

Principles governing F1 hybridization in the genera Aeonium and Greenovia in La Gomera, Canary Islands

Author: Arango, Octavio

Source: Willdenowia, 53(3): 257-268

Published By: Botanic Garden and Botanical Museum Berlin (BGBM)

URL: https://doi.org/10.3372/wi.53.53305

BioOne Complete (complete.BioOne.org) is a full-text database of 200 subscribed and open-access titles in the biological, ecological, and environmental sciences published by nonprofit societies, associations, museums, institutions, and presses.

Your use of this PDF, the BioOne Complete website, and all posted and associated content indicates your acceptance of BioOne's Terms of Use, available at www.bioone.org/terms-of-use.

Usage of BioOne Complete content is strictly limited to personal, educational, and non - commercial use. Commercial inquiries or rights and permissions requests should be directed to the individual publisher as copyright holder.

BioOne sees sustainable scholarly publishing as an inherently collaborative enterprise connecting authors, nonprofit publishers, academic institutions, research libraries, and research funders in the common goal of maximizing access to critical research.



Principles governing F1 hybridization in the genera *Aeonium* and *Greenovia* in La Gomera, Canary Islands

Octavio Arango¹

Version of record first published online on 23 January 2024.

Abstract: A phytochorological and ecobiological study of the nine species, two subspecies and two varieties of the genera *Aeonium* and *Greenovia* in La Gomera is carried out, with special attention to F1 hybridization in nature. To the ten hybrids already described on the island, seven new hybrids were discovered, which are the subject of a separate publication. In all the known hybrids I determined whether the parents were sympatric or allopatric. The geographical coincidence of the parents was a sine qua non requirement for hybridization to occur in all the hybrids known so far, since pollen transfer by biotic and abiotic agents does not function between disjunct areas. The two prezygotic reproductive barriers that currently prevent interspecific hybridization in *Aeonium* or intergeneric hybridization between *Aeonium* and *Greenovia* are the geographic separation of populations and the different flowering times. The ability to survive and reproduce successfully of F1 hybrids in *Aeonium* and *Greenovia* is usually lower than that of the parent species and they generally disappear from the environment in the first filial generation without leaving offspring. The fundamental principles that govern F1 hybridization in La Gomera are extracted, and after verifying that they were also fulfilled in the other mountainous islands, they were synthesized in eight principles, which provide an updated vision of hybridization in the genus *Aeonium*.

Keywords: Aeonium, Canary Islands, chorology, ecology, Greenovia, hybridization, La Gomera, reproductive biology, sympatry

Article history: Received 13 April 2023; peer-review (1st round) completed 31 May 2023; received in revised form 4 and 18 July 2023; peer-review (2nd round) completed 22 August 2023; received in revised form 23 September 2023; accepted for publication 10 October 2023.

Citation: Arango O. 2024: Principles governing F1 hybridization in the genera *Aeonium* and *Greenovia* in La Gomera, Canary Islands. – Willdenowia 53: 257–268. https://doi.org/10.3372/wi.53.53305

Introduction

This research is based on the phytochorological study of Voggenreiter (1999), where he describes in detail the distribution of all species, subspecies, and hybrids that compose the genera *Aeonium* Webb & Berthel. and *Greenovia* Webb & Berthel. in La Gomera, and locates cartographically each of the taxa in grids of 1×1 km. In addition, he establishes the relationships of the species with the hybrids known up to that moment. His study shows how the taxa of both genera are distributed throughout practically the entire island, and more than half of them are endemic to La Gomera. The remarkable adaptability of these plants has allowed them to colonize the large variety of ecological niches on the island, making it an optimal natural laboratory for chorological, ecological, and reproductive biology studies.

The genus *Greenovia* was recognized as different from *Aeonium* a long time ago (1836–1840) based on significant morphological characters, such as flower parts or placentation. With the introduction of molecular studies two facts appeared. First, the *Greenovia* members formed a tight monophyletic cluster and, second, this cluster ap-

peared within the larger *Aeonium* group (Mes 1995; Jorgensen & Frydenberg 1999; Mort & al. 2002). Since it is desirable that a genus should be monophyletic, Mes suggested to include *Greenovia* as a section of *Aeonium* and therefore *Aeonium* was not paraphyletic anymore.

Both facts have been maintained as more detailed and robust trees were built, the latest and most complete being that of Messerschmid & al. (2023). Changing the name of *Greenovia* to *Aeonium* is certainly an easy way out. However, I think it is premature. Despite the many nucleotides included in the latter study, the tree was not well resolved. And several taxa appear in different branches when the trees are built with different criteria. This means that there still is not enough information to properly describe the group phylogeny. Until the branching order of the whole group is well established, changing the name of a genus is premature. And several authors have gone along this line (Lodé 2010; Schönfelder & Schönfelder 2018; Sauerbier & al. 2023).

Of all the genera that make up the rich and varied flora of the Canary Islands, *Aeonium* has the largest number of species and subspecies (Bramwell & Bramwell 2001; Acebes-Ginovés & al. 2009; Schönfelder & Schönfelder

1 Cl. Loreto 24–26, Escalera B, 4° 2ª, 08029 Barcelona, Spain. Author for correspondence: Octavio Arango, oja.oja@hotmail.com 2018; BIOTA 2023; Sauerbier & al. 2023) and it has generated the largest number of natural hybrids in the entire archipelago. In addition, intergeneric hybrids are formed frequently between *Aeonium* and *Greenovia*. The works of Praeger (1929, 1932) were the seminal papers on hybridization in the genus *Aeonium* in the Canary Islands. They were later joined by the publications of Lems (1960), Liu (1989), Marrero (1992), Voggenreiter (1999), and Jorgensen & Olsen (2001). Since then, very little progress has been made in understanding the eco-biological mechanisms governing hybridization in the genera *Aeonium* and *Greenovia*, with most subsequent publications focusing only on the taxonomic aspects of hybrids.

Marrero (1992) established three different groups of plants in the Canary flora according to the hybridization process, depending on the level of differentiation that the taxa had reached during the geographic isolation to which they had been subjected, and the capacity to hybridize when they coincide again. In addition, he analysed the effectiveness of the reproductive barriers acquired during isolation and the viability of the hybrids they generate. The first group includes allopatric species that normally hybridize when they meet again due to various factors, such as Cheirolophus Cass., Crambe L. and Limonium Mill. The second group includes isolated or sympatric species, in which hybridization is more or less viable depending on the taxa, such as Echium L., Micromeria Benth. and Sideritis L. And the third group, the most numerous, consists of sympatric species but with sporadic hybridizations and generally unviable offspring such as Aeonium, Argyranthemum Webb and Sonchus L.

To advance knowledge of this difficult and complex group of plants, this research project has three main objectives: (1) to continue the chorological study of Voggenreiter (1999) on hybridization in the genera *Aeonium* and *Greenovia* and to validate his results; (2) to synthesize the fundamental principles that govern F1 hybridization in the genera *Aeonium* and *Greenovia* in La Gomera, and to explore if they are fulfilled in the other mountainous islands in the Canary Archipelago; (3) to identify new hybrids in their habitat and to study the particularities in which they occur in nature.

The overall results of this research have been divided into two articles. In the present paper, the results related to the natural history of F1 hybrids and the fundamental principles governing hybridization in the genera *Aeonium* and *Greenovia* are presented. In the second article (Arango 2023c), the taxonomic results are provided, and seven new hybrids discovered in the fieldwork are described and illustrated.

Material and methods

Based on the areas of distribution of the species that make up the genera *Aeonium* and *Greenovia* provided by Voggenreiter (1999), fieldwork was carried out in

many parts of the island of La Gomera. The search for new hybrids was carried out according to the targeted search method developed previously (Arango 2021c). The method consists of three steps: (1) the distribution of all the species in the island was taken from Voggenreiter (1999) and later authors; (2) for each pair of species with overlaps in their areas of distribution, the literature was searched to check whether hybrids between them had been described; (3) for those species pairs that did not have hybrids described, the areas of overlap were extensively searched for potential hybrids.

Data were collected on the morphology, chorology, ecology, and reproductive biology of the taxa that make up both lineages and their hybrids, with special attention to the identification of nothotaxa unknown to science (see Appendix 1). New nothotaxa were recognized by being found living inter parentes and by the distinctive features they presented, according to the critical morphological characters with taxonomic value established by Praeger (1932) and Liu (1989) for the genera *Aeonium* and *Greenovia*.

For the hybridization experiments, cuttings of the different species were grown in the garden until they bloomed. The flowers that were going to act as receptors were emasculated while still in bud stage, and the stigmas covered to prevent unwonted pollination. They were left 24–48 hours for the carpels to become receptive. Pollen from the donor plants was then collected and placed on the stigma. Finally, the pollinated flowers were covered again to prevent further pollinations. After fructification, seeds were collected and planted in the next fall. The seeds were processed and planted according to the technique described by Schulz (2007).

Finally, each of the observational data collected in the field and the laboratory was assigned a score of 0, 1 or 2 (0: the parameter does not exist in the hybrids; 1: it exists but is not constant; 2: it exists and is shared by all natural hybrids), which allowed me to know the common characters in natural hybrids of *Aeonium* and *Greenovia*, and from this information I synthesized the principles governing the phenomenon of hybridization in La Gomera. Then, the analysis of the information stored in the database obtained from the observation of numerous natural hybrids in all the islands and my publications confirm that these principles can be extrapolated to the rest of the mountainous islands of the Canary Archipelago (Arango 2015, 2016a, 2016b, 2017, 2019a, 2019b, 2019c, 2021a, 2021b, 2021c, 2023a).

Results and Discussion

Hybrids in La Gomera

There are 13 taxa of *Aeonium* and *Greenovia* in La Gomera (Fig. 1, Table 1). Therefore, if all could hybridize among them, there would be 78 potential hybrids.

	A. appendiculatum	A. canariense subsp. <i>latifolium</i>	A. castello-paivae	A. decorum	A. decoum var. alucense	A. gomerense	A. holochrysum	A. holochrysum var. rubrolineatum	A. saundersii	A. sedifolium	A. spathulatum	A. lindleyi subsp. viscatum	G. diplocycla
Aeonium appendiculatum													
A. canariense subsp. latifolium	P												
A. castello-paivae	P	D											
A. decorum	D	D	D										
A. decorum var. alucense	X	Х	X	Р									
A. gomerense	X	Р	D	N	X								
A. holochrysum	X	X	Р	Р	X	X							
A. holochrysum var. rubrolineatum	Р	Р	Р	Р	X	X	Р						
A. saundersii	Р	D	Р	N	X	X	X	Р					
A. sedifolium	X	X	X	P	P	X	X	X	X				
A. spathulatum	Р	Р	Р	Р	X	Р	Р	D	N	X			
A. lindleyi subsp. viscatum	X	D	D	N	Р	P	Р	Р	P	Р	X		
Greenovia diplocycla	Р	Р	N	D	X	Р	Р	Р	N	X	N	Р	

= parents sympatric; = parents allopatric; D = hybrids previously described; N = hybrids newly recognized; P = hybrids theoretically possible; X = hybrids not theoretically possible.

Fig. 1. Matrix of all potential crosses among taxa of genera *Aeonium* and *Greenovia* living in La Gomera. For each cross, green background indicates sympatry and, therefore, crossing is theoretically possible, while orange background indicates allopatry and, therefore, crossing is not theoretically possible.

However, as I will argue below, hybridization in nature will only occur when the two parental species coexist in sympatry. Therefore, out of the 78 possible crosses, 27 are not expected to occur due to allopatric distribution of the parents, while 51 potentially exist. Up to now, 10 have been described in the literature and seven additional ones are published in Arango (2023c) (Table 2).

Of the seven new hybrids contributed by Arango (2023c), three are intergeneric between *Aeonium* and

Greenovia. The only representative of the genus Greenovia in La Gomera is G. diplocycla Webb ex Bolle (Praeger 1929, 1932; Bramwell & Bramwell 2001; Lodé 2010). However, it is a fairly frequent taxon that is distributed throughout almost the entire island and grows intermixed with different species of the genus Aeonium. Out of the ten hybrids previously described by other authors, only one was intergeneric, ×Greenonium laxiflorum J. M. Macarrón & Bañares. The targeted and systematic

search method I have been using favoured the finding of three new intergeneric hybrids of *G. diplocycla* with *A. castello-paivae* Bolle, *A. saundersii* Bolle, and *A. spathulatum* (Hornem.) Praeger (Table 2). Of these, ×*Greenonium cabrerae* A. Santos had already been described in the island of La Palma (Santos 1983). In La Gomera, I found a vicariant nothovariety of the latter in the vicinity of Roque de Ojila, a site where the parents are sympatric.

In a recent study, Messerschmid & al. (2023), in an extensive analysis of the genus Aeonium s.l. (including Greenovia) concluded that "those islands with the longest history of habitation by Aeonium had the lowest percentages of co-occurring and hybridizing taxon pairs compared to islands where Aeonium arrived later". They based this conclusion on their counts of taxa and hybrids in different islands resulting in Gran Canaria (more recently colonized by the genus) having a larger percent of hybrids than Tenerife (colonized earlier). Unfortunately, it appears that these authors finished their sampling in 2019 and have not considered all the novel taxa validly published of the genera Aeonium (two species, one subspecies, and ten nothospecies) and Greenovia (two species and one nothospecies) since then (Arango 2019a, 2019b, 2019c, 2019d, 2021a, 2023a, 2023b). When all the available data are considered, the numbers of hybrids known in every island are the following: Tenerife 30, La Palma 18, La Gomera 17, Gran Canaria nine and El Hierro six. Moreover, Tenerife (21 taxa and 30 hybrids) has about three times more taxa than Gran Canaria and three times more hybrids (eight taxa vs nine hybrids). Since the number of known hybrids is simply proportional to the number of taxa on an island, the more complex assumption of age of colonization is unnecessary and does not reflect reality.

Experimental crosses

Controlled experiments in the laboratory were essential to understand how F1 hybridization works in the genera Aeonium and Greenovia. Whenever mature seeds of the new F1 hybrids found in nature were available, a germination test was performed in the laboratory to determine the fertility of the seeds. In addition, to find out if genetic barriers prevented hybridization between species of the genera Aeonium and Greenovia, different artificial crosses were made between endemic species from different Canary Islands, Madeira, and Cabo Verde (Appendix 2), and all produced fertile seeds and viable embryos. Unfortunately, the two East African taxa, A. leucoblepharum Webb ex A. Rich. and A. stuessyi H.-Y. Liu, have not flowered in cultivation and similar crosses could not be attempted with Macaronesian species. However, I am not aware of any publication reporting reproductive incompatibility between species of these genera. Therefore, in principle, there are no genetic barriers to hybridization among any species and the seeds of the F1 hybrids are fertile.

Table 1. Total taxa and potential and described hybrids in La Gomera.

Hybrids	Number	Percent
Total number of species and subspecies	13	
Total number of possible crosses (all against all)	78	100
Hybrids not expected due to allopatry	27	34
Hybrids potentially existing due to sympatry	51	65
Described hybrids	17	21
Hybrids described previously	10	13
Hybrids described in Arango (2023c)	7	9

The need for sympatry

As can be seen in Fig. 1, all the described hybrids fulfil the condition that the parental species are sympatric. This can also be seen in Messerschmidt & al. (2023: fig. 5). To the best on my knowledge, no hybrid has been described between allopatric species. This can be extrapolated to all the islands. Moreover, this observation appears in most of the descriptions of other authors, in which the comment "found living among the parents" is constantly included. These findings support the hypothesis that parent sympatry is a sine qua non condition for F1 hybridization to occur in the genera *Aeonium* and *Greenovia* in the Canary Islands.). This will constitute the first principle (see below).

One may ask why is sympatry necessary if both pollinating insects and the wind, in principle, seem to have enough capacity to transport pollen to distant places and, as shown in the previous section, there are no barriers to hybridization. Cross-pollination between species of the genera Aeonium and Greenovia can be carried out by both biotic and abiotic agents, given that most Canary Island plants, including Aeonium and Greenovia, are considered to have a generalist pollination syndrome. Among the former, insects, lizards, bats and birds have been shown to be active (Bernardello & al. 2001; Olsen & al. 2002; Valido & al. 2004; Esfeld 2009; Padrón & al. 2009; Traveset & al. 2009; Nelson 2010; Severio & Rodríguez-Rodríguez 2011; Ortega-Olivencia & al. 2012; Arango 2013). The wind, in turn, usually only produces the effective dispersal of pollen between individuals located at a short distance (Levin & Kerster 1974). Although some insects can travel several kilometres in their foraging activities and wind can transport pollen over long distances (Heinrich 1979; Roubick 2001; Ollerton & al. 2011), the results of this work show that pollen transfer through these vectors does not work between allopatric species. As mentioned above, all F1 hybrids in La Gomera always occurred when the geographical areas of distribution of the parents coincided (Arango 2023c). This was also the case for the 28 new hybrids studied in other islands of the archipelago (T, GC, P, H) (Arango 2015, 2016a, 2016b, 2017, 2019a, 2019b, 2019c, 2021a, 2021b, 2021c, 2023a).

Table 2. Aeonium and Greenovia hybrids discovered so far in La Gomera.

Hybrids	Parent 1	Parent 2	Year
Hybrids previously described			
A. ×aguajilvense Bañares	A. castello-paivae Bolle	A. gomerense (Praeger) Praeger	1996
A. ×beltranii Bañares	A. canariense subsp. latifolium (Burchard) Bañares	A. decorum Webb ex Bolle var. decorum	1986
A. ×bravoanum Bramwell & G. D. Rowley ex Heath	A. castello-paivae	A. lindleyi subsp. viscatum (Bolle) Bañares	1992
A. ×castellodecorum Bañares	A. castello-paivae	A. decorum var. decorum	1986
A. $\times castelloplanum$ Bramwell & G. D. Rowley ex Heath	A. canariense subsp. latifolium	A. castello-paivae	1992
A. ×holospathulatum nothovar. sanchezii (Bañares) Bañares	A. arboreum var. rubrolineatum (Svent.) H. Y. Liu	A. spathulatum (Hornem.) Praeger	1996
A. ×perezii Bañares	A. appendiculatum Bañares	A. decorum var. decorum	1990
A. ×sancti-sebastianii Bramwell & G. D. Rowley ex Heath	A. canariense subsp. latifolium	A. saundersii Bolle	1992
A. ×vegamorae Bramwell & G. D. Rowley ex Heath	A. canariense subsp. latifolium	A. lindleyi subsp. viscatum	1992
×Greenonium laxiflorum Macarrón & Bañares	A. decorum var. decorum	Greenovia diplocycla Webb ex	2007
New hybrids (Arango 2023c)		Bolle	
A. ×dendroides O. Arango	A. decorum var. decorum	A. saundersii	2023
A. ×pendulum O. Arango	A. saundersii	A. spathulatum	2023
A. ×praedictum O. Arango	A. decorum var. decorum	A. gomerense	2023
A. ×rhombifolium O. Arango (1)	A. decorum var. decorum	A. lindleyi subsp. viscatum	2023
$\times Greenonium\ cabrerae$ nothovar. $ojilense$ O. Arango $^{\mbox{\tiny (1)}}$	A. spathulatum	Greenovia diplocycla	2023
×Greenonium haeckelii O. Arango	A. castello-paivae	G. diplocycla	2023
×Greenonium lajense O. Arango	A. saundersii	G. diplocycla	2023

⁽¹⁾ Only the vegetative part is known.

Possibly the large inflorescences of Aeonium, formed by numerous brightly coloured flowers, extraordinarily nectariferous and polliniferous, exert an effect of attraction and loyalty to the pollinator, therefore preventing them from having to move to other areas in search of the reward they are looking for (Rademaker & De Jong 1998; Santos & al. 2004; Ares 2019). However, other authors have found low pollen transfer between pairs of sympatric Aeonium species and explain their results by a specific behaviour of pollinators during foraging, such as the floral preference of a given species for the reward it receives (Esfeld & al. 2009). They agree with Campbell & al. (2002) that the quality and quantity of pollen transfer are primarily affected by pollinator behaviour rather than variable flower characteristics such as colour, scent, and reward regime.

Finding the missing natural hybrids

The species richness and taxonomic complexity of the genus *Aeonium*, together with its formidable hybridogenic capacity, has generated the largest number of natural hybrids in the entire Canary Island flora, with at least 80 nothotaxa known to date. As mentioned above, there are still many theoretically possible hybrids to be discov-

ered in La Gomera (and in other islands), with at least 34 crosses between species that share the same distribution areas. However, it is becoming increasingly difficult to find new hybrids on the island, possibly due to wellknown factors such as (1) the progressive anthropogenization and fragmentation of the natural environment; (2) the serious damage caused by the numerous introduced herbivores (goats and sheep, both domestic and feral) that graze uncontrolled in many parts of the island; (3) the devastating fires that the island has suffered in recent decades; and (4) the negative effects that climate change is causing to the island environment (Caujapé-Castells & al. 2010; Arango 2021d). These factors have led to a critical conservation status (CR) for endemics such as A. gomerense, A. saundersii and A. decorum Webb ex Bolle var. alucense (Gómez 1996; Moreno 2008, 2011; IUCN 2012); and if effective solutions are not adopted in the short term, other taxa such as A. appendiculatun Bañares, whose distribution area was severely affected by the 2012 fire, probably will have to be added to the list of endangered species.

Generally, recognition of interspecific hybrids in the first filial generation (F1) in the genus *Aeonium* or intergeneric hybrids between *Aeonium* and *Greenovia* is easy, since they are usually found intermixed with the parents and usually express morphological characters of both. However, determining hybrids in the F2, F3, F4, and successive generations is extremely complex since the characters they express hide inherited traits from the parents. Likewise, the introgressive gene flow caused by backcrossing with the parental species generates progeny with very little morphological differentiation regarding the parents, with which they are generally confused.

Three particular cases illustrate the vagaries of hybrid discovery. An interesting result of this work has been to actually confirm how the distribution areas of two species that originally did not coincide spatially, Aeonium decorum var. decorum Webb ex Bolle and A. gomerense (Praeger) Praeger, have been brought into contact by human action. This has occurred due to the phenomenon that Voggenreiter (1999) called "anthropogenic translocation following the roads" This brought the populations of both parents closer together due to the progressive colonization of the roadside by A. decorum; and, as predicted by the German botanist, hybridization between them has eventually occurred. The new nothotaxon, A. xpraedictum Arango, was found on one of the edges of the GM-1 road near La Degollada de San Sebastián, the site where the areas of the parents are close to each other. It is a well-known fact that roadsides in the Canary Islands constitute an optimal microhabitat for the propagation of numerous endemism (Delgado & al. 2007; Hengstum 2012).

On the other hand, the cross between A. decorum and A. sedifolium (Webb ex Bolle) Pithard & Proust, described in Tenerife as A. xpuberulum Bañares, has not yet been found in La Gomera. However, it is not ruled out that it may exist in the inaccessible rocky areas of the Puntallana Special Nature Reserve in the east of the island, where both species grow intermingled (Casanova 1990; Voggenreiter 1999). In this case, extremely arid conditions (del Arco & Rodríguez, 2019) may not be favourable for the development of the hybrids, as I have observed that a very high proportion of Aeonium seedlings that germinated during the rainy season do not survive very dry and hot summers. At any rate, finding this hybrid in the western part of Tenerife was unexpected, since A. decorum is extraordinarily rare in Tenerife (Ceballos & Ortuño 1976; Hernández 1998). Perhaps the environmental conditions of Masca in Tenerife are more favourable to the subsequent development of the progeny.

Finally, in a few cases a hybrid was found away from the parental populations. Usually, the parents were found upwind and/or uphill from the hybrid, suggesting that the seeds were blown away from the parent populations by the wind. The three cases found were *Aeonium* ×*bravoanum* (*A. castello-paivae* × *A. lindleyi* subsp. *viscatum* (Bolle) Bañares in La Gomera; *A. xbornmuelleri* (*A. canariense* subsp. *canariense* (L.) Webb & Berthel. × *A. urbicum* subsp. *urbicum* (C. Sm. ex Hornem.) Webb & Berthel. in Tenerife; *A. canariense* subsp. *canariense* × *A. liui* Arango in Tenerife.

Persistence of the hybrids

Reproduction of natural F1 hybrids by vegetative propagules is rare in *Aeonium* and *Greenovia* and only occurs when the parents are stoloniferous such as *A. simsii* (Sweet) Stearn or *A. cuneatum* Webb & Berthel. In my exhaustive explorations, I have only found two natural hybrids with this property, *A. ×stoloniferum* Arango in Gran Canaria and *A. ×acebesii* Arango in Tenerife (Arango 2019a, 2021c).

Occasionally, on the five mountainous islands of the Canary archipelago (T, P, G, GC, H), I have found some Aeonium forming small hybrid swarms composed of individuals with the ability to reproduce by fertile seeds, possibly obtained by autogamy, backcrossing or introgressive crosses with other species, without ruling out apomixis, which has not been studied in the genus Aeonium (Fig. 2). The number of different parental pairs producing these F2 and successive generations does not exceed eight crosses, so they can be considered an exception. Rarer are stable hybrids with the capacity to colonize larger geographical areas, an occurrence which I have only observed on the island of La Palma, where the phenomenon of hybridization is very complex. An example of this type of hybridization can be observed in El Time, a site on the west of La Palma, where it is difficult to find morphologically representative specimens of A. davidbramwellii H. Y. Liu, and most of the population is composed of hybrid specimens. Probably, this population has reached stability by successive backcrosses with the parental species or by introgressive crosses with other species with which it shares its distribution area This phenomenon is well known in other taxonomically complex groups such as Biscutella L. (Cruciferae). Some authors have chosen to consider these stabilized hybrids as species of hybridogenic origin in the absence of data that would place them definitively in the nothospecies rank (Mateo 2023).

Principles governing hybridization in the genera *Aeonium* and *Greenovia*

Analysis of the results of this work and those of Voggenreiter (1999), plus studies of numerous natural hybrids on other mountainous islands of the archipelago (GC, H, P, T), and methodical observations (Arango 2015, 2016a, 2016b, 2017, 2019a, 2019b, 2019c, 2021a, 2021b, 2021c, 2023a) have provided essential information to better understand the natural history of F1 hybrids of the genera *Aeonium* and *Greenovia*, which I have synthesized into the following eight principles. The principles governing hybridization in the F1 generation in the genera *Aeonium* and *Greenovia* have been expressed as principles and not as laws because, in natural sciences, laws are universal, while principles are subject to biological evolution and changes in the natural environment and may have exceptions (Elgin 2003; Martínez & al. 2017).



Fig. 2. Seedbed of a hybrid swarm of *Aeonium ×junoniae* Bramwell & G. D. Rowley ex Heath from La Palma, obtained with seeds from nature, which confirms occasional fertility of some hybrids in F2 generation or later.

First principle — F1 hybridization in nature only occurs between species that share the same geographic area, or at least where the areas of distribution of both parents come into contact at some point. Pollen transfer through biotic and abiotic agents does not work between disjunct areas.

Second principle — When two species that share the same geographical area have overlapping blooming phenology, F1 hybridization always ends up occurring between them.

Third principle — Crosses between the different species produce fertile seeds that give rise to F1 hybrids since there are no genetic barriers between them to prevent this. However, the fertility of F1 hybrids is quite variable, ranging from absolute sterility to complete fertility.

Fourth principle — Seeds of F1 hybrids usually germinate where they originate, but given their small size they are, in rare cases, dispersed by the wind and germinate between hundreds of meters and a few kilometres away from the site where they were formed.

Fifth principle — The evolutionary aptitude of F1 hybrids in nature, understood as the capacity to survive or reproduce successfully, is usually low or null, and they generally disappear from the natural environment in the F1 generation. However, some crosses do generate fertile hybrids with a certain stability and ability to form small swarms, but they are the exception.

Sixth principle — The number of hybrid individuals generated by the different crosses in nature does not depend on the size of the populations of the parental species or on the taxonomic affinity between them, such as belonging to the same section or to the same molecular clade. The most common is that they occur in a low frequency.

Seventh principle — Generally, F1 hybrids express the morphological characters inherited from both parents in similar proportions. But in some cases, the alleles of one parent are dominant over those of the other, making the offspring more closely resemble the dominant parent.

Eighth principle — Despite the low fertility of the F1 hybrids, back-crossings of these with the parental species occur in nature sporadically. When this occurs, it produces genic introgression and morphological convergence between the progeny and the parents.

From these principles, it can be deduced that the only two effective prezygotic barriers that prevent intraspecific F1 hybridization in the genus Aeonium and intergeneric hybridization between Aeonium and Greenovia in the Canary Islands are the geographical separation of the populations and the different flowering times. Probably, the time they have been separated has not been long enough to establish such reproductive barriers. Moreover, the efficacy of different flowering times as a reproductive barrier is decreasing due to climate change effects in the Canary Islands. Nowadays, it is increasingly common to observe earlier flowering, lengthening of the flowering period or extemporaneous flowering (unpublished data of the author). This fact is evident when comparing the phenological data I have collected since 2010 with the flowering dates provided by Liu (1989), many of them obtained from historical herbarium sheets. These changes in floral phenology will make the appearance of new natural hybrids possible.

Final comments

Although individual hybrids in the genera *Aeonium* and *Greenovia* usually disappear from nature in the first filial generation, they can produce gene dilution in the species with which they coexist, as pollen viability can reach 80–90% in some hybrids (Liu 1989). Hence the importance of recognizing hybrids in nature, describing them, and depositing the types in herbarium collections. In addition, they should be included in all Canary databases, as is done by the website Endémicas Canarias (2023), because they can contribute to a better understanding of the evolutionary history and phylogeny of the flora of the Canary Islands.

A better understanding of the chorology, ecology, and reproductive biology of the genera *Aeonium* and *Greenovia* may favour the development of new conservation strategies to ensure the long-term survival of these plants. Increasingly, importance is being given to the detrimental effects of hybrids on the species due to the loss of genetic identity, which compromises their reproduction and makes them more vulnerable to various threats in the natural environment. It is also well known that genes, evolution, and environmental adaptation always go together (Levin & al. 1996; Francisco-Ortega & al. 2000; Brilhante & al. 2021). Perhaps understanding these phenomena can help to explain why the Canary

Island Network of Protected Natural Spaces (Law 12/1994 of 19 December 1994), after three decades of operation, has failed to protect *Aeonium* species, which are in a critical state of conservation category CR (IUCN 2012), since they are all found within protected areas, and none have recovered, neither in number of individuals nor in area covered. It would be interesting to find out why these areas have not achieved such a basic objective as in situ conservation of threatened species (Chape & al. 2005; Reyes-Betancort & al. 2008; Moreno 2015; IUCN 2012).

Acknowledgements

I thank Águedo Marrero Rodríguez, Department of Plant Systematics and LPA Herbarium, Jardín Botánico Viera y Clavijo, for his interesting comments and masterly teachings on the phenomenon of hybridization in the Canary Island flora. I also thank Carlos Pedrós-Alió, professor and research scientist at CSIC in Madrid for his time spent on critical reading of the manuscript, as his comments improved this paper considerably, and for his "insistence" on where and how this paper should be published. I am grateful to Cristina González Montolongo of the Department of Botany, Ecology and Plant Physiology of the University of La Laguna for the access she has always given me to consult the sheets of the TFC Herbarium, the most complete collection of natural hybrids of the genera Aeonium and Greenovia the world. Finally, an anonymous reviewer is thanked for providing comments on two earlier versions of this article. Likewise, I thank the editor for his careful revision and useful comments.

References

- Acebes-Ginovés J. R., León-Arencibia M. C., Rodríguez-Navarro L., del Arco-Aguilar M., García-Gallo A., Pérez de Paz P. L., Rodríguez-Delgado O., Martín-Osorio V. E. & Wildpret de la Torre W. 2010: *Pteridophyta*, *Spermatophyta*. Pp. 119–172 in: Arechavaleta M., Rodríguez S., Zurita N. & García A. (coord.), Lista de especies silvestres de Canarias. Hongos, plantas y animales terrestres. 2009. Santa Cruz de Tenerife: Gobierno de Canarias.
- Arango O. 2013: Consideraciones sobre la polinización ornitófila del bejeque *Aeonium arboreum* ssp. *holo-chrysum* en el Parque Nacional de La Caldera de Taburiente (La Palma). – Makaronesia 15: 98–105.
- Arango O. 2015: *Aeonium x claperae (Crassulaceae)*, un nuevo híbrido natural de Tenerife (Islas Canarias). Vieraea **43:** 329–334.
- Arango O. 2016a: Confirmación experimental del rango de especie para *Aeonium mascaense* Bramwell (*Crassulaceae*). Bot. Macarones. **29:** 25–34.

- Arango O. 2016b: *Aeonium* x *wildpretii* Bañares (*Crassulaceae*): descripción de la flor, nueva localidad y actualización taxonómica. El Hierro, Islas Canarias. Bot. Macarones. **29:** 83–89.
- Arango O. 2017: *Aeonium x monteaquaense*, *A. x afurense* y *A. x marreroi (Crassulaceae*), tres nuevas notoespecies de las islas Canarias. Vieraea **45:** 267–284. https://doi.org/10.31939/vieraea.2017.45.15
- Arango O. 2019a: Descripción de cuatro nuevas notoespecies del género *Aeonium* (*Crassulaceae*) en las Islas Canarias y aportaciones corológicas y nomenclaturales al género. – Vieraea **46:** 115–146. https:// doi.org/10.31939/vieraea.2019.46.tomo01.06
- Arango O. 2019b: *Aeonium liui (Crassulaceae)*: una nueva especie de Tenerife, islas canarias. Bot. Macarones. **30:** 7–22.
- Arango O. 2019c: Descripción de *Aeonium* x *exsul* (*Crassulaceae*). Un viejo nombre para un nuevo híbrido natural de Tenerife (Islas Canarias). Bot. Macarones. **30:** 143–152.
- Arango O. 2019d: Descripción de *Aeonium urbicum* ssp. *boreale* (*Crassulaceae*), una nueva subespecie de uno de los endemismos más representativos de Tenerife, Islas Canarias. Vieraea **46:** 11–28. https://doi.org/10.31939/vieraea.2019.46.tomo01.02
- Arango O. 2021a: *Greenovia millennium (Crassulaceae)*: una nueva especie y sus híbridos. Tenerife, Islas Canarias. Bot. Macarones. **31:** 11–32.
- Arango O. 2021b: Confirmación experimental del rango de especie de *Aeonium mascaense* (*Crassulaceae*). (Addenda). Bot. Macarones. **31:** 33–40.
- Arango O. 2021c: Nuevas aportaciones al género *Aeonium* (*Crassulaceae*): descripción de cinco notoespecies nuevas y ampliación corológica. Islas Canarias. Vieraea 47: 79–114. https://doi.org/10.31939/vieraea.2021.47.07
- Arango O. 2021d: El «mal de las hojas moradas» en el género *Aeonium (Crassulaceae*): etiología, síntomas y evolución. La Gomera, islas Canarias. Rev. Acad. Canar. Cienc. **33:** 21–34.
- Arango O. 2023a: Descripción de *Aeonium* x *gulliveri* (*Crassulaceae*), un nuevo y singular híbrido de La Palma. Bot. Macarones. **32:** 167–174.
- Arango O. 2023b: *Greenovia ignea* y *Aeonium calderense* (*Crassulaceae*): dos nuevas especies de La Palma, Islas Canarias. Bot. Macarones. **32:** 145–166.
- Arango O. 2023c: Contribution to the knowledge of the natural hybrids of the genera *Aeonium* and *Greenovia* (*Crassulaceae*) in La Gomera, Canary Islands. Collect Bot. **42**(e008). https://doi.org/10.3989/collectbot .2023.v42.008
- Ares R. 2019: Los atractivos de las flores para los polinizadores.
 En: La conducta de las plantas: etología botánica.
 Buenos Aires: Editorial Fundación de Historia Natural Félix de Azara.
- Bernardello G., Anderson G. J., Stuessy T. F. & Craw-Ford D. J. 2001: A survey of floral traits, breeding

systems, floral visitors, and pollination systems of the angiosperms of the Juan Fernández Islands (Chile).

– Bot. Rev. **67:** 255–308. https://doi.org/10.1007/BF02858097

- Biota 2023: Banco de datos de biodiversidad de Canarias.
 Published at https://www.biodiversidadcanarias.es/biota/especies [accessed various times in 2021 and 2023].
- Bramwell D. & Bramwell Z. 2001: Flores silvestres de las Isla Canarias. 4º edición. Madrid: Editorial Rueda.
- Brilhante M., Roxo G., Catarino S., Dos Santos P., Reyes-Betancort A., Caujapé- Castells J., Meneses Sequeira M., Talinhas P. & Romeiras M. M. 2021: Diversification of *Aeonium* species across Macaronesian archipelagos: correlations between genome-size variation and their conservation status. Frontiers Ecol. Evol. 9(607338). https://doi.org/10.3389/fevo.2021.607338
- Campbell D. R., Waser N. M. & Pederson G. T. 2002: Predicting patterns of mating and potential hybridization from pollinator behavior. – Amer. Naturalist 159: 438–450. https://doi.org/10.1086/339457
- Casanova J. 1990: Sobre la presencia de *Aeonium sedi*folium (*Crassulaceae*) en la isla de La Gomera. – Vieraea **19:** 139–140.
- Caujapé-Castells J., Tye A., Crawford D. J., Santos-Guerra A., Sakai A., Beaver K., Lobin W., Florens F. B. V., Moura M., Jardim R., Gómes I. & Kueffer C. 2010: Conservation of oceanic island floras: present and future global challenges. Perspect. Pl. Ecol. Evol. Syst. 12: 107–129. https://doi.org/10.1016/j.ppees.2009.10.001
- Ceballos L. & Ortuño F. 1976: Estudio sobre la vegetación y flora forestal de las Canarias Occidentales. Tenerife: Ed. Excmo. Cabildo Insular.
- Chapes S., Harrison J., Spalding M. & Lysenko I. 2005: Measuring the extent and effectiveness of protected areas as an indicator for meeting global biodiversity targets. Philos. Trans., Ser. B. **360:** 443–455. https://doi.org/10.1098/rstb.2004.1592
- del Arco M. & Rodríguez O. 2019: Vegetation of the Canary Islands. Plant and vegetation **16.** Utrecht: Springer. https://doi.org/10.1007/978-3-319-77255-4
- Delgado J. M., Arroyo N. L. & Fernández-Palacios J. R. 2007: Efectos del borde de las carreteras sobre la temperatura, la luz, la cobertura del dosel y la altura del dosel en bosques de laurisilva y pinos (Tenerife, Islas Canarias). – Plan Urbano Paisajístico 81: 328–340.
- Elgin M. 2003: Biology and a priori laws. Philos. Sci. **70:** 1380–1389. https://doi.org/10.1086/377415
- Endémicas Canarias 2023: Especies vegetales de Canarias. Published at https://endemicascanarias.com/index.php/es/ [accessed 2 Feb 2023].
- Esfeld K., Koch M. A., van der Niet T., Seifan M. & Thiv M. 2009: Little interspecific pollen transfer despite overlap in pollinators between sympatric *Aeonium* (*Crassulaceae*) species pairs. Flora **204:** 709–717. https://doi.org/10.1016/j.flora.2008.10.002

- Francisco-Ortega J., A. Santos-Guerra A., Kim S. C. & Crawford D. J. 2000: Plant genetic diversity in the Canary Islands: a conservative perspective. Amer. J. Bot. **87:** 909–19. https://doi.org/10.2307/2656988
- Gómez C., del Arco M., Acebes J. R., Bañares A., Beltrán
 E., Betancort M. J., Cabrera M. A. & Fernández M.
 1996: Libro rojo de especies vegetales amenazadas de las Islas Canarias. Santa Cruz de Tenerife: Gobierno de Canarias.
- Hengstum T. van, Lachmuth S., Oostermeijer J. G. B.,
 Dennijs H. C. M., Meirmans P. G. & van Tienderen
 P. H. 2012: Human-induced hybridization among congeneric endemic plants on Tenerife, Canary Islands. Pl. Syst. Evol. 298: 1119–1113. https://doi.org/10.1007/s00606-012-0624-6
- Heinrich B. 1979: Bumblebee economics. Cambridge & London: Harvard University Press.
- Hernández E. 1998: Nota sobre la presencia de *Aeonium decorum* Webb *ex* Bolle (*Crassulaceae*) en la isla de Tenerife. Anales Jard. Bot. Madrid **56:** 158–159.
- IUCN 2012: IUCN Red List categories and criteria. Version 3.1. Second edition. Prepared by the IUCN Species Survival Commission. – Gland & Cambridge: IUCN. https://www.iucnredlist.org/resources/categories-and-criteria
- Jorgensen T. H. & Olsen J. M. 2001: Adaptative radiation of island plants: evidence from *Aeonium (Crassulaceae)* of the Canary Islands. Perspect. Pl. Ecol. Evol. Syst. **4:** 29–42. https://doi.org/10.1078/1433-8319-00013
- Lems K. 1960: Botanical notes on the Canary Islands II. The evolution of plant forms in the islands: *Aeonium*. Ecology **41:** 1–17. https://doi.org/10.2307/1931934
- Levin D. A., Francisco-Ortega J. & Jansen R. K. 1996: Hybridization and extinction of rare plant species. Conservation Biol. **10:** 10–16. https://doi.org/10.1046/j.1523-1739.1996.10010010.x
- Levin D. A. & Kerster H. W. 1974: Gene flow in seed plants. – Evol. Biol. 7: 139–220. https://doi.org /10.1007/978-1-4615-6944-2_5
- Liu H.-Y. 1989: Systematics of *Aeonium* (*Crassulaceae*).

 Special Publ. Natl. Mus. Nat. Sci. Taiwan 3.
- Lodé J. 2010: Plantas suculentas de las Islas Canarias. Guía de identificación fácil. – Santa Cruz de Tenerife: Publicaciones Turquesa.
- Marrero Á. 1992: Evolución de la Flora Canaria. Pp. 62–63 in: Kunkel G. (ed.), Flora y vegetación del Archipiélago Canario. Tratado florístico de Canarias, Primera parte. Las Palmas de Gran Canaria: Editora Regional de Canarias.
- Martínez M., García E. & Bernal C. 2017: Reduccionismo, leyes naturales y complejidad: diferentes estrategias de investigación y explicación científica. Sci. Stud. **15:** 243–262. https://doi.org/10.11606/51678 -31662017000200003

- Mateo G. 2023: Sobre el género *Biscutella* L. (*Cruciferae*) en la Cordillera Ibérica. Flora Montiber. **85:** 16–26.
- Mes T. H. M. 1995: Phylogenetic and systematic implications of chloroplast and nuclear spacer sequence variation in the Macaronesian *Sempervivoidea* and related *Sedoideae* (*Crassulaceae*). Pp. 30–44 in: Hart H. 't & Eggli U. (ed.), Evolution and systematics of the *Crassulaceae*. Leiden: Backhuys.
- Messerschmid T. F. E., Avrahamczyk S., Bañares Baudet A., Brilhante M. A., Eggli U., Hühn P., Kadereit J. W., dos Santos P., de Vos J. M. & Kadereit G. 2023: Inter- and intra-island speciation and their morphological and ecological correlates in *Aeonium (Crassulaceae)*, a species-rich Macaronesian radiation. Ann. Bot. **131:** 697–721. https://doi.org/10.1093/aob/mcad033
- Moreno J. C. (coord.) 2008: Lista Roja 2008 de la flora vascular española. Dirección General de Medio Natural y Política Forestal (Ministerio de Medio Ambiente, y Medio Rural y Marino, y Sociedad Española de Biología de la Conservación de Plantas). Madrid: Gobierno de España.
- Moreno J. C. 2011: Lista roja de la flora vascular española. Adenda 2010. Actualización con los datos de la adenda 2010 al atlas y libro rojo de la flora vascular amenazada. Sociedad Española de Biología de la Conservación de Plantas. Madrid: Gobierno de España.
- Moreno J. C., Domínguez F., Marrero M. & Bañares A. 2015: Application of the Red List index for a conservation assessment of Spanish vascular plants. Conservation Biol. **29:** 910–919. https://doi.org/10.1111/cobi.12437
- Mort M. E., Soltis D. E., Soltis P. S., Francisco-Ortega J. & Santos-Guerra A. 2002: Phylogenetics and evolution of the Macaronesian clade of *Crassulaceae* inferred from nuclear and chloroplast sequence data. Syst. Bot. 27: 271–188.
- Nelson E. C. 2010: Lizards on *Aeonium lancerottense* in Lanzarote, Canary Islands: a new example of pollination by reptiles? Bradleya **28:** 15–18. https://doi.org/10.25223/brad.n28.2010.a3
- Nyffeler R. 2003: *Aeonium (Crassulaceae)*. Pp. 15–23 in: Eggli U. (ed.), Illustrated handbook of succulent plants: *Crassulaceae*. Berlin: Springer.
- Ollerton J., Winfree R. & Tarrant S. 2011: How many flowering plants are pollinated by animals? Oikos **120:** 321–326. https://doi.org/10.1111/j.1600-0706 .2010.18644.x
- Olsen J. M., Skildsen L. I. & Venkatasamy S. 2002: Invasion of pollination networks on oceanic islands: importance of invader complexes and endemic super generalists. Diversity & Distrib. 8: 181–192. https://doi.org/10.1046/j.1472-4642.2002.00148.x
- Ortega-Olivencia A., Rodríguez-Riaño T., Pérez-Bote J. L., López J., Mayo C., Valtueña F. J. & Navarro-

- Pérez M. 2012: Insects, birds and lizards as pollinators of the largest-flowered *Scrophularia* of Europe and Macaronesia. Ann. Bot. **109**: 153–167. https://doi.org/10.1093/aob/mcr255
- Padrón B., Traveset A., Biedenweg T., Díaz D., Nogales M. & Olsen J. N. 2009: Impact of alien plant invaders on pollination networks in two archipelagos. PlosOne 4(e6275). https://doi.org/10.1371/journal.pone.0006275
- Praeger R. L. 1929: Semperviva of the Canary Islands area. Proc. Roy. Irish Acad., B **38**: 454–499.
- Praeger R. L. 1932 [reprint 2012]: An account of the *Sempervivum* group. Stuttgart: Plant monograph reprints J. Cramer & H. K. Swann.
- Rademaker M. C. J. & De Jong T. J. 1998. Effects of flower number on estimated pollen transfer in natural populations of three hermaphroditic species: an experiment with fluorescent dye. J. Evol. Biol. 11: 623–641. https://doi.org/10.1046/j.1420-9101.1998.11050623.x
- Reyes-Betancort A., Santos Guerra A., Guma I. R., Humphries C. J. & Carine M. A. 2008: Diversity, rarity and the evolution and conservation of the Canary Islands endemic flora. Anales Jard. Bot. Madrid 65: 25–45. https://doi.org/10.3989/ajbm.2008.v65.i1.244
- Roubik D. W. 2001: Ups and downs in pollinator populations: when is there a decline? Ecol. & Soc. **5(1)**(2). https://doi.org/10.5751/ES-00255-050102
- Santos J. M., Bentabol A., Hernández Z. & Modino D. 2004: Catálogo de flora de interés apícola de Tenerife. Descripción morfológica de sus pólenes. Santa Cruz de Tenerife: Ediciones Casa de la Miel. Exmo. Cabildo de Tenerife.
- Santos Guerra A. 1983: Flora y vegetación de La Palma.
 Santa Cruz de Tenerife: Ed. Interinsular Canaria S.A.
- Sauerbier H., Cabrera F. & Muer T. 2023: Flora vascular de Canarias. Santa Cruz de Tenerife: Publicaciones Turquesa.
- Schönfelder P. & Schönfelder I. 2018: Flora Canaria. Guía de identificación. Edición en español. – Santa Cruz de Tenerife: Editorial Turquesa.
- Schulz R. 2007: *Aeonium* in habitat and cultivation. San Bruno: Schulz.
- Severio F. & Rodríguez-Rodríguez M. C. 2011: *Gallotia caesaris* (Caesar's lizard). Nectarivory. Herpetol. Rev. **42:** 602–603.
- Thiede J. & Eggli U. 2007: Crassulaceae. Pp. 83–119 in: Kubitzki K., Bayer C. & Stevens P. (ed.), The families and genera of vascular plants. Edited by K. Kubitzki. Volume IX. Flowering plants. Eudicots. Berberidopsidales, Buxales, Crossosomatales, Fabales p.p., Geraniales, Gunnerales, Myrtales p.p., Proteales, Saxifragales, Vitales, Zygophyllales, Clusiaceae alliance, Passifloraceae alliance, Dilleniaceae, Huaceae, Picramniaceae, Sabiaceae. Berlin Heidelberg: Springer.

Traveset A., Nogales M. & Navarro L. 2009: Mutualismos planta-animal en islas: influencia en la evolución y mantenimiento de la biodiversidad. – Pp. 157–180 in: Mendel R., Nogales M. & Navarro L. (ed.), Ecología y evolución de interacciones planta-animal. – Santiago de Chile: Editorial Universitaria.

Turland N. J., Wiersema J. H., Barrie F. R., Greuter W., Hawksworth D. L., Herendeen P. S., Knapp S., Kusber W.-H., Li D.-Z., Marhold K., May T. W., McNeill J., Monro A. M., Prado J., Price M. J. & Smith G. F. (ed.) 2018: International Code of Nomenclature for algae, fungi, and plants (Shenzhen Code) adopted by the Nineteenth International Botanical Congress Shenzhen, China, July 2017. – Regnum Veg. 159. – Glashütten: Koeltz Botanical Books. https://doi.org/10.12705/Code.2018

Valido A., Dupont Y. L. & Olsen J. M. 2004: Bird-flower interactions in the Macaronesian islands. – J. Biogeogr. **31:** 1945–1953. https://doi.org/10.1111/j.1365 -2699.2004.01116.x

Voggenreiter V. 1999: Fitocorología de las 11 especies y sus híbridos de *Aeonium* Webb & Berth. y de *Greenovia diplocycla* Webb *ex* Bolle en La Gomera, Islas Canarias (*Crassulaceae*). – Vieraea **27:** 27–44.

Appendix 1

Additional material studied — Spain, Canary Islands, La Gomera: *Aeonium ×aguajilvense* Bañares (*A. castello-paivae* Bolle × *A. gomerense* (Praeger) Praeger), Aguajilva, 700 m, May 1991, Á. *Bañares* (TFC 37963). *Aeonium ×beltranii* Bañares (*A. canariense* subsp.

latifolium (Burchard) Bañares × A. decorum Webb ex Bolle), El Bailadero, 950 m, Mar 1981, Á. Bañares (TFC 19946). Aeonium ×bravoanum Bramwell & G. D. Rowley ex Heath (A. castello-paivae × A. lindleyi subsp. viscatum (Bolle) Bañares), El Bailadero, 950 m, May 1983, Á. Bañares (TFC 19944); Tamargada, 350 m, Jun 1983, C. Rios Jordan (TFC 9951). Aeonium xcastellodecorum Bañares (A. castello-paivae × A. decorum), El Bailadero, 950 m, May 1983, Á. Bañares (TFC 19944). Aeonium decorum Webb ex Bolle, Barranco del Agua, Valle de San Sebastian, Apr 1845, Bourgeau, (FI 000088). Aeonium gomerense (Praeger) Praeger, without locality, 8 May 1951, E. R. Sventenius (ORT 5621); ibidem, Cumbre Carbonera, 950m, Jun 1996, Á. Bañares & al. (TFC 38718). Aeonium xperezii Bañares (A. appendiculatum Bañares × A. decorum), Barranco de Benchejigua, 600 m, Apr 1985, Á. *Bañares* (TFC 25024). Aeonium sedifolium (Webb ex Bolle) Pit. & Proust, Masca, E. R. Sventenius (ORT 12400). *Greenonium laxiflorum Macarrón & Bañares (A. decorum × Greenovia diplocycla Webb ex Bolle), Barranco de La Laja, 550 m, Mar 1987, Á. Bañares (TFC 25003). — EL HIERRO: Aeonium ×holospathulatum Bañares (A. arboreum var. holochrysum H. Y. Liu × A. spathulatum (Hornem.) Praeger), near San Andrés, Mar 1989, Á. Bañares (TFC 3974); ibidem, San Andrés, 1020 m, 29 May 2012, O. Arango (LPA). — LA PALMA: *Greenonium cabrerae A. Santos (A. spathulatum \times Greenovia diplocycla), Las Manchas (supra Jedey), 1000 m, Jul 1990, Á. Bañares (TFC 37964). — TENERIFE: Aeoniumm ×holospathulatum Bañares nothovar. holospathulatum (A. arboreum var. $holochrysum \times A$. spathulatum), Arafo, 950 m, Mar 1981, Á. Bañares (TFC 19949).

Appendix 2

Experimental crosses between allopatric species of *Aeonium* and *Greenovia*. All crosses resulted in fertile seeds and viable embryos. - (CV) = Cabo Verde; (G) = La Gomera; (GC) = Gran Canaria; (H) = El Hierro; (L) = Lanzarote; (M) = Madeira; (T) = Tenerife.

Parent 1	Parent 2			
Inter archipelagos				
A. arboreum (L.) Webb & Berthel. subsp. arboreum (GC)	A. glutinosum (Aiton) Webb & Berthel. (M)			
A. arboreum subsp. arboreum (GC)	A. gorgoneum J. A. Schmidt (CV)			
A. glandulosum Webb & Berthel. (M)	A. urbicum (C. Sm. ex Hornem.) Webb & Berthel. subsp. urbicum (T)			
A. glutinosum (M)	A. haworthii Salm-Dyck ex Webb & Berthel. (T)			
A. glutinosum (M)	A. sedifolium (Webb ex Bolle) Pit. & Proust (G)			
A. glutinosum (M)	A. tabuliforme (Haw.) Webb & Berthel. (T)			
A. gorgoneum (CV)	A. haworthii (T)			
Inter islands				
A. canariense (L.) Webb & Berthel. subsp. canariense (P)	A. simsii (Sweet) Stearn (GC)			
A. canariense subsp. christii (Burchard) Bañares (P)	A. pseudourbicum Bañares (T)			
A. canariense subsp. christii (P)	G. millennium O. Arango (T)			
A. davidbramwellii H. Y. Liu (P)	A. lancerottense (Praeger) Praeger (L)			
A. davidbramwellii (P)	A. liui O. Arango (T)			
A. decorum Webb ex Bolle var. decorum (G)	A. haworthii (T)			
A. decorum var. decorum (G)	A. simsii (GC)			
A. goochiae Webb & Berthel. (P)	A. volkeri E. Hern. & Bañares (T)			
A. haworthii (T)	A. nobile (Praeger) Praeger (P)			
A. lindleyi subsp. viscatum (Bolle) Bañares (G) (1)	A. nobile (P)			
A. liui (T)	A. percarneum (R. P. Murray) Pit. & Proust (GC)			
A. mascaense Bramwell (T)	A. sedifolium (G)			
A. mascaense (T)	A. simsii (GC)			
A. pseudourbicum (T) (1)	A. saundersii Bolle (G)			
A. pseudourbicum (T)	A. simsii (GC)			
A. saundersii (G)	A. simsii (GC)			
A. sedifolium (G)	A. simsii (GC)			
Intra islands				
A. arboreum var. holochrysum H. Y. Liu (H)	A. valverdense (Praeger) Praeger (H)			
A. canariense subsp. christii (P)	A. spathulatum (Hornem.) Praeger (P)			
A. haworthii (T)	A. smithii (Sims) Webb & Berthel. (T)			
A. holochrysum var. rubrolineatum (Svent.) H.Y.Liu (G) $^{(1)}$	A. lindleyi subsp. viscatum (G)			
A. mascaense (T)	A. pseudourbicum (T)			
A. mascaense (T)	G. aurea (C. Sm. ex Hornem.) Webb & Berthel. (T)			

⁽¹⁾ Hybrid created by Carles Jiménez Box.

Willdenowia

Open-access online edition bioone.org/journals/willdenowia BioOne



Online ISSN 1868-6397 · Print ISSN 0511-9618 · 2022 Journal Impact Factor 1.900 Published by the Botanic Garden and Botanical Museum Berlin, Freie Universität Berlin © 2024 The Author · This open-access article is distributed under the CC BY 4.0 licence