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

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¹ US Environmental Protection Agency, Office of Research and Development, National Health and Environmental Effects
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² Big Name University, Department of R, City, BN, 01020, USA

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¹⁰
³ Estacion Biologica de Donana (EBD-CSIC), E-41092 Sevilla, Spain

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* 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 experiment to understand how phylogenetic relatedness and plant traits mediate the impacts of heterospecific pollen transfer. We conducted 4XXXX crosses by experimentally transferring pollen (50% and 100% ratio) with reciporcal crosses between 10 species belonging to three different families: 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 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 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
- ²⁸ Carvalheiro et al. (2014) or negative for others Pauw (2013), depending on the facilitation gradient. An
- 29 increasing number of visits often correlates with higher chances of fertilization Engel and Irwin (2003).
- 30 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
- $_{32}$ deposited on the stigma is thus highly relevant to the pollination success of the plant Aizen and Harder

(2007).

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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

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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
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   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). 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). The relative amounts of closely/far related pollen that arrives to a stigma gives a
   first snapshoot where pollen competition can occur and improves the understanding of how important
   the reduction of fitness can be from heterospecific pollen where closely related species cand lead to
   greater detrimental effects Arceo-Gómez and Ashman (2016).
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 $_{91}$ Further, a majority of plant species are generalist and thus receive visits from multiple different

pollinators. Given these are generally the ones that receive greater loads of heterospecific pollen Fang and Huang (2013) and unrelated species are more likely to coexist with other species due to less niche overlap (Ref), understanding the role of foreign pollen from distantly related species thus deserves greater attention in understanding coexistence blah blahXXXXX refs.. Notwithstanding, the effect of heterospecific pollen of far and close related species at community level remains to be explored beyond single pairwise interactions. Moreover, heterospecific pollen studies in nature have the complexity 97 added great environmental variability which can lead to confounding interpretations and diverse floral structures and some of them complex such as Asteraceae species, which involve tedious work to be able to study foreign pollen effect. For this reason, we investigated how floral reproductive traits and 100 relatedness mediate the impact of heterospecific transfer by asking the following research questions: To 101 what extent do (i) floral reproductive traits and (ii) relatedness, mediate the impacts of heterospecific 102 pollen on seed set. We do this by creating an artificial co-flowering community with 10 species 103 belonging to three different families with different traits. 104

$_{\scriptscriptstyle{5}}$ 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)I would
explain more the bauty of our nested dessign to ensure close and far distance simultaneously,
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).

5 Table 1

Family	Genus	Species
Brassicaceae	Brassica	Brassica rapa

Family	Genus	Species
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
Solanaceae	Solanum	Solanum lycopersicum
Solanaceae	Solanum	Solanum melongena

116 Hand-pollination

Foreign pollen effect was studied through two different treatments, one with 50% conspecific pollen and 117 50% heterospecific pollen and a second one with 100% foreign pollen (N=10) this second I don't get, 118 maybe explain it's ultility.. Therefore, 180 different combinations were performed with N=10 per 119 combination. Seed set was the proxy of effect for all our treatments. Moreover, hand cross pollination 120 (between individuals of the same species), hand self pollination, apomixis (bagged emasculated flowers) 121 and natural selfing were tested for each species (N=10). For the treatments with foreign pollen and 122 hand cross pollination, flowers were emasculated the day prior anthesis and hand pollinated next day 123 with a toothpick. Hand-pollination was conducted with 3-4 gentle touches on the stigma surface. The mixes of pollen were realized on an eppendorf based on the pollen counts made with Neubaeur chamber 125 (each anther was counted 4 times for 20 different anthers per species)-IB explain better and give a bit 126 more of detail. In order to confirm that the treatments applied were 50-50 percent pollen, for each focal 127 species the total stigmatic load of pollen was counted from one donor of each family (N=3). 128

29 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 explain was

tested. Pollen was counted for 20 anthers of each species with 4 replicates per sample with an
hemocytometer. Previously, anthers were squashed on a known solution with the pippete tip and
homogeneize with a vortex for 30 seconds. Ovule number was counted with the help of an
stereomicroscope and a small grid over a petri dish from 15 randomly selected flowers. The different
morphometrical traits were measured with a digital stereomicrospe. Levels of self incompatibility were
estimated by dividing the fruit set of hand self pollination by hand cross pollination Lloyd and Schoen
(1992).

139 Analysis

We used the statistical language R (R Core Team 2018) for all our analyses. Differences of seed set between treatments and hand cross pollination for each species was tested through mixed linear models. 141 For the following analysis we scaled the values of seed production for all the species with mean 0 and sd of 1. To test the effect of heterospecific pollen, we substracted to the seed set of hand cross pollination 143 the seed set of heterospecific pollen treatments. In order to see correlations between heterospecific 144 pollen effect and traits we performed Mantel test between the matrix of heterospecific pollen effect and 145 the distance matrix of each trait (euclidean distances). Moreover, Mantel test was also conducted 146 between heterospecific pollen effect and the square root of the matrix of phylogenetic distance due to 147 improvement in the statistical power (Letten & Cornwell 2014). all is here, but I would break it by questions and give a bit more detail, to avoid overwhelm the reader We explored also the relations 149 between traits and heterospecific pollen effect through generalized mixed models where the response 150 variable was heterospecific pollen effect, the independent variable the different traits and the random 151 effects the different treatments per species [Here I think you should think if this controls for the non 152 independency of donors and recipients. I think not. Maybe look onto matrix regresions?). Moreover, pairwise evolutive distances were calculated with MEGA7 for two kinds of markers: 1) Internal 154 transcribed spacer (ITS) and 2) ribulose-bisphosphate carboxylase (RBCL). The sequences of interest 155 were downloaded from NCBI GenBank and the phylogenetic tree constructed by maximum likelihood 156 with MEGA7. Make a section on how you contriucted phylogeny. 157

I would explain three test. 0) treatment effects with GLM's, 1) Mantels: relative effects, 2) GLM's or matrix models: Absolute effects and explain them in three independent paragraphs including a

160 rationale of why

Phylogenetic signal of traits?

162 RESULTS

Results of hand cross pollination, self hand pollination, natural selfing and apomixis are presented in 163 Table 2. Heterospecific pollen reduced seet set significatively with the 50-50% heterospecific pollen 164 treatments for 65% of the pairwise interactions p<0.05. Across families we found a very similar effect 165 but when species where look at species level they respond differently even within the same family 166 rephrase and maybe test statistically?, for instance for two species of the Brassicaceae family Brassica oleracea and Eruca versicaria we found very contrasting effects of foreign pollen where for the first one, 168 all donors reduce seed set significatively and for the second, just two species did out of nine. The 100% 169 foreign pollen treatments barely produced seeds or fruits and just for Sinapis alba we did not find 170 significant differences between the hand cross pollination and one treatment with pollen from a 171 confamilial- IB Unclear. Solanaceae species with berry fruit type developed small fruits or even normal 172 fruits in some cases under which treatment. S. lycopersicum seems to produced small fruits (35% of the 173 treatments) independently of pollen and pollen donor due to also apomictic treatments did, never 174 normal size. C. annuum produced some fruits (9%) of both small and normal size and finally S. 175 melongena produced seedless normal fruits with just confamilial pollen (3%), for both species seems 176 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 178 of the treatments effect would be cool, or at least in the appendices

Table 2. 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). turn into a figure somehow?

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

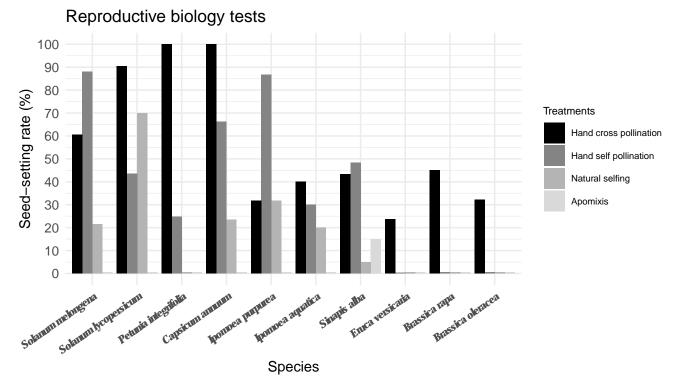


Figure 1: Barplot with the different treatments that provide information of the reproductive biology of the ten species. The y axis is the proportion of ovules converted to seed in percentage. The different treatments (N=10) which are presented in the legend are, hand cross pollination, hand self pollination, 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

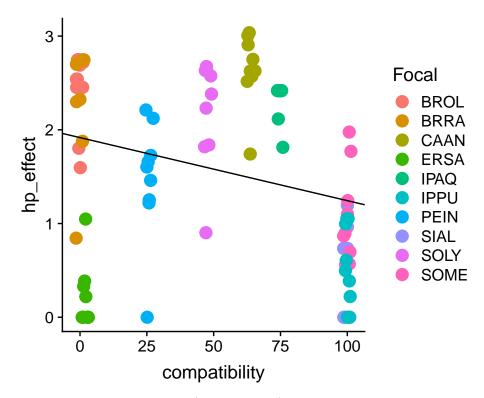


Figure 2: The effect of heterospecific pollen (scaled see set) is represented in function of the compatibility system (self/cross*100) for the different species. Each coulored dot represents the interaction of a focal species with a different pollen donor.

Compatibility index don't multiply per 100 from Lloyd

191 DISCUSSION

- Discussion 192 Herbs vs tress, annual vs perennial... Many flowers vs few flowered species; structural composition on 193 a system 194 What are the implications of the findings? Ideas about pollen size in heterospecific pollen effect. (still have to develop it more...) 196 Let's classify pollen size in three groups in order to understand the interaction between pollen donor 197 and recipient: 1) Donor pollen size < Recipient pollen size 2) Donor pollen size = Recipient pollen size 198 3) Donor pollen size > Recipient pollen size Now I try to develop each part 200 1) Donor pollen size < Recipient pollen size 201 Effect: 202 • Donor's pollen could clogg the stigma 203

• Chemical inhibition

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• Recipient's pollen have faster pollen tube growth (example with my data)

• Reduction in number of ovules (Also with my species)

Traits associated with bigger pollen of the recipient:

- Big differences in pollen size can be traduced in low relatednes therefore less likely of pollen germination on a far related stigma.
- 2) Donor pollen size = Recipient pollen size
- Very relatedness dependant this point

- Similar probabilities of taken space on the stigma
- 3) Donor pollen size > Recipient pollen size
- 214 Effect:
- 215 -In small stigmas big pollen grains can occupy great part of the stigmatic area.
- -small pollen grains can get embeded
- ²¹⁷ IB: Think also on using tree analysis to test if hp effect depends on complex trait combinations. Tree
- 218 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
- 220 Science. Ask me if you want more details or code examples.

221 CONCLUSIONS

222 ACKNOWLEDGEMENTS

223 REFERENCES

- ²²⁴ 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.
- ²²⁶ Arceo-Gómez, G., and T.-L. Ashman. 2011. Heterospecific pollen deposition: Does diversity alter the
- consequences? New Phytologist 192:738–746.
- ²²⁸ 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
- 230 104:1003-1008.
- Ashman, T.-L., and G. Arceo-Gómez. 2013. Toward a predictive understanding of the fitness costs of

- 232 heterospecific pollen receipt and its importance in co-flowering communities. American Journal of
- 233 Botany 100:1061–1070.
- Bartomeus, I., J. Bosch, and M. Vilà. 2008. High invasive pollen transfer, yet low deposition on native
- 235 stigmas in a carpobrotus-invaded community. Annals of Botany 102:417–424.
- ²³⁶ 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
- ²⁴¹ 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:
- ²⁴⁵ 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.
- ²⁵¹ 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
- 261 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
- 263 28:30-37.
- ²⁶⁴ R Core Team. 2018. R: A language and environment for statistical computing. R Foundation for
- ²⁶⁵ 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.
- ²⁶⁸ 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.
- Whitehead, M. R., R. Lanfear, R. J. Mitchell, and J. D. Karron. 2018. Plant mating systems often vary
- ²⁷³ 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
- 275 rates in rhododendron and their influence on hybridization. Sexual Plant Reproduction 3:7–17.

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