# working title Compatibility system and stigma size are the main predictors of heterospecific pollen effect

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

<sup>4</sup>
<sup>5</sup>
<sup>1</sup>US 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

<sup>7</sup>
<sup>8</sup>
<sup>2</sup>Big Name University, Department of R, City, BN, 01020, USA

<sup>9</sup>
<sup>10</sup>
<sup>3</sup>Estacion Biologica de Donana (EBD-CSIC), E-41092 Sevilla, Spain

11

\* corresponding author: barragansljose@gmail.com

Pollinator sharing can have negative consequences for plant fitness with the arrival of foreign pollen. However, the costs of heterospecific pollen are not yet well understood. We conducted a glasshouse 14 experiment to understand how phylogenetic relatedness and plant traits mediate the impacts of 15 heterospecific pollen transfer. We conducted 1800 crosses by experimentally transferring pollen (50%) 16 and 100% ratio) with reciprocal crosses between 10 species belonging to three different families: 17 Brassicaceae, Solanaceae and Convolvulaceae. Seed set was used as proxy of plant fitness. We found that for 65% of the treatments with 50% mix reduced seed set. Moreover, the reduction in seed set 19 was dependent on the degree of relatedness and reproductive traits of the pollen recipient and not the pollen donor. Our results show that certain traits, particularly compatibility system, are critical in 21 understanding the costs of heterospecific pollen.

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

#### 25 INTRODUCTION

- In most ecosystems, plant species normally coexist and share their floral visitors with other species
  Waser et al. (1996). From the plants' perspective, pollinator sharing can be positive for some plants
- Carvalheiro et al. (2014) or negative for others Pauw (2013), depending on the facilitation gradient. An

increasing number of visits often correlates with higher chances of fertilization Engel and Irwin (2003).

- However this is not always the case, among these possible flower visitors there are also nectar robbers
- and pollen thieves Inouye (1980); Magrach et al. (2017). Receiving both sufficient quantity and quality
- deposited on the stigma is thus highly relevant to the pollination success of the plant Aizen and Harder
- зз (2007).

3

By visiting many plant species, many pollinators are responsible for conspecific pollen loss and the transport of foreign pollen, both of which can have important detrimental effects on species fitness Morales and Traveset (2008); Ashman and Arceo-Gómez (2013); Arceo-Gómez and Ashman (2016). Foreign pollen arrival can play an important role in plant species fitness but outcomes are variable and 37 appear to be context dependent as there is not always a decrease in fitness Morales and Traveset (2008). Some of this variation is likely due to the enormous variability of foreign pollen transferred across systems ranging from 0 to 75 percent. However, most studies report ranges of heterospecific pollen 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), yet even these relatively low 42 amounts of heterospecific pollen transferred can decrease fitness greatly Thomson et al. (1982). While 43 we now have some understanding of the impacts of heterospecific pollen quantity, we have less understanding of other factors that could be driving the variation in impacts upon fitness. Ashman and Arceo-Gómez (2013) postulated the first predictive framework that identifies a need to understand how plant traits might mediate heterospecific pollen effect, whereby mating system and pollen size were 47 predicted to potentially mediate the impact of foreign pollen transfer on plant fitness. This concept is supported by specific case studies, such as Tong and Huang (2016) that demonstrate an asymmetrical effect in 6 species of *Pedicularis* whereby the pollen of long styled species was able to grow the full length of the style on short styled species but not vice versa. While this suggests that the impacts of heterospecific pollen may differ among pollen donor and recipient, few studies have been conducted to ascertain whether this pattern is in fact a general trend or to identify the extent to which other plant traits are critical to heterospecific pollen impacts. Plant traits are crucial to understand heterospecific pollen effect but the multifactorial nature of the traits that are involve in the pollen-pistil interaction make difficult to unravel what are the main traits in driving the effect. These traits can be seen from a male perspective of both donor and recipient where pollen size, pollen aperture number and pollen allelopathy are key components to understand the outcome of foreign pollen arrival Murphy and Aarssen (1995); Ashman and Arceo-Gómez (2013). In 59 Ashman and Arceo-Gómez (2013) small pollen is predicted to cause a greater fitness decrease, although this can be true there are also other possibilities to consider which can obscure a predictive framework like big pollen can clogg small stigmas with fewer pollen grains, bigger stigmas are less likely to be

clogged by small pollen grains and bigger pollen can outcompete smaller pollen grains due to faster pollen tube growth rate Williams and Rouse (1990). Moreover, to understand the different mechanical or chemical effects of pollen also the female traits of the pollen recipient have to be considered, from the literature these main traits are: stigma size, style length, number of ovules, incompatibility system and also a structural trait such as flower morphology Montgomery and Rathcke (2012); Ashman and 67 Arceo-Gómez (2013); Tong and Huang (2016). For example, greater stigmatic area is positively correlated with greater heterospecific pollen deposition Montgomery and Rathcke (2012) and therefore possibly with an increase in negative effect. For species that are self-incompatible the barriers towards heterospecific pollen are stronger than self-compatible species Ashman and Arceo-Gómez (2013). Nonetheless, 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)). Although past research has progress in the understanding of what traits can mediate the effect as we have shown here, there are multiple traits involved and multiple possible scenarios still to be explored empirically for a full understanding of the importance of heterospecific pollen effect in nature. For the understanding at what level or intensity the interference of pollen can occur is important to consider the relatedness of the interacting species. Closely related species are more likely to have similar traits Letten and Cornwell (2015). The similarity in traits between closely related species can lead to higher chances of ovule usurpation/abortion Arceo-Gómez and Ashman (2011) and therefore studies predict and show a greater negative effect of closely related species Ashman and Arceo-Gómez (2013); Arceo-Gómez and Ashman (2016). Few studies however, have focused on the impacts of heterospecific pollen on fitness of distantly related species Galen and Gregory (1989); Neiland and Wilcock (1999). Despite the fact that far related species are also able to decrease species fitness (REFS). Yet, most insects and most stigmas have been found to carry multiple species of foreign pollen with little attention to degree of relatedness (Arceo-Gómez and Ashman (2016); Fang and Huang (2013). Understanding the role of foreign pollen from distantly related species thus deserves greater attention. The relatedness of foreign pollen gives a first snapshoot of where the pollen competition can occur and therefore could be a proxy of effect. With that said, Arceo-Gómez and Ashman (2016) is the only work which has proven a greater effect of close related species through a meta-analysis but with

- low sample sizes and lack of significance. Therefore, there is a need to study the effect of heterospecific pollen of far and close related species at community level beyond single pairwise interactions.
- Heterospecific pollen studies in nature have the complexity added of great environmental variability which can lead to confounding interpretations. Moreover, the great diversity of floral structures and small sizes of some of them such as Asteraceae species make tedious or almost impossible the studies of heterospecific pollen on the field. For this reason, we investigated how floral reproductive traits and relatedness mediate the impact of heterospecific pollen by creating an artificial co-flowering community in a glasshouse with 10 species belonging to three different families with different traits. Our study tries to answer the following questions: (i) Do heterospecific pollen reduce seed set? To what extent do (ii) floral reproductive traits and (iii) relatedness, mediate the impacts of heterospecific pollen on seed set.

## 102 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 104 the species with day and night temperature differences. The experimental design had species from 105 three different families: Solanaceae, Brassicaceae and Convolvulaceae (Table 1). The species of the 106 study had different reproductive traits and different degree of relatedness where the reciprocal crosses 107 between species allowed us to have multiple different scenarios of both traits and relatedness. Moreover, 108 the species selected had fast life cycle and low structural flower complexity in order to perform the 109 pollination treatments and grow the different species from seeds. For the purpose of the experiment all 110 the species where considered as pollen recipient and as pollen donor (see interaction matrix, fig 2). 111 Species were watered once or twice per day and fertilized weekly (NPK 23: 3.95: 14). 112

#### 113 Table 1

Family	Genus	Species
Brassicaceae	Brassica	Brassica rapa
Brassicaceae	Brassica	Brassica oleracea

Family	Genus	Species	
Brassicaceae	Eruca	Eruca versicaria	
Brassicaceae	Sinapis	Sinapis alba	
Convolvulaceae	Ipomoea	Ipomoea aquatica	
Convolvulaceae	Ipomoea	Ipomoea purpurea	
Solanaceae	Capsicum	Capsicum annuum	
Solanaceae	Petunia	Petunia integrifolia	
Solanaceae	Solanum	Solanum lycopersicum	
Solanaceae	Solanum	Solanum melongena	

#### 114 Hand-pollination

Foreign pollen effect was studied through two different treatments, one with 50% conspecific pollen and 115 50% heterospecific pollen and a second one with 100% foreign pollen (N=10) in order to see if foreign 116 pollen can trigger fruit production by itself or even seeds through ovule usurpation. Therefore, 180 117 different combinations were performed with N=10 per combination. Seed set was the proxy of effect for 118 all our treatments. Moreover, hand cross pollination (between individuals of the same species), hand 119 self pollination, apomixis (bagged emasculated flowers) and natural selfing were tested for each species 120 (N=10). For the treatments with foreign pollen and hand cross pollination, flowers were emasculated 121 the day prior anthesis and hand pollinated next day with a toothpick. Hand-pollination was conducted with 3-4 gentle touches on the stigma surface. For each species 20 anthers were collected and their 123 pollen counted with a hemocytometer, each anther was counted 4 times and then an average of these 124 counts was performed. Once, the average number of pollen grains per anther was known, the 125 proportion of anthers per mix was calculated in order to achieve a 50-50% mix. In order to confirm 126 that the treatments applied were the desire proportions, the total stigmatic load of pollen was counted 127 and the proportions calculated between the two species of the mix. Therefore, 20 different 128 heterospecific crosses 50-50% were counted for all the focal species with crosses of just one species of 129 the other two families (N=3). 130

#### Traits and evolutive distance

The traits measured for each species were pollen per anther, number of ovules and stigma, style, ovary width and length. For the stigma, the stigmatic area was also measured and moreover the stigmas were divided in wet/dry type. All the morphometrical measurements were performed with a stereophotomicroscope. Pollen was counted for 20 anthers of each species with 4 replicates per sample 135 with an hemocytometer. Previously, anthers were squashed on a known solution with the pippete tip 136 and homogeneize with a vortex for 30 seconds. Ovule number was counted with the help of a 137 stereomicroscope and a small grid over a petri dish from 15 randomly selected flowers. Fruits per 138 number of flowers treated were counted for Solanaceae species and for the rest, the number of seeds 139 produced per average number of ovules of each species. Levels of self incompatibility were estimated by 140 dividing the fruit set of hand self pollination by hand cross pollination Lloyd and Schoen (1992).

#### 142 Analysis

To evaluate heterospecific pollen effect on seed production we performed linear mixed models. The
distinct heterospecific pollination treatments were compared through relevelling each variable with the
cross pollination treatment which was our control for optimum seed production for all the species. The
different replicates of each treatment were considered as random effects. Seed production was scaled for
all the species with mean 0 and standard deviation of 1 prior to the analysis. All the analysis were
conducted with the statistical language R (R Core Team 2018).

In order to test the relative effect of traits on seed production with foreign pollen we performed Mantel 149 test between the assymetrical matrix of heterospecific pollen effect (10 by 10 matrix) with the different 150 distance matrices of traits (Euclidean distances). Heterospecific pollen effect was obtained through the 151 subtraction of seed production by hand cross pollination minus seed production of the different 152 heterospecific pollen treatments. To find a model with the best explanatory traits we used the function bioenv from R. We also conducted Mantel test between the matrix of heterospecific pollen effect and 154 the distance matrix from all the traits. Moreover, we explored also the correlation between traits and 155 heterospecific pollen effect through generalized mixed models where the response variable was 156 heterospecific pollen effect and the explanatory variable the different traits. In addition, we tested the 157 correlation between the total amount of pollen deposited on the stigma with heterospecific pollinations and the stigma size through Pearson's correlation. 159

We conducted mantel test to check for correlations between heterospecific pollen effect and phylogenetic distance. Due to improvents in statistical power we used the square root of the phylogenetic distance (Letten & Cornwell 2014). Two different phylogenetic distances were used from two kinds of markers: 1)

Internal transcribed spacer (ITS) and 2) ribulose-bisphosphate carboxylase (RBCL). The sequences were obtained from GenBank (https://www.ncbi.nlm.nih.gov/genbank/, accessed 20 Oct. 2018). The sequences were aligned with clustal omega and the pairwise evolutive distances calculated with MEGA7.

Phylogenetic signal of traits?

167 Add pollen on stigma counts to methods

168 [Next bit to results, also divided it with]

Total pollen deposited on stigmas was significantly correlated with the stigmatic area, pearsons correlation=0.57 and p-value=0.008. Proportions of pollen talk about it to 50-50 mix? Talk about this in methods too.

Add also plot with fruit production and pot it in appendix. Also the plots of ratios to appendices.

## 173 RESULTS

Results of hand cross-pollination, self hand-pollination, natural selfing and apomixis are presented in Figure 1 (see appendix 1 for table with values). Heterospecific pollen reduced seet set significatively 175 with the 50-50% heterospecific pollen treatments for 65% of the pairwise interactions p<0.05. Across families we found a very similar effect but when species where look at species level they respond 177 differently even within the same family rephrase and maybe test statistically?, for instance for two 178 species of the Brassicaceae family Brassica oleracea and Eruca versicaria we found very contrasting effects of foreign pollen where for the first one, all donors reduce seed set significatively and for the 180 second, just two species did out of nine. The 100% foreign pollen treatments barely produced seeds or 181 fruits and just for Sinapis alba we did not find significant differences between the hand cross pollination 182 and one treatment with pollen from a confamilial- IB Unclear. Solanaceae species with berry fruit type

developed small fruits or even normal fruits in some cases under which treatment. S. lycopersicum
seems to produced small fruits (35% of the treatments) independently of pollen and pollen donor due to
also apomictic treatments did, never normal size. C. annuum produced some fruits (9%) of both small
and normal size and finally S. melongena produced seedless normal fruits with just confamilial pollen
(3%), for both species seems that fruit formation was induced by pollen on the stigma because of lack
of fruit production with treatments that tested for apomixis.clarify this descriptive statistics part- Also
a figure with a summary of the treatments effect would be cool, or at least in the appendices

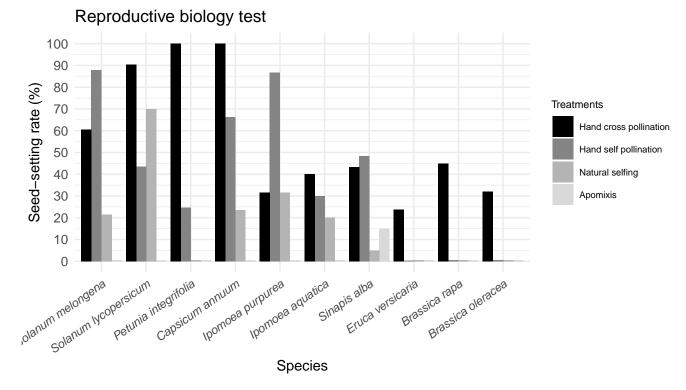


Figure 1: Barplot with the different treatments that provide information of the reproductive biology of the ten species. The y axis is the proportion of ovules converted to seed in percentage. The different treatments (N=10) which are presented in the legend are, hand cross pollination, hand self pollination, natural selfing and apomixis. More information about these treatments can be found in Methods and Appendices.

Mantel test indicates that a possible?? It exists! correlation exist between heterospecific pollen effect and the evolutive relative distances, for ITS and RBCL markers we had r coefficients of 0.29 and 0.25 respectively p<0.05 think on a figure - maybe using NMDS. Moreover, Mantel test indicates that also a possible?? correlation between stigma width and stigma type exist (stats??). Trait correlations were also explored with GLMM

I have done it at the moment just for Compatibility system Also I have to fix from mixed linear model to GLMM, just realize that

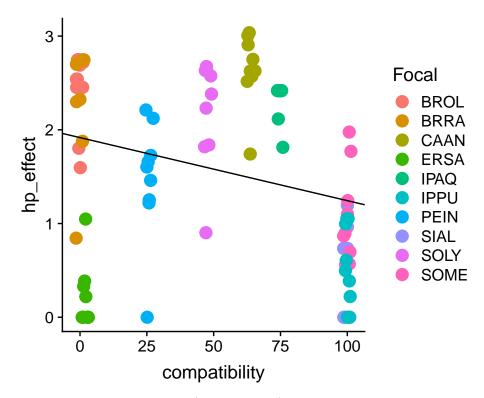


Figure 2: The effect of heterospecific pollen (scaled see set) is represented in function of the compatibility system (self/cross\*100) for the different species. Each coulored dot represents the interaction of a focal species with a different pollen donor.

8 Compatibility index don't multiply per 100 from Lloyd

# 199 DISCUSSION

Discussion Herbs vs tress, annual vs perennial... Many flowers vs few flowered species; structural composition on 201 a system 202 What are the implications of the findings? Ideas about pollen size in heterospecific pollen effect. (still have to develop it more...) 204 Let's classify pollen size in three groups in order to understand the interaction between pollen donor 205 and recipient: 1) Donor pollen size < Recipient pollen size 2) Donor pollen size = Recipient pollen size 206 3) Donor pollen size > Recipient pollen size Now I try to develop each part 1) Donor pollen size < Recipient pollen size 209 Effect: • Donor's pollen could clogg the stigma 211 • Chemical inhibition 212 Traits associated with bigger pollen of the recipient: 213 • Recipient's pollen have faster pollen tube growth (example with my data) 214 • Reduction in number of ovules (Also with my species) 215

• Very relatedness dependant this point

germination on a far related stigma.

2) Donor pollen size = Recipient pollen size

216

217

218

11 February 28, 2019

• Big differences in pollen size can be traduced in low relatednes therefore less likely of pollen

- Similar probabilities of taken space on the stigma
- 3) Donor pollen size > Recipient pollen size
- 222 Effect:
- 223 -In small stigmas big pollen grains can occupy great part of the stigmatic area.
- -small pollen grains can get embeded
- 225 IB: Think also on using tree analysis to test if hp effect depends on complex trait combinations. Tree
- 226 analysis are great when two different strategies lead to the same outcome. This would never been pick
- up by GLMs. The r package is party{} . You can see an example applied to birds is Sol et al 2010
- Science. Ask me if you want more details or code examples.

## 229 CONCLUSIONS

## 230 ACKNOWLEDGEMENTS

## 231 REFERENCES

- <sup>232</sup> 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.
- <sup>234</sup> Arceo-Gómez, G., and T.-L. Ashman. 2011. Heterospecific pollen deposition: Does diversity alter the
- consequences? New Phytologist 192:738–746.
- <sup>236</sup> 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
- 238 104:1003-1008.
- Ashman, T.-L., and G. Arceo-Gómez. 2013. Toward a predictive understanding of the fitness costs of

- 240 heterospecific pollen receipt and its importance in co-flowering communities. American Journal of
- 241 Botany 100:1061–1070.
- <sup>242</sup> 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.
- <sup>244</sup> Carvalheiro, L. G., J. C. Biesmeijer, G. Benadi, J. Fründ, M. Stang, I. Bartomeus, C. N.
- <sup>245</sup> 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
- <sup>249</sup> Journal of Botany 90:1612–1618.
- <sup>250</sup> 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:
- 253 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.
- Letten, A. D., and W. K. Cornwell. 2015. Trees, branches and (square) roots: Why evolutionary
- relatedness is not linearly related to functional distance. Methods in Ecology and Evolution 6:439–444.
- Lloyd, D. G., and D. J. Schoen. 1992. Self-and cross-fertilization in plants. i. functional dimensions.
- <sup>259</sup> International Journal of Plant Sciences 153:358–369.
- Magrach, A., J. P. González-Varo, M. Boiffier, M. Vilà, and I. Bartomeus. 2017. Honeybee spillover
- reshuffles pollinator diets and affects plant reproductive success. Nature ecology & evolution 1:1299.
- 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.
- Murphy, S. D., and L. W. Aarssen. 1995. Reduced seed set in elytrigia repens caused by allelopathic
- pollen from phleum pratense. Canadian Journal of Botany 73:1417–1422.
- Neiland, M., and C. Wilcock. 1999. The presence of heterospecific pollen on stigmas of nectariferous
- <sup>269</sup> 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
- 271 28:30-37.
- 272 R Core Team. 2018. R: A language and environment for statistical computing. R Foundation for
- 273 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.
- <sup>276</sup> 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.
- Waser, N. M., L. Chittka, M. V. Price, N. M. Williams, and J. Ollerton. 1996. Generalization in
- pollination systems, and why it matters. Ecology 77:1043–1060.
- <sup>280</sup> Whitehead, M. R., R. Lanfear, R. J. Mitchell, and J. D. Karron. 2018. Plant mating systems often vary
- <sup>281</sup> widely among populations. Frontiers in Ecology and Evolution 6:38.
- Williams, E., and J. Rouse. 1990. Relationships of pollen size, pistil length and pollen tube growth
- rates in rhododendron and their influence on hybridization. Sexual Plant Reproduction 3:7–17.

# APPENDIX

285 1.

Table S1. Perecentage of seeds produced per ovule for the ten species used in the experiment. The treatments presented are hand cross pollination, hand self pollination, natural selfing and apomixis (emasculated flowers).

Species	Cross	Self	Natural_selfing	Apomixis
Brassica oleracea	32.06897	0.0000000	0.00000	0
Brassica rapa	44.97041	0.0000000	0.00000	0
Eruca versicaria	23.75000	0.4166667	0.00000	0
Sinapis alba	43.33333	48.3333333	5.00000	15
Ipomoea aquatica	40.00000	30.0000000	20.00000	0
Ipomoea purpurea	31.66667	86.6666667	31.66667	0
Capsicum annuum	100.00000	66.2240664	23.48548	0
Petunia integrifolia	100.00000	24.7727273	0.00000	0
Solanum lycopersicum	90.38043	43.4782609	70.00000	0
Solanum melongena	60.47525	87.9702970	21.56436	0

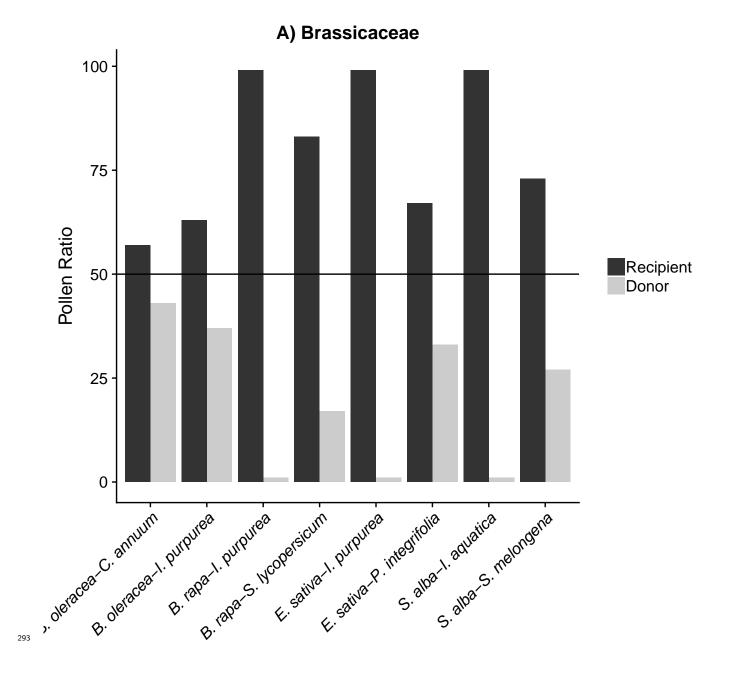
2.

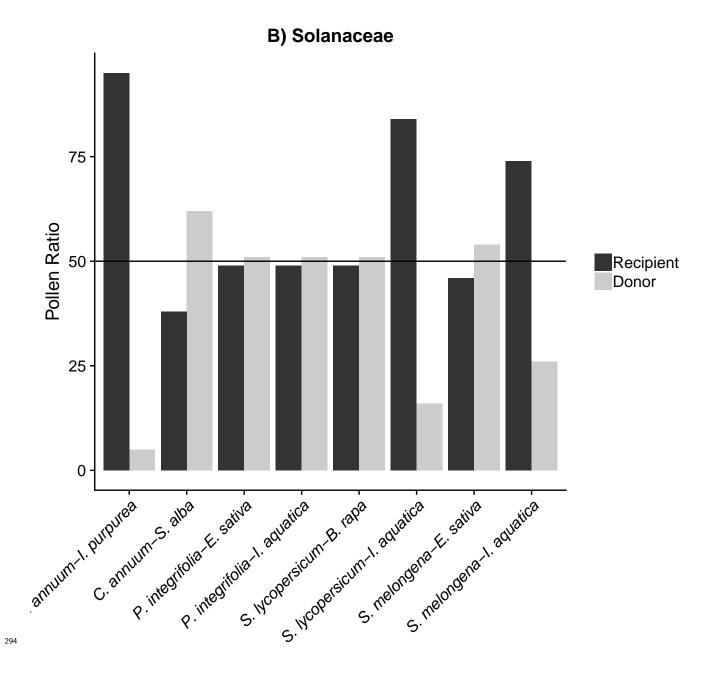
289

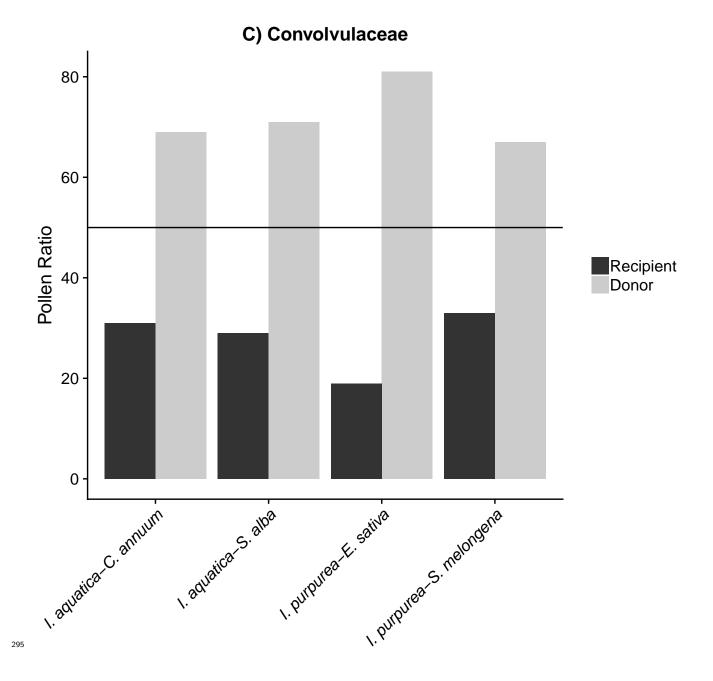
Figure S1. Pollen ratios separated by family A) Brassicaceae, B) Solanaceae, C) Convolvulaceae.

Pollen was counted for crosses with species of the other two families (N=3). To prepare these ratios all

292 the pollen grains on the stigma were counted.







# List of Tables

# List of Figures

298	1	Barplot with the different treatments that provide information of the reproductive biology	
299		of the ten species. The y axis is the proportion of ovules converted to seed in percentage.	
300		The different treatments (N=10) which are presented in the legend are, hand cross	
301		pollination, hand self pollination, natural selfing and apomixis. More information about	
302		these treatments can be found in Methods and Appendices	9
303	2	The effect of heterospecific pollen (scaled see set) is represented in function of the	
304		compatibility system (self/cross*100) for the the different species. Each coulored dot	
305		represents the interaction of a focal species with a different pollen donor	10