

<sup>1</sup> Drought regimens predict life history strategies in *Heliophila*

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## Abstract

12

## Summary

13        Explaining variation in life history strategies is an enduring goal of evolutionary biology  
14      and ecology. Early theory predicted that for plants, annual and perennial life histories reflect  
15      adaptation to environments that experience alternative drought regimens. Nevertheless,  
16      empirical support for this hypothesis from comparative analyses remains lacking.

17        Here, we test classic life history theory in *Heliophila* (Brassicaceae), a diverse genus of  
18      flowering plants native to Africa, controlling for phylogeny and integrating 34 years of  
19      satellite-based drought detection with 2,192 herbaria occurrence records.

20        We find that the common ancestor of *Heliophila* species was likely an annual, and that  
21      perenniability and annuality have repeatedly evolved, an estimated seven and five times,  
22      respectively. By comparing historical drought regimens, we show that annuals occur in  
23      environments where droughts are significantly more frequent than perennial species. We also  
24      provide evidence that annual plants adapt to predictable drought regimens by escaping  
25      drought prone seasons as seeds.

26        These results yield compelling support for longstanding theoretical predictions by  
27      revealing the importance of drought frequency and predictability to explain plant life history.  
28      More broadly, this work highlights scalable approaches integrating herbaria records and  
29      remote sensing to address outstanding questions in evolutionary ecology.

30        *Keywords:* drought adaptation, herbaria records, *Heliophila*, life history evolution,  
31      phylogeography, remote sensing

32 Drought regimens predict life history strategies in *Heliophila*

33 **Introduction**

34 Understanding the causes and consequences of life history variation is a longstanding  
35 goal of ecology and evolutionary biology (Cole, 1954). In plants, life histories are especially  
36 diverse, with some species completing their life cycle in a number of weeks to others that live  
37 for thousands of years (Brown, 1996). Along this continuum in angiosperms an important  
38 division exists distinguishing annuals which complete their seed to seed life cycle within a  
39 single calendar year from perennials which can persist over multiple years. Annual plants do  
40 not need to survive through the full range of seasonal environmental variation and spend at  
41 least some portion of the year as a seed where they are relatively protected from  
42 environmental stress. In contrast, perennial plants can continue vegetative growth over  
43 multiple years and must survive conditions experienced during all seasons but can also  
44 benefit from competitive advantages and, if iteroparous, multiple bouts of reproduction.  
45 These represent fundamentally different life history strategies and predicting their occurrence  
46 is important for community, ecosystem, and agricultural ecology. However, the  
47 environmental factors that explain their evolution and distributions remain empirically  
48 unresolved (Friedman & Rubin, 2015).

49 Classical theory predicts shorter life spans in environments where adult mortality is  
50 high (Charnov & Schaffer, 1973; Stearns, 1992; Franco & Silvertown, 1996). Because lack of  
51 water is perhaps the greatest threat to survival during vegetative or reproductive growth in  
52 plants, this theory has been extended to the hypothesis that annuality is adaptive when it  
53 allows plants to escape drought (Schaffer & Gadgil, 1975). Indeed, adaptation to drought,  
54 defined as episodes of increased aridity causing plant stress (Passioura, 1996), is often invoked  
55 as an explanation for the success of annual species. And while a few cases are cited where  
56 annuality appears to be more common in environments with greater aridity (Stebbins Jr,

57 1952; Morishima *et al.*, 1984), this hypothesis has yet to be supported while controlling for  
58 the effect of common ancestry (phylogeny) on life habit. In one previous study where this  
59 question was addressed phylogenetically, (Evans *et al.*, 2005) annuals were not found to be  
60 associated with environments that experience more drought. This could be explained by the  
61 relatively small number of species studied and the reliance on a limited number of weather  
62 stations to characterize environments, highlighting the need to develop more scalable  
63 methods to study the geographic distributions of traits such as life history. Thus, in this  
64 study we leverage thousands of herbaria specimens among dozens of species and  
65 high-resolution remote sensing to study the distributions and environmental factors  
66 potentially driving the evolution and distribution of life history.

67 It is also critical to consider another dimension of drought adaptation: the expectation  
68 that annuality is most adaptive when droughts are not only frequent but also predictable.  
69 That is, when the frequency of drought is particularly high during certain seasons. Such  
70 predictability is important for selection to favor and escape strategy during those seasons  
71 which are particularly drought prone. While there has been at least one example of annuality  
72 associated with environments qualitatively classified as “predictable” in a general sense  
73 (Datson *et al.*, 2008), the seasonal predictability of drought experienced by annuals has yet  
74 to be rigorously studied. As such, further empirical work is needed to support the model of  
75 annuality as a mechanism of drought adaptation via escape from drought prone seasons.  
76 Here we study herbarium collection dates to ask whether annuals indeed exhibit evidence of  
77 an escape strategy from seasons with elevated drought frequency.

78 In addition to drought escape in annuals as a mechanism of adaptation to frequent and  
79 predictable droughts, droughts may be necessary for the success of annuals more generally by  
80 acting as episodes of disturbance that provide opportunities for annuals to establish and  
81 compete with sympatric perennial species. Indeed, there is evidence that perennials dominate  
82 in environments where disturbance events are infrequent (Rees & Long, 1992; Corbin &

83 D'Antonio, 2004; Clary, 2012). The resulting prediction from this hypothesis is that in the  
84 absence of frequent drought, perenniability should evolve. However, little is known about this  
85 component of life history evolution because previous work has almost entirely focused on the  
86 origins of annuality rather than perenniability (Friedman & Rubin, 2015). This highlights the  
87 need to study taxa which have seen transitions from annual to perennial life histories as well.

88 Here we combine a long-term global dataset of satellite detected drought events with  
89 metadata from natural history collections to test these classic hypotheses within the African  
90 endemic mustard genus, *Heliophila* L. (Brassicaceae). If annuality is an adaptive strategy  
91 allowing plants to escape drought prone seasons, then drought frequency should predict the  
92 distribution of life history strategies across landscapes, and annual species should be more  
93 commonly associated with drought prone regions than perennial species. Additionally, if  
94 perenniability offers competitive advantage in the absence of drought, associations between life  
95 history and drought frequency should be significant when phylogenies include transitions  
96 from annual to perennial life history strategy. Finally, if annual species have adapted to  
97 escape predictably drought prone seasons, observations of growing annual species  
98 (i.e. occurring in forms other than seed) should be rare during seasons when drought  
99 frequency is highest. Phylogenetic relatedness can influence tests of associations between  
100 species' traits and their environments by confounding common environments caused by  
101 selection from common environments caused by ancestry. (Felsenstein, 1985; Barrett *et al.*,  
102 1996). Therefore, we assessed the relationship between life history distribution and drought  
103 frequency while controlling for phylogeny.

104

## Materials and Methods

105 **Data**

106       **Data availability.** All analyses were performed using R. All data and the source

107 code to produce this manuscript are available at <https://github.com/greymonroe/heliophila>.

108       **Life history data for *Heliophila*.** *Heliophila* is a genus of flowering plants

109 endemic to southern Africa including the Cape Floristic and Succulent Karoo Regions. These

110 are among the most botanically diverse environments on Earth and the *Heliophila* species

111 occurring there are considered to be among the most phenotypically diverse genera of the

112 family Brassicaceae (Mummenhoff *et al.*, 2005; Mandáková *et al.*, 2012). This genus includes

113 both perennial and annual species, and transitions between life history strategy may have

114 occurred multiple independent times (Appel & Al-Shehbaz, 1997; Mummenhoff *et al.*, 2005).

115 Furthermore, the fine scale climatic heterogeneity of southern Africa is ideal for studying the

116 distribution of traits in relation to environmental parameters (Sayre *et al.*, 2013). We used

117 life histories reported by Mummenhoff *et al.* (2005), grouping species into annual or

118 perennial life history categories. Perenniality was defined as any form of perennial life

119 history (e.g., herbs, shrubs, mixed, etc). Because the nature of species reported with mixed

120 traits were unknown (i.e. plasticity vs. genetic variation), we classified these species here as

121 perennial since they can maintain vegetative across multiple years at least to some capacity.

122       ***Heliophila* occurrence records.** To characterize the distributions of annual

123 (studied here, n = 21) and perennial (studied here, n = 21) *Heliophila* species, all (8670)

124 records for the genus *Heliophila* were downloaded from the Global Biodiversity Information

125 Facility (gbif.org) on July 21, 2018 (GBIF, 2018). Herbaria records such as these provide a

126 rich data sources to characterize the geographical distributions of species (Thiers, 2016;

127 Willis *et al.*, 2017; Lang *et al.*, 2018). And as they become digitized (Soltis, 2017), herbaria

128 collections have been used to study relationships between trait distributions, geography, and

129 climate (Davis *et al.*, 2015; Stropp *et al.*, 2016; Wolf *et al.*, 2016; Václavík *et al.*, 2017).

130       **Sequence data for phylogeny.** An alignment of ITS I and II sequences for 21

131 annual and 21 perennial *Heliophila* species was obtained from the authors of Mandáková *et*

<sup>132</sup> *al.* (2012). Individual ITS I and II sequences for *Aethionema grandiflorum*, *Alliaria petiolata*,  
<sup>133</sup> *Cardamine matthioli*, *Chamira circaeoides*, and *Rorippa amphibia* were downloaded from  
<sup>134</sup> Genbank.

<sup>135</sup> **Satellite-detected drought data.** Remotely sensed data is a powerful tool for  
<sup>136</sup> characterizing seasonal patterns in drought because it is less limited in spatial and temporal  
<sup>137</sup> scope and resolution than weather stations or field observations (AghaKouchak *et al.*, 2015).  
<sup>138</sup> From an ecological perspective, droughts are best defined as episodes of plant stress caused  
<sup>139</sup> by elevated aridity (Passioura, 1996). Thus remote sensing offers the additional benefit for  
<sup>140</sup> studying drought as an agent of natural selection because plant stress caused by drought can  
<sup>141</sup> be observed from space (Kogan, 1995a). The remotely sensed Vegetative Health Index (VHI)  
<sup>142</sup> is one such metric, which detects landscape scale reductions in plant cover and temperature  
<sup>143</sup> conditions characteristic of drought (Kogan, 2001). Generated from data collected by NOAA  
<sup>144</sup> AVHRR satellites since 1981, the VHI is a composite index combining Normalized Difference  
<sup>145</sup> Vegetation Index (NDVI) derived quantification of vegetative stress (Vegetative Condition  
<sup>146</sup> Index - VCI) with temperature stress indicated by anomalies in thermal spectra  
<sup>147</sup> (Temperature Condition Index - TCI). These indices were developed to create an unbiased  
<sup>148</sup> quantification of drought across ecosystem types. The VHI of year  $y$  during week  $w$  of [1, 52]  
<sup>149</sup> at pixel  $i$  is derived from the following equations, where  $n$  is the number of years observed.

$$VCI_{y,w,i} = 100 \frac{NDVI_{y,w,i} - NDVI_{min,w,i}}{NDVI_{max,w,i} - NDVI_{min,w,i}}$$

<sup>150</sup> Low values of VCI indicate episodes when plant cover is particularly low for a given  
<sup>151</sup> location during a given time of the year. Thus, it controls for the location and season in  
<sup>152</sup> quantifying plant stress.

$$TCI_{y,w,i} = 100 \frac{T_{max,w,i} - T_{y,w,i}}{T_{max,w,i} - T_{min,w,i}}$$

153 Similarly, low TCI values indicate episdoes of high thermal stress shown to be negatively  
154 correlated with precipitation and soil moisture (AghaKouchak *et al.*, 2015).

$$VHI_{y,w,i} = 0.5(VCI_{y,w,i}) + 0.5(TCI_{y,w,i})$$

155 By combining VCI and TCI, the VHI distinguishes drought from other forms of vegetative  
156 stress (Kogan, 1995b). The use of the VHI to detect drought has been validated globally and  
157 across ecosystem types (AghaKouchak *et al.*, 2015), including in southern Africa, the focal  
158 region of this study (e.g. Figure S1). To date, the VHI has most often been applied for  
159 evaluating drought risk for agricultural research (e.g., Rojas *et al.*, 2011; Kogan *et al.*, 2016).  
160 But it also presents a tool to study seasonal patterns in the frequency of drought across  
161 environments and to test hypotheses about the effect of drought on ecological and  
162 evolutionary processes (Kerr & Ostrovsky, 2003). As such, the VHI has been applied recently  
163 to study drought related ecology of natural species and proven useful for predicting  
164 intraspecific variation in drought tolerance traits and genes (Mojica *et al.*, 2016; Dittberner  
165 *et al.*, 2018; Monroe *et al.*, 2018b). Here, we accessed VHI data at  $16km^2$  resolution from  
166 1981 to 2015 ([https://www.star.nesdis.noaa.gov/smcd/emb/vci/VH/vh\\_ftp.php](https://www.star.nesdis.noaa.gov/smcd/emb/vci/VH/vh_ftp.php)) to  
167 characterize the seasonal drought frequencies experienced by annual and perennial *Heliphila*  
168 species across their native range of southern Africa.

## 169 Analyses

170 **Drought frequency calculations.** To characterize drought regimens across the  
171 distributions of annual and perennial species of *Heliphila*, we calculated drought during  
172 different seasons at the location of observations for *Heliphila* records using the VHI.  
173 Specifically, we created maps of the frequencies of observing drought conditions between  
174 years (VHI<40, NOAA) during the winter (quarter surrounding winter solstice), spring  
175 (quarter surrounding spring equinox), summer (quarter surrounding summer solstice) and

176 fall (quarter surrounding fall equinox) from 1981 to 2015 across the range of *Heliophila*.  
177 From these maps, the drought frequency (the number of times drought is observed divided  
178 by the total number of years, 34) during the winter, spring, summer, and fall were extracted  
179 for the locations of all GBIF records.

180 **Filtering of occurrence records.** To avoid instances with spurious location data,  
181 we filtered raw GBIF records by restricting our analyses to include only:

- 182 • records for species with reported life history
- 183 • records with geospatial data
- 184 • records without known geospatial coordinate issues (i.e., coordinates reported are those  
185 of herbarium)
- 186 • records from collection sites classified as land pixels in the VHI dataset
- 187 • records from Africa (to exclude locations of cultivation)
- 188 • records without duplicates (i.e., identical species, location, collection date)

189 **Phylogeny construction and ancestral state estimation.** Outgroup (

190 *Aethionema grandiflorum*, *Alliaria petiolata*, *Cardamine matthioli*, *Chamira circaeoides*, and  
191 *Rorippa amphibia*) and ingroup *Heliophila* ITS I and II sequences were aligned using  
192 MAFFT (Katoh *et al.*, 2002) with strategy G-INS-I, offset value 0.1, and all other options  
193 set as default. The *GTR + Γ* model of nucleotide substitution was determined to best fit the  
194 data based on AIC using jModelTest2 (Guindon & Gascuel, 2003; Darriba *et al.*, 2012). A  
195 maximum clade credibility tree with branch lengths as relative time was estimated by  
196 summarizing data from six runs of 100,000,000 generations of Bayesian Markov chain Monte  
197 Carlo conducted in BEAST 2 (Bouckaert *et al.*, 2014). Model selection and phylogenetic  
198 analyses were conducted through the CIPRES Science Gateway (Miller *et al.*, 2010).

199 Ancestral state estimation was performed in R using the package phytools (Revell, 2012) to  
200 generate 10,000 stochastic character maps simulated under an equal rates model of character  
201 evolution for the trait life habit (annual or perennial).

**202 Comparison of drought frequency between annual and perennial species.**

203 To evaluate the hypothesis that annual and perennial life history strategies reflect  
204 adaptations to alternative drought regimens, we tested the corresponding prediction that the  
205 observed distributions of annual and perennial *Heliophila* species would be significantly  
206 associated with historic drought frequency. We tested for a relationship between drought  
207 frequency and life history, season, and their interaction by analysis of variance while  
208 including species as a random effect using the lme4 package in R (???) and compared  
209 annuals and perennials using by Tukey adjusted post-hoc contrasts. We next calculated the  
210 mean drought frequency during the winter, spring, summer and fall for each species. Because  
211 shared evolutionary history of closely related species can lead to spurious associations  
212 between traits and environments (Felsenstein, 1985), we tested for a relationship between life  
213 history strategy and drought frequency while controlling for phylogeny using phylogenetic  
214 logistic regression (Ives & Garland, 2010). This statistical approach is designed to control for  
215 the confounding effects of common ancestry's influence on demographic features such as  
216 geospatial relationships when addressing hypotheses about the role of natural selection on  
217 trait distributions.

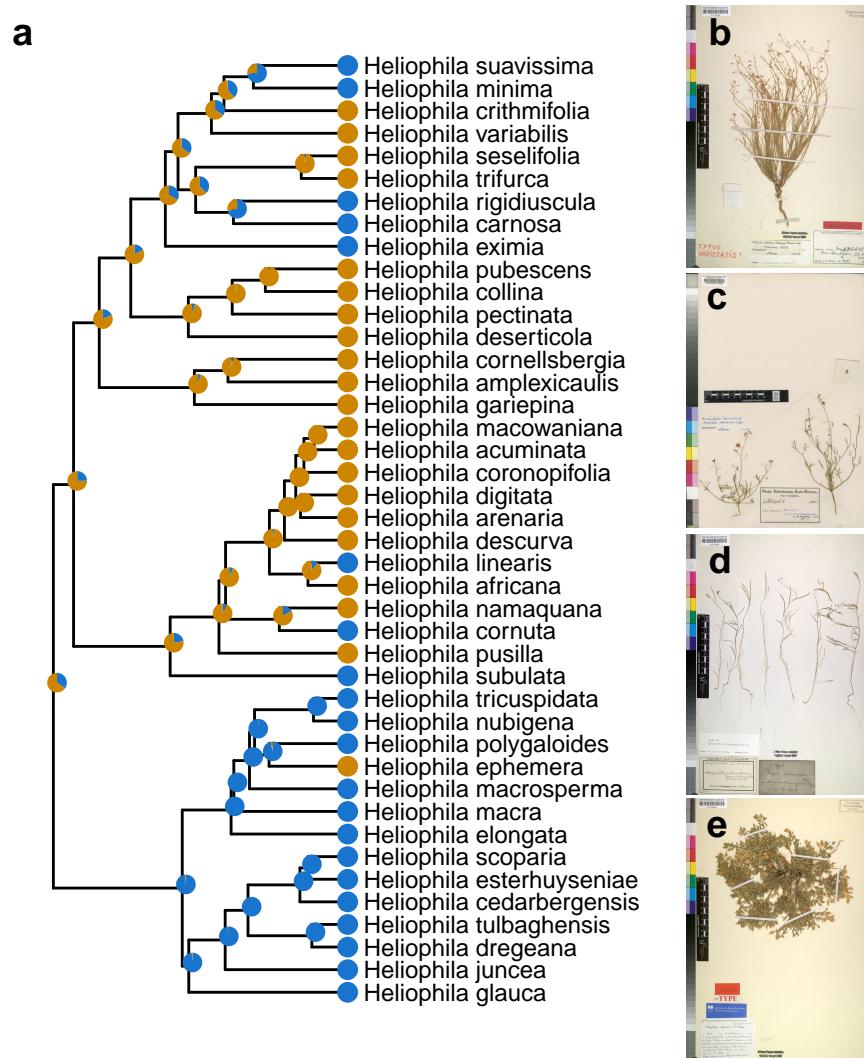
**218 Collection dates.** To test the hypothesis that annual species have adapted to

219 escape drought prone seasons as seeds, collection dates for herbarium specimens were  
220 compared between annual and perennial species. Comparisons of distributions were made by  
221 Two-sample Kolmogorov-Smirnov test and Barlett variance test.

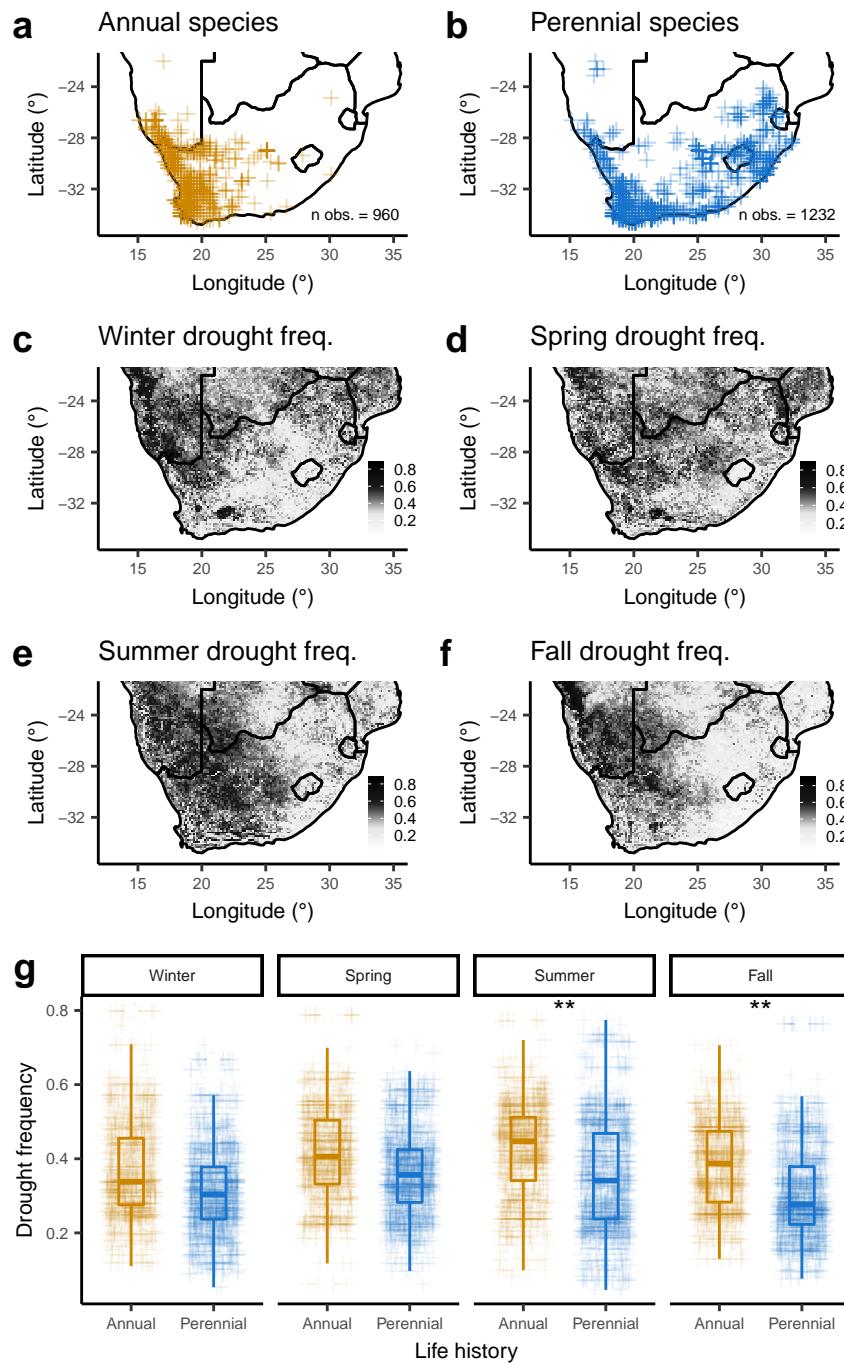
**222 Results**

223 The topology of the estimated *Heliophila* phylogeny was consistent with previous

224 studies (Mummenhoff *et al.*, 2005; Mandáková *et al.*, 2012). Based on 10,000 stochastic  
225 character maps simulated under an equal rates model of character evolution in life history,  
226 an average of approximately seven changes from annual to perennial and five changes from



*Figure 1.* Species and examples of herbaria specimens of *Heliophila* (a) Phylogeny and life history strategies of species studied. Orange circles at branch tips mark annual species and blue circles mark perennial species. At internal nodes, pie charts indicate the estimated posterior probability of being annual versus perennial. Example herbaria specimens accessed via GBIF of (a) *H. minima*, (b) *H. deserticola*, (c) *H. coronopifolia* and (d) *H. ephemera*. Images (b,d,e) courtesy of The Bavarian Natural History Collections (CC BY-SA 4.0) and (c) The London Natural History Museum (CC BY 4.0). Links to images are found in the supplement.



*Figure 2.* Locations of occurrence records of (a) annual and (b) perennial *Heliophila*. Drought frequency during the (c) winter, (d) spring, (e) summer and (f) fall detected using the VHI. (g) Drought frequencies during each season at the observation locations of annual and perennial *Heliophila* (Post-hoc contrasts annuals and perennials from ANOVA, \*\* =  $p < 0.01$ ).

227 perennial to annual are observed per stochastic character map (Figure 1a). These results  
228 suggest that the ancestral state of *Heliophila* was annual and that both character states have  
229 arisen independently multiple times.

230 Out of 8670 *Heliophila* GBIF records, 6634 were for species with reported life history  
231 (Mummenhoff *et al.*, 2005), 2856 had geospatial data, 2833 did not have geospatial issues,  
232 2684 were located on pixels classified as land having drought measurements, 2543 were  
233 located in Africa, 2192 were not duplicated. Thus, after all filtering steps, 2192 records for  
234 42 species (Figure 1, Table S1) passed for further analyses. The number of samples varied  
235 between species, with a mean of 52.19 samples per species. *H. rigidiuscula* had the most  
236 records, 201, and *H. cornellsbergia* the fewest, 2 (Table S1).

237 There were clear visual differences between the distributions of the 960 annual and the  
238 1232 perennial *Heliophila* observation records (see Figure S2 for maps of individual species).  
239 While annual species were generally found in the western regions of South Africa and  
240 Namibia, primarily in the Cape Floristic Region and Succulent Karoo (Figure 2a), the  
241 occurrence of perennials extended to the southern and eastern coast of South Africa (Figure  
242 2b).

243 The frequency of drought varied considerably across the ranges of *Heliophila* species  
244 (Figure 2c-f). This heterogeneity is expected, given that this is one of the most climatically  
245 diverse regions of the Earth (Sayre *et al.*, 2013). It is worth noting the east to west cline in  
246 drought frequency observed during the summer, which distinguishes the high drought  
247 frequency of the Kalahari Sands and Namib Desert phytogeographic regions from the low  
248 drought frequency of the Drakensberg Mountains and Coastal Zambesian phytogeographic  
249 regions. In the Cape phytogeographic region there was finer scale heterogeneity in drought  
250 frequency during the summer.

251 We found that the frequency of drought was significantly higher at the locations of

occurrence records for annual species. When comparing across all occurrence records (all records rather than species means, Figure 2g), a mixed-model analysis of variance which included species as random effect revealed a significant relationship between drought frequency and life history ( $p < 0.01$ ), season ( $p < 0.01$ ) and their interaction ( $p < 0.01$ ) (Table S2). Post-hoc contrasts showed that the frequency of drought was significantly higher at the location of annuals during the summer (z ratio = 3.93,  $p = < 0.01$ ), and fall (z ratio = 4.06,  $p < 0.01$ ). Because a comparison across all occurrence records does not account for variation in the number of records per species (Table S1) or species relatedness (Figure 1a), we also tested whether mean drought frequency values of each species were significantly different between annuals and perennials using phylogenetic logistic regression. We found that the mean drought frequencies were significantly higher ( $\alpha = 0.05$ ) in annual species during the spring, summer, and fall (Table 1, Figure 3a,b). These findings indicate that common ancestry alone does not explain differences in the drought frequencies experienced between the environments of annual and perennial *Heliophila*.

The preceding results indicate that annual species are found in environments where droughts are significantly more frequent, especially in the summer and fall. Classic life history theory hypothesizes that annuality reflects adaptation to such environments because it allows species to escape predictable stressful conditions. If this is the case, we would expect that annuals spend the drought prone seasons of summer and fall as seeds. To test this hypothesis, we compared the dates of occurrence records between annual and perennial *Heliophila* species. The distributions reveal a considerable difference in the timing of observation of these two life histories. In comparison to perennials, which appear to be collected throughout the year, annuals are almost exclusively observed during the winter and spring (Figure 3b). The differences between the distribution of collection dates were significant by all tests ( $ks.test D = 0.25, p < 0.01$ ;  $bartlett.test K2 = 503.18, p < 0.01$ ) This is consistent with a model of life history in which annual species flower in the spring, set seed, senesce, and die before the summer. Thus, these annual species are likely to remain

Table 1

*Phylogenetic logistic regressions between life history, and the mean drought frequency observed at specimen sites of *Heliophila* species the winter, spring, summer, and fall.*

Predictor	Estimate	P
Intercept	0.7231	0.6636
Winter drought freq.	-1.5452	0.7274
Intercept	5.0107	0.0534
Spring drought freq.	-12.9014	0.0464
Intercept	7.7093	0.0054
Summer drought freq.	-19.9056	0.0042
Intercept	7.0162	0.0082
Fall drought freq.	-20.8174	0.0067

*Note.* Annual species were scored as 0 and perennial species as 1.

<sup>279</sup> dormant during the summer and fall, when drought is the strongest predictor of the  
<sup>280</sup> distributions of annual and perennial life histories (Figure 3a).

## Discussion

<sup>282</sup> To test the hypothesis that annual and perennial plants reflect adaptation to  
<sup>283</sup> alternative drought environments we examined the landscape distribution of life history  
<sup>284</sup> strategies in the large and diverse mustard genus, *Heliophila*. Using metadata of 2192  
<sup>285</sup> occurrence records and a 34 year dataset of satellite-detected droughts, we tested the

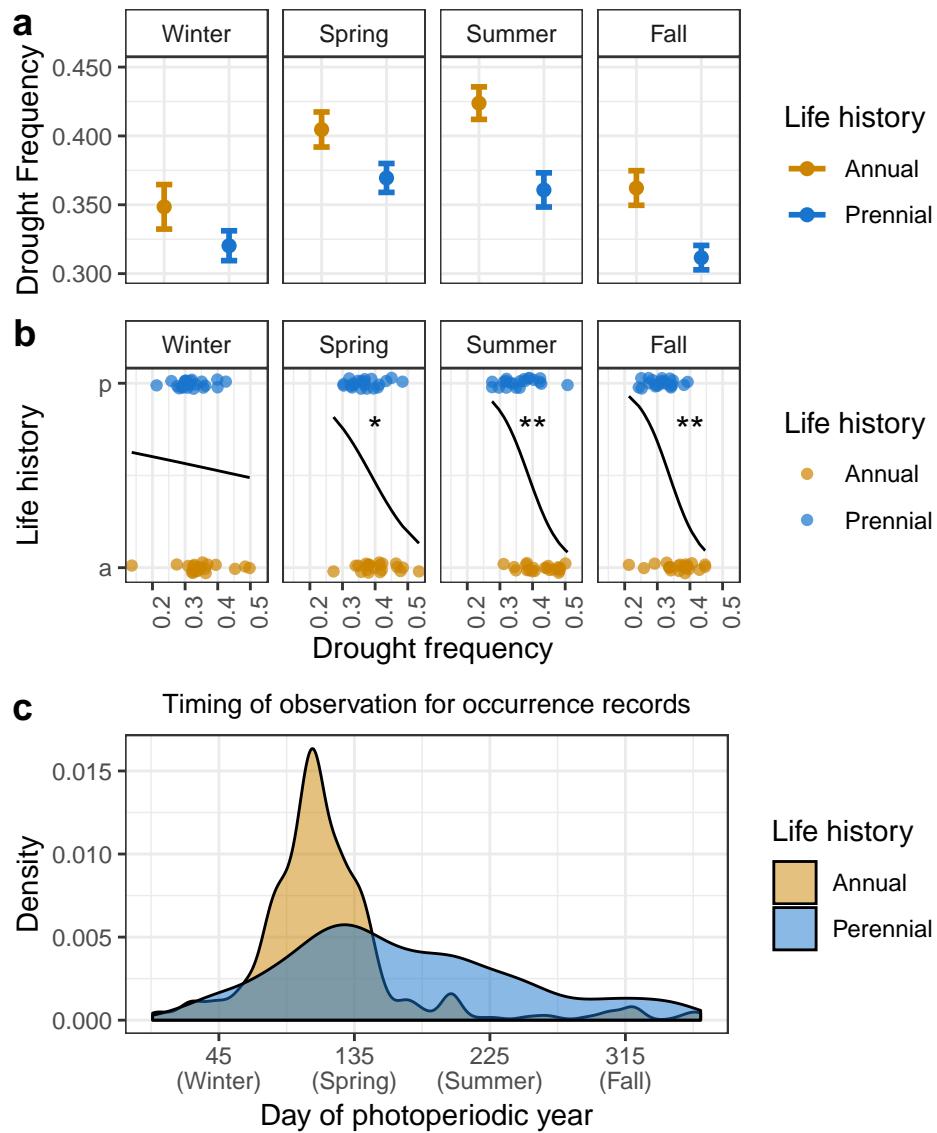


Figure 3. (a) Comparison (mean + SE) of drought frequency across seasons calculated at the occurrence locations of GBIF records of annual and perennial species of *Heliophila*. (b) Results from phylogenetic logistic regression, where the line shows the model fit and \* =  $p < 0.05$ , \*\* =  $p < 0.01$ . Annuals were scored as 0 and perennials as 1. (c) Collection dates of GBIF records of annual and perennial species of *Heliophila* in relation to the photoperiodic calendar where day 1 is intermediate to the fall equinox and winter solstice.

<sup>286</sup> prediction that annual species are more often observed in drought-prone locations than  
<sup>287</sup> perennial species, when controlling for phylogenetic relatedness. We found that drought

frequency is significantly different between the distributions of annual and perennial species, with annuals being found in environments with more frequent drought, and that this signal is strongest during the seasons when annuals are likely escaping via seed dormancy. These results remain significant while controlling for the phylogenetic relationships of *Helophilus* species, yielding support for the role that natural selection has played in driving contemporary distributions of these alternatives strategies in relation to drought regimens.

We cannot eliminate the possibility that confounding traits or environmental variables are the causative factors explaining variation in the distributions of annual and perennial species. Nevertheless, these results provide quantitative support for the classic prediction that annual species are found in environments that experience more frequent drought than perennial species, building on previous investigations of associations between life history and climate (Morishima *et al.*, 1984; Evans *et al.*, 2005; Datson *et al.*, 2008; Cruz-Mazo *et al.*, 2009). To our knowledge this is the first study to demonstrate a significant association between life history and drought in a phylogenetic context informed by large scale species distribution data and long term drought detection.

Unfortunately, herbarium collections and their associated data do not represent systematic or random sampling of a species distribution. Significant biases in collecting exist, which we have not necessarily controlled for here, and may have some effect on our findings, such as a bias toward collecting near roads or near the locations of natural history collections (Daru *et al.*, 2018). Future research will benefit from systematic sampling efforts to avoid these noted biases. However, the ecosystems of southern Africa include several biodiversity hotspots and are among the most botanically well sampled regions on Earth (Daru *et al.*, 2018), suggesting that this may currently be the optimal region for our analyses of life history distribution. Indeed, we were able to use 2192 occurrence records to study 42 species, which represents a significant advance over relying on personal observations to characterize species distributions.

314 These findings empirically support classical theoretical predictions about the adaptive

315 value of annual and perennial life history strategies. Taken together, they suggest that in

316 *Heliophila*, annual species are adapted to environments with predictable droughts by

317 escaping drought prone seasons during the dormant seed phase of their life cycle. They also

318 suggest that perenniability is adaptive in environments where droughts are less frequent.

319 While most previous work has focused on describing the evolutionary origins of annuality

320 (Barrett *et al.*, 1996; Conti *et al.*, 1999; Andreasen & Baldwin, 2001; Verboom *et al.*, 2004;

321 Friedman & Rubin, 2015) there are at least a few other cases where perenniability appears to

322 have arisen from an annual ancestor (Bena *et al.*, 1998; Tank & Olmstead, 2008). And while

323 early theory predicted selection for annuality when adult mortality is high (Stearns, 1992),

324 we also find evidence that perenniability could be explained by reduced frequency of drought.

325 This is supported by the theoretical prediction that perenniability is advantageous in stable

326 habitats. The phylogeny reveals several transitions from annual to perennial life history

327 (Figure 1a) and the distributions of perennial *Heliophila* extend into regions where drought

328 frequency is low (Figure 2b, Figure S2). Perennials may be able to out compete annual

329 relatives in environments where the infrequency of drought favors strategies that allow plants

330 to benefit from growth over many seasons. This also suggests that annuals rely on drought

331 as a source of disturbance for seedling recruitment when competing with perennials (Corbin

332 & D'Antonio, 2004). Indeed, no annual species were observed in the low drought regions of

333 eastern South Africa (Figure 2, Figure S2).

334 These findings suggest that species with locally adaptive life history strategies could be

335 threatened by rapidly changing drought regimens (Dai, 2011). In light of the findings here,

336 forecasted reductions in rainfall across eastern South Africa (Service & Comission, 2017)

337 could be particularly impactful to plant community compositions. Here we found that this

338 region is currently dominated by derived perennial species of *Heliophila*. However, a scenario

339 in which droughts become more frequent in this region may allow for the establishment of

340 annuals. Such changes in selection patterns and shifts in plant functional diversity could

341 have impacts on ecosystem functioning and processes such as carbon cycling (Garnier *et al.*,  
342 1997; Roumet *et al.*, 2006; Monroe *et al.*, 2018a). Furthermore, the changes in frequency of  
343 drought may be an important factor when considering the use of perennial cropping systems  
344 (Parry *et al.*, 2005; Lelièvre & Volaire, 2009).

345 In conclusion, we find strong support for classic life history theory that predicts that  
346 annuality is adaptive in environments with frequent and predictable droughts. We report  
347 evidence consistent with a life history model in annuals in which they escape drought-prone  
348 seasons during the seed phase of their life cycle. Finally, we find evidence that the  
349 distributions of perennial lineages may indicate a competitive advantage in areas where  
350 droughts are infrequent. More broadly, this work highlights the irreplaceable value of natural  
351 history collections and demonstrates the power of combining such information with large  
352 scale remote sensing data to address outstanding classic hypotheses in ecology and evolution.

353

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357

### Author contributions

358 JGM, BG, KGT and JKM contributed to the design of the research, interpretation,  
359 and writing the manuscript. JGM, BG, and KGT contributed to the performance of the  
360 research and data analysis.

361

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515

**Supplement**

516

**Images used.** <https://www.gbif.org/occurrence/1099023487>

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<https://www.gbif.org/occurrence/1057389408>

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<https://www.gbif.org/occurrence/1099023562>

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<https://www.gbif.org/occurrence/1099023490>

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**Supplementary tables and figures.**

Table S1

*Heliophila* species records and the mean drought frequencies during different seasons at the location of records

Species	LH	n	Winter	Spring	Summer	Fall
<i>Heliophila acuminata</i>	a	28	0.32	0.38	0.41	0.36
<i>Heliophila africana</i>	a	91	0.33	0.35	0.34	0.34
<i>Heliophila amplexicaulis</i>	a	60	0.32	0.36	0.39	0.33
<i>Heliophila arenaria</i>	a	65	0.34	0.37	0.38	0.34
<i>Heliophila carnosa</i>	p	129	0.33	0.37	0.39	0.31
<i>Heliophila cedarbergensis</i>	p	3	0.40	0.43	0.32	0.27
<i>Heliophila collina</i>	a	16	0.35	0.47	0.48	0.45
<i>Heliophila cornellsbergia</i>	a	2	0.33	0.42	0.35	0.21
<i>Heliophila cornuta</i>	p	101	0.35	0.40	0.40	0.34
<i>Heliophila coronopifolia</i>	a	40	0.37	0.42	0.40	0.37
<i>Heliophila crithmifolia</i>	a	97	0.35	0.42	0.45	0.38
<i>Heliophila descurva</i>	a	12	0.36	0.38	0.38	0.29
<i>Heliophila deserticola</i>	a	133	0.48	0.48	0.46	0.45
<i>Heliophila digitata</i>	a	30	0.33	0.38	0.44	0.38
<i>Heliophila dregeana</i>	p	17	0.33	0.37	0.33	0.32
<i>Heliophila elongata</i>	p	82	0.26	0.32	0.30	0.25
<i>Heliophila ephemera</i>	a	3	0.14	0.27	0.31	0.26
<i>Heliophila esterhuyseniae</i>	p	3	0.21	0.30	0.37	0.27
<i>Heliophila eximia</i>	p	12	0.42	0.41	0.32	0.34
<i>Heliophila gariepina</i>	a	12	0.50	0.53	0.48	0.41
<i>Heliophila glauca</i>	p	35	0.29	0.35	0.34	0.33
<i>Heliophila juncea</i>	p	150	0.32	0.37	0.39	0.35
<i>Heliophila linearis</i>	p	94	0.32	0.33	0.28	0.30
<i>Heliophila macowaniana</i>	a	31	0.33	0.38	0.44	0.39

<i>Heliophila macra</i>	p	22	0.30	0.30	0.32	0.29
<i>Heliophila macrosperma</i>	p	5	0.28	0.36	0.35	0.25
<i>Heliophila minima</i>	p	35	0.36	0.45	0.51	0.39
<i>Heliophila namaquana</i>	a	16	0.39	0.46	0.48	0.39
<i>Heliophila nubigena</i>	p	19	0.31	0.36	0.43	0.38
<i>Heliophila pectinata</i>	a	16	0.27	0.34	0.50	0.34
<i>Heliophila polygaloides</i>	p	12	0.40	0.48	0.42	0.34
<i>Heliophila pubescens</i>	a	9	0.31	0.40	0.48	0.39
<i>Heliophila pusilla</i>	a	45	0.32	0.38	0.38	0.34
<i>Heliophila rigidiuscula</i>	p	201	0.30	0.33	0.28	0.24
<i>Heliophila scoparia</i>	p	106	0.31	0.37	0.36	0.31
<i>Heliophila seselifolia</i>	a	80	0.36	0.42	0.45	0.40
<i>Heliophila suavissima</i>	p	92	0.30	0.39	0.42	0.31
<i>Heliophila subulata</i>	p	103	0.29	0.33	0.31	0.29
<i>Heliophila tricuspidata</i>	p	8	0.28	0.33	0.38	0.30
<i>Heliophila trifurca</i>	a	77	0.45	0.48	0.48	0.43
<i>Heliophila tulbaghensis</i>	p	3	0.36	0.41	0.36	0.35
<i>Heliophila variabilis</i>	a	97	0.35	0.41	0.40	0.37

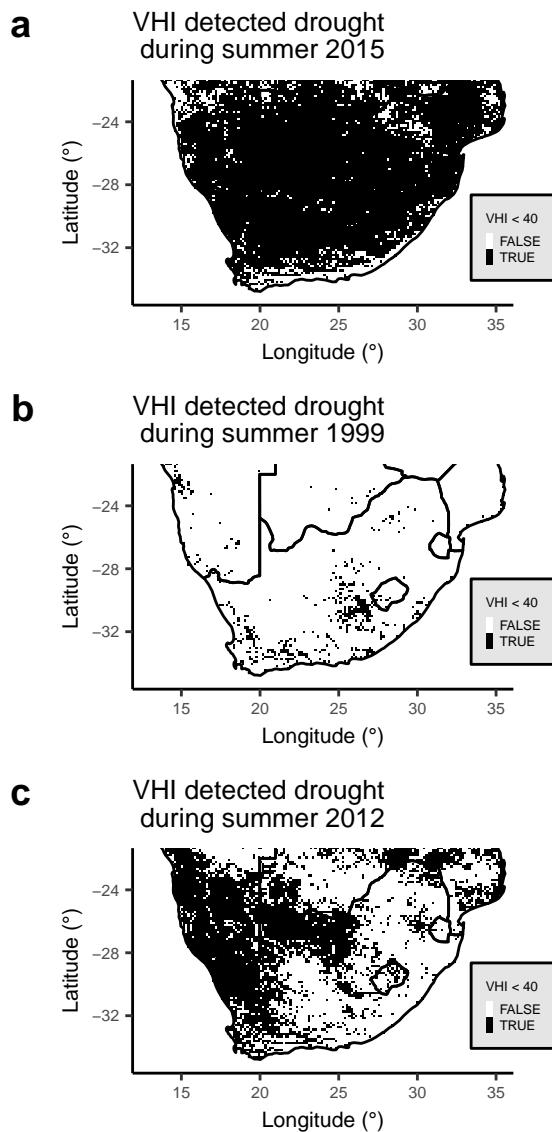
*Note.* LH = Life history (a = annual, p = perennial). n=sample size of GBIF records. Seasons are mean drought frequencies observed at locations of records.

Table S2

*Analysis of variance (ANOVA) to compare drought frequency as a function of life history, season, and their interaction while including species as a random effect.*

predictor	Sum.Sq	Mean.Sq	NumDF	DenDF	F.value	p-value
life history	0.1715	0.1715	1	36.9156	12.2117	0.0013
season	4.3906	1.4635	3	8,718.5058	104.2028	0.0000
life history x season	0.2035	0.0678	3	8,718.5058	4.8301	0.0023

*Note.* Type III Analysis of Variance Table with Satterthwaite's method



*Figure S1.* Example years experiencing contrasting degrees of drought in southern Africa. Vegetative Health Index (VHI) values below 40 indicate remotely sensed drought. Drought detection during these years is validated by previously reported precipitation based estimates of drought occurrence (Monyela, 2017) which confirm that while (a) 2015 was one of the worst drought years on record, (b) 1999 was one of the wettest, and (c) 2012 was typical in terms of precipitation patterns. It is worth noting that drought can be detected using the VHI across ecosystems, including those inhabited by perennial rather than annual *Heliophila* species.



*Figure S2.* Maps of occurrence records for individual species. Orange points indicate annual species. Blue points indicate perennial species.