution

- 191 28:30-37.
- 192 R Core Team. 2018. R: A language and environment for statistical computing. R Foundation for
- 193 Statistical Computing, Vienna, Austria.
- Thomson, J. D., B. J. Andrews, and R. Plowright. 1982. The effect of a foreign pollen on ovule
- development in diervilla lonicera (caprifoliaceae). New Phytologist 90:777–783.
- Tong, Z.-Y., and S.-Q. Huang. 2016. Pre-and post-pollination interaction between six co-flowering
- pedicularis species via heterospecific pollen transfer. New Phytologist 211:1452–1461.
- Whitehead, M. R., R. Lanfear, R. J. Mitchell, and J. D. Karron. 2018. Plant mating systems often vary
- widely among populations. Frontiers in Ecology and Evolution 6:38.
- 200 Xie, Y. 2014. Knitr: A comprehensive tool for reproducible research in R. in V. Stodden, F. Leisch,
- ²⁰¹ and R. D. Peng, editors. Implementing reproducible computational research. Chapman; Hall/CRC.
- ²⁰² Xie, Y. 2015. Dynamic documents with R and knitr. 2nd editions. Chapman; Hall/CRC, Boca Raton,
- 203 Florida.
- ²⁰⁴ Xie, Y. 2018. Knitr: A general-purpose package for dynamic report generation in r.

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