

Evolution of reproductive isolation in a long-term evolution experiment with *Drosophila melanogaster*: 30 years of divergent life history selection.

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Reproductive isolation, experimental evolution, life history, development time, sexual conflict, *Drosophila*

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Author contributions

CR and HT contributed equally to this work and are joint lead authors. CR, HT, and AKC developed the overall theme of the study; HT and AKC designed the experiments; CR, HT & AKC carried out the experiments; CR and HT analysed the data; CR and HT wrote the first draft of the paper; CR, HT, and AKC contributed to the final draft and revisions.

Conflict of interest statement

The authors declare no competing interests.

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ABSTRACT

We ask if three decades and over 1 500 generations of divergent life history selection on age at reproduction has resulted in the evolution of reproductive isolation (RI) between laboratory populations of *Drosophila melanogaster*. We tested for premating, postmating-prezygotic and postzygotic reproductive isolation between 3 replicate population pairs. Large, evolved differences in body size between selection treatments suggested the potential for prezygotic barriers driven by sexual selection or physical incompatibilities between the sexes. Although a simple prediction would be preference for larger size, creating directional isolation, our results from individual mate choice trials indicate that populations from both selection treatments show a marked bias towards homotypic mate choice; indicative of prezygotic RI driven by sexual selection or sexual conflict. Hybridization between the focal populations resulted in the production of viable adult flies with intermediate size and developmental traits. We observed a suggestive but statistically non-significant trend of fitness decline in the F2 generation of hybrids, but no significant evidence suggesting the evolution of postmating-prezygotic or postzygotic RI. Our findings are in accord with extant literature that posits that premating RI evolves before postmating forms of RI.

19 INTRODUCTION

20
21 Understanding the mechanisms underlying the evolution of reproductive isolation (RI) and
22 the timing of their origination is central to the study of speciation. With the fossil record yielding
23 limited information about this process on microevolutionary timescales, there has been a focus
24 on extant incipient species, either where populations naturally intermingle in contact zones or
25 through experimental crosses. Such populations, part way through the process of RI, can help to
26 illuminate the nature of divergence, as indicated by premating isolation or hybrid inferiority due,
27 for example, to antagonistic epistasis between incompatible alleles (Dobzhansky 1937; Muller
28 1942) and hence offer a snapshot of speciation in action. But what selective agents or historical
29 contingencies have driven the divergence may or may not be apparent, particularly as replication
30 and controls may be absent or limited, and partial isolation and changes in the environment can
31 lead to reversals in the development of RI (Lackey & Boughman 2017), complicating the
32 inference of sequence in the evolution of isolating mechanisms.

33
34 Here we focus on early stages of ecological speciation. Ecological speciation is the process
35 through which reproductive isolation evolves as a consequence of local adaptation with limited
36 gene flow. Divergent selection pressures can contribute to this process by acting directly, or
37 through pleiotropy or tight linkage with genes that are directly responsible for reproductive
38 isolation (Rice and Hostert 1993; Rundle and Nosil 2005). Divergent evolved responses between
39 ecotypes may result in (1) changes in morphology, mating behaviours or physiological characters
40 that hinder mating attempts mechanically or via mate choice (pre mating reproductive isolation);
41 (2) changes in gametes/reproductive machinery that inhibit fertilization (post mating-prezygotic
42 reproductive isolation); and/or (3) genetic incompatibilities that reduce survival or reproductive
43 success of hybrid individuals (postzygotic reproductive isolation).

44 In addition to divergent natural selection in allopatric populations, divergent intersexual
45 coevolution via sexual selection has long been recognized to have a potentially significant role in
46 the evolution of reproductive isolation (Lande 1981; Maan and Seehausen 2011; Panhuis *et al.*
47 2001; Ritchie 2007; Singh and Singh 2014). Likewise, interlocus sexual conflict – where traits
48 that increase the fitness of one sex directly cost the other – can result in rapid coevolution (an
49 “evolutionary chase” or “arms race”) between male and female reproductive traits (Rice &
50 Holland 1997; Parker & Partridge 1998; Rice 1998) and potentially accelerate the process of
51 speciation (Ritchie 2007; Gavrilets 2014; Syed *et al.* 2017); the evolution of female resistance to
52 reduce the direct costs of mating is predicted to contribute to assortative mate choice (Gavrilets
53 *et al.* 2001).

54 While natural systems such as hybrid zones have advanced our understanding of the
55 evolution of RI immensely, researchers are frequently limited to indirect inferences, such as
56 using estimates of gene flow and clines to infer reproductive isolation. Beyond knowledge of the
57 evolutionary history, these systems limit the number RI barriers that can be assessed, and

generally lack replication at the population-pair level. Notable exceptions include stickleback fishes, where repeated invasion of freshwater from marine ancestors, and subsequent differentiation into limnetic-benthic and lake-stream ecotypes provide extensive replication (Lackey & Boughman 2017); and storm-petrels, where repeated separation of populations by breeding time (allochrony) provides replicated evidence of reproductive isolation in sympatry (Wallace *et al.* 2017). Experimental evolution can be a valuable tool for more direct investigation, allowing us to probe potential drivers of speciation between populations with well understood evolutionary histories. Experimental studies in speciation have attempted to test if (1) non-ecological drivers like drift, (2) negative selection against hybridization between ecotypes, (3) divergent selection on populations or (4) sexual conflict produced reproductive isolation (reviewed in Rice and Hostert 1993; Coyne and Orr 2004; Fry 2009; White *et al.* 2020). An obvious limitation to experimental evolutionary studies of speciation is time: research projects rarely exceed granting cycles and few organisms lend themselves to experimental evolution. It is little surprise then that *Drosophila* is far and away the genus most investigated in the laboratory.

Early divergent selection studies produced seemingly equivocal results with respect to creating reproductive isolation. A closer examination suggested that divergent selection was highly effective, given a pre-existent tendency to mate assortatively on the basis of the trait selected upon (Rice and Hostert 1993; Fry 2009). Divergent selection upon behavioural traits resulted in the correlated evolution of reproductive isolation (del Solar 1966; Soans *et al.* 1974; Hurd and Eisenberg 1975; Lofdahl *et al.* 1992) while divergent selection on other traits like bristle number (Barker and Cummins 1969; Santibañez and Waddington 1958) produced negative results. More recently, experimental evolution studies have focused on selection regimes that do not directly involve characters that are known to influence intra-population assortment. For example, a number of studies have investigated the importance of local adaptation to distinct nutrition sources for mating assortment. Divergent selection regimes with different nutritional environments result in premating isolation through (1) the evolution of signalling traits and mating preferences in *Drosophila serrata* (Rundle *et al.* 2005), (2) the evolution of symbiotic microbiota in inbred strains of *D. melanogaster* (Najarro *et al.* 2015; Sharon *et al.* 2010), (3) competitive ability differences between controls and selected individuals in *D. melanogaster* (Belkina *et al.* 2018), (4) positive assortative mating preferences in *D. melanogaster* (albeit unstable) (Nash *et al.* 2019) and in *Saccharomyces cerevisiae* (Mahilkar *et al.* 2023); and (5) postzygotic reproductive isolation through genetic incompatibilities in *Saccharomyces cerevisiae* (Dettman *et al.* 2007) and *Neurospora* (Dettman *et al.* 2008).

Life history traits are less frequently studied in the context of reproductive isolation, but have also received some attention – predominantly in studies of allochronic speciation. Powell *et al.* (2020) note how adaptation to differences in seasonal timing has been shown to contribute to reproductive isolation in a wide variety of organisms including plants (Savolainen *et al.* 2006; Lowry *et al.* 2008), insects (Tauber *et al.* 1977; Wood & Guttman 1982; Horner *et al.* 1999; Eubanks *et al.* 2003; Abbot & Withgott 2004; Forbes *et al.* 2009; Ordning *et al.* 2009; Wadsworth

et al. 2013; Hood et al. 2015) and vertebrates (Friesen et al. 2007). Speciation through other life-history traits and correlated changes in intersexual coevolution has received less attention, due to the myriad challenges associated with studying these systems in nature. Divergent selection on pre-adult development time has been shown to produce pre-mating isolation through (1) disturbances in circadian rhythm resulting in changes in phases of mating in the melon fly, *Bactrocera cucurbitae* (Miyatake and Shimizu 1999), and (2) evolved body size differences, and correlated levels of sexual conflict, in *Drosophila melanogaster* (Ghosh and Joshi 2012). Likewise, divergent selection on body size results in premating isolation, through mechanical incompatibilities during attempted copulations between *Columbicola* feather mites (Villa et al. 2019).

Age-at-reproduction has been extensively manipulated using experimental evolution in *Drosophila* to investigate correlated changes in longevity, development time, body size and a slew of associated traits (Prasad and Joshi 2003). In this study, we use an exceptionally long term evolution experiment (LTEE) with population pairs separated by over 1,500 generations and three decades under divergent selection in allopatry. The two *D. melanogaster* life-history selection regimes used in this study are selected to reproduce late in life (4 weeks from egg; CO) and extremely early in life (9 days from egg; Accelerated CO / ACO). The ACO populations have evolved small body size, rapid development time, and reduced lifespan (Chippindale et al. 1997a; Chippindale et al. 2004a). On the other hand, the CO flies are big, develop slowly and likely have an increased ability to resist copulations as females and to inflict harm as males (Chippindale et al. 2004b; Chippindale et al. 1997b; Mital et al. 2022, Verma et al. 2022).

In the speciation literature, body size is identified as a crucial trait that can drive the evolution of reproductive isolation (Servedio et al. 2011). In *D. melanogaster*, body size is strongly correlated with juvenile development time (Chippindale et al. 1997a; Nunney 1996; Prasad and Joshi 2003; Prasad et al. 2000; Zwaan et al. 1995) and life-history selection on development has been shown to result in the evolution of body size (Chippindale et al. 1997a; Prasad et al. 2000). An important sexually selected trait for both sexes, large females tend to be more fecund (Roff 2002; Stearns 1992) and attract higher courtship effort from males (Byrne and Rice 2006); while large males have increased competitive abilities (Markow and Ricker 1992; Partridge and Farquhar 1983), and secure more frequent copulations as a consequence of female preference (Markow 1986; Partridge et al. 1987, but see Prasad et al. 2007). Given the evolved size differences between the ACOs and COs, a simple prediction is that individuals of both population-types display a mating preference for individuals from the larger CO populations, producing conditions of partial, directional, reproductive isolation driven by sexual selection on the basis of body size, along the lines observed by Ghosh and Joshi (2012).

In *D. melanogaster*, divergent reproductive strategies between the sexes have long been recognized (Bateman 1948) potentially resulting in interlocus sexual conflict. Frequent matings increase male fitness but can decrease female fitness as a result of male harassment and the harmful effects of genital wounding and seminal fluid (Chapman et al. 1995; Bonduriansky et al.

2008; Travers *et al.* 2015). The harm experienced by females as a result of mating, measured by lifespan and egg production rate, is correlated with male body size (Pitnick and García-González 2002; Friberg and Arnqvist 2003). In the context of our two selection treatments, we predict that male harm and female resistance traits will be stronger in the CO populations compared to the ACO populations, due to the difference in body size between the large COs and smaller ACOs. In experimental hybridization between CO and ACO lines, this difference could drive premating isolating barriers to evolve in one direction, with CO females resistant to mating with the smaller ACO males, and ACO females vulnerable to unwanted mating with CO males. Indeed, working with a similar system, Ghosh and Joshi (2012) found such one-directional isolation, driven by large females resisting mating attempts by small males, and a preference for (or lack of resistance to) large males in females of both treatments. On the other hand, especially with the large number of generations in allopatry, mating signals and preferences may have diverged between populations in our selection experiment creating the possibility for positive assortment in mate choice experiments.

We investigated mate choice through individual female, male, and group mating choice assays in the ACO/CO complex of selected lines. We further looked for potential gametic incompatibilities (postcopulatory prezygotic isolation) by comparing egg hatchability in hybrid matings to the parental populations. Finally we looked for postzygotic effects (hybrid breakdown or vigour) in developmental traits, body size, and adult fitness.

METHODS

Experimental populations

Our experiments employed the accelerated development (ACO) (Chippindale *et al.* 1997a) and control-old (CO) (Rose *et al.* 1992) populations. All populations are maintained under standard laboratory conditions: moderate density (larval density of 80 - 120 larvae / vial), 25°C, 12:12 light:dark cycle on banana/agar/killed-yeast medium, and with a census population size of approximately 2,000 individuals per generation.

The five CO populations (CO₁-CO₅) were derived from like-numbered O populations in 1989 (Rose *et al.* 1992). Each CO population is maintained on a 28-day discrete life cycle, with flies placed into cages at a density of ~1000 flies/cage. On day 26, yeast paste is added to the medium to stimulate egg-laying, and on day 28/day 0, eggs are collected at 80-120 eggs per vial. In 1991, one A population (ACO₁-ACO₅) was derived from each of the five CO populations and initially subjected to direct selection for accelerated development. After generation 175, these accelerated populations were maintained on a discrete life cycle of 9 days.

Experimental populations were derived from three of the five pairs of ACO and CO populations (ACO₁, ACO₃, ACO₅ and CO₁, CO₃, CO₅). To remove environmental effects and synchronise experiments, all populations were maintained over a control 14 day life cycle for two generations prior to any assay. Prior to experiment, life cycles were staggered in light of evolved development time differences (ACO << CO) with a 48h difference in egg collection time to synchronise eclosion time of adults. Each assay was repeated for three replicate population pairs: CO₁-ACO₁, CO₃-ACO₃, and CO₅-ACO₅.

Female mate choice assay

To test for the existence of premating reproductive isolation between populations as mediated by female preference, we evaluated the mate choice of ACO and CO females. Flies were sorted by sex and collected as virgins within 6 hours of eclosion under light CO₂ anaesthesia. A food dye technique was used to differentiate between males from each regime (developed from Verspoor *et al.* 2015). Dye techniques were calibrated based on tests of mortality, and the volume of dye used did not result in detectable harmful side-effects, measured through changes in lifespan. Vials were dyed three days prior to the assay using 6 drops of red or blue food dye. One day prior to the experiment, male flies were added to coloured vials at a density of 10 flies/vial and left to ingest the dye for 24 hours. Females were acclimated to individual vials for at least 24 hours prior to the assay. At the time of the assay one male of each population identity and colour was mouth aspirated into an empty vial, then flipped into a vial containing an acclimatised female. Mating latency (time to amplexus formation), colour of the mated male and mating duration were recorded. Under the conditions we used, copulations lasted on average between 15-20 minutes. A viable mating was considered one that lasted for more than

two minutes, and vials in which mating did not occur within 60 minutes were discarded. This represents a commonly employed design to test mate choice (Dukas 2005, Ghosh & Joshi 2012, Arbuthnott *et al.* 2017). Vials were randomised so that observers were unaware of female identity. Trials were colour balanced so that in half of the trials the ACO male was blue and the CO male red, and in the other half the alternate dye pattern was applied. Flies were 2-3 days old at the time of the assay.

This assay was repeated over two days, with female choice tested in both a morning (9am-12pm) and afternoon (1pm-4pm) trial each day. In each of the 4 trials, we tested the choice of 50 females from both selection regimes for each of the 3 replicate populations. The ratio of homotypic to heterotypic matings was calculated across the 50 replicate vials for each combination of replicate population and selection regime.

Male mate choice assay

It is difficult to distinguish the influence of male competition and courtship effort from female preference and resistance. By testing individual male mate choice in addition to female choice and contrasting the results, we hoped to uncover the source of any observed non-random mating pattern. From the time of virgin collection, males were maintained at 10 males/vial. Two females, one from each selection regime, were acclimatised to the same vial 24 hours prior to the assay. Female flies from ACO and CO selection treatments could be identified reliably based upon size alone. On the day of the experiment, one male was “pootered” (mouth aspirated) into a vial containing two females. As with the female choice assay, observers recorded mating latency, the size of the mated female (large or small) and mating duration and all other procedures were the same.

Group mate choice assay

To test for the existence of premating reproductive isolation between ACO and CO populations when mating occurred in a group environment, we evaluated female mate choice in vials containing 10 females and 12 males from each regime. Flies were handled and marked using the procedures outlined above for individual assays. On the day of the assay, male vials of paired ACO and CO populations were combined and then flipped into a female vial. Vials were observed until at least 8 (out of a possible 10) simultaneously copulating pairs were observed, and were then frozen for 60 minutes without disruption to the copulating pairs. The frozen pairs were observed, and the colour of each mated male was recorded. This methodology was repeated over two trials. During each trial, 10 vials were tested for each regime within each replicate population.

Hatchability

233
234 In order to determine the existence of postmating prezygotic isolation we compared the
235 hatch rate of eggs produced from F1 and F2 crosses $CO_i\text{♀} \times ACO_i\text{♂}$ and $ACO_i\text{♀} \times CO_i\text{♂}$ as well
236 as the parental crosses $CO_i\text{♀} \times CO_i\text{♂}$, and $ACO_i\text{♀} \times ACO_i\text{♂}$ ($i = 1,3,5$). Crosses were performed
237 in vials with 12 virgin females and 10 virgin males. A period of at least 4 hours was allowed for
238 mating and egg laying. From each cross, male and female progeny were collected as virgins and
239 maintained in sex specific vials. Hybrid and parental crosses were staggered in an attempt to
240 synchronise time of eclosion.

241
242 For tests of hatchability, 100 flies of each sex were deposited in a collector bottle with a
243 plate of yeasted food. After an egg laying period of 3 hours the plate was removed. Eggs were
244 transferred into vials using fine brushes and egg collection solution (Ashburner 2005). Each vial
245 contained 90 eggs arranged in a grid pattern to facilitate easy counting. 9 such vials were created
246 for each cross identity, for each of the three replicate populations. 24 hours after egg collection,
247 the proportion of hatched eggs was determined. Vials continued to be maintained under standard
248 conditions to be used in tests of viability, development time, and size.

249
250 Larvae to adult viability

251
252 To uncover any survival differences, we compared the larvae to adult viability of hybrids
253 compared to flies from parental crosses. Twelve days after egg collection, the number of eclosed
254 adults in each vial was recorded. The difference between the number of viable eggs and eclosed
255 adults was used as a measure of larvae to adult viability. 9 vials were tested for each cross
256 identity in each replicate population.

257
258 Development time

259
260 Development time of flies from hybrid and parental crosses was also analysed as an
261 indicator of hybrid viability. Each vial was observed for eclosion over a period of 4 days,
262 beginning 24 hours prior to the time of expected peak eclosion for each line. For the first 2 days,
263 vials were observed every 6 hours. For the following 2 days, observations were made every 12
264 hours. During each observation, the number of eclosed flies was recorded. An average eclosion
265 time was then calculated for the vial. 9 vials were tested for each cross identity across each
266 replicate population.

267
268 Body Size

269
270 Body size was measured using dry weight as a proxy. During peak eclosion, 20 female flies
271 were collected from each cross identity. These flies were given 1 hour to physically mature and
272 then were frozen. Flies were divided into 4 groups of 5 females and placed into tinfoil dishes to

be dried in an oven at 70 degrees Celsius for 24 hours. Each dish was measured in a microbalance four times: twice with the flies and twice without. These totals were averaged and divided by 5 to provide an estimate of the dry weight of an individual fly.

Fertility - Competitive reproductive fitness

An adult fitness assay was completed to test for parental line and hybrid fertility differences. In this assay, red-eyed focal flies competed with control flies marked with a recessive brown-eye marker (bw^1) derived from the baseline. The IVb population was selected as these flies are a common ancestor to both the focal populations, also having an intermediate life-history selection protocol. The recessive nature of bw allowed the proportion of red eyed progeny to be used as a metric of paternity share.

Focal F1 and F2 flies of the hybrid crosses $CO_i \text{♀} \times ACO_i \text{♂}$, $ACO_i \text{♀} \times CO_i \text{♂}$ as well as the parental crosses $CO_i \text{♀} \times CO_i \text{♂}$, and $ACO_i \text{♀} \times ACO_i \text{♂}$ ($i = 1, 3, 5$) were tested. Crosses were performed in vials with 12 virgin females and 10 virgin males. A period of at least 4 hours was allowed for mating and egg laying. From each cross, male and female progeny (focal flies) were collected as virgins and maintained in sex specific vials at a density of 10 flies/vial.

Focal flies were anaesthetized under CO_2 and combined with four IVb flies of the same sex (competitors) and four IVb flies of the opposite sex (potential mates). Vials were supplemented with a sparse amount of yeast. Female focal and IVb flies were acclimated to vials 24 hours prior to the assay. After 48 hours, flies were flipped into new vials where oviposition occurred. Flies were 5-7 days old at the time of the assay. After 18 hours flies were discarded. At the time of eclosion, the number of red eyed and brown eyed offspring produced from the competition assay were counted. Thirty vials for each sex and cross identity were tested for each of the 3 replicate populations.

Statistical Analysis

Statistical analysis was conducted in R version 4.2.3 -- "Shortstop Beagle" (R Core Team 2023). Data was visually assessed for residual normality and heterogeneity. Where generalised linear models were used, residual dispersion was tested using the DHARMa package (Hartig 2022).

Female, male and group mate choice data was analysed with a repeated G-test for goodness of fit (Ghosh & Joshi 2012, McDonald 2014). In each case, the ratio of homotypic to heterotypic matings was tested for deviation from a 1:1 null expectation separately for each replicate population and overall for each selection regime. To study the effect of the male and female identity on mating latency and duration, we used linear mixed effects models (LMMs) fit

313 with mated male identity and mated female identity as fixed effects and trial ID and replicate
314 population as crossed random effects. Data from female choice and male choice experiments
315 were analysed independently. In both experiments, mating latency models with raw data
316 produced skewed residual distributions. To correct for this, we used mixed linear models using
317 $\log(\text{mating latency} + 1)$ as the response variable.

318
319 To determine the effect of cross identity on fly development time and body size, LMMs
320 were fit with cross identity as a fixed effect and replicate population as a random effect. To
321 determine the effect of cross identity on hatchability, viability and the fertility of flies, binomial
322 generalised linear mixed effects models (GLMMs) were fit with cross identity included as fixed
323 effect and parental replicate included as a random effect. Fertility data was analysed separately
324 for males and females. Due to over-dispersion issues, we remodelled the viability data and the
325 fertility data for both sexes with beta-binomial generalised linear mixed effects models using the
326 'glmmTMB' package (Brooks *et al.* 2017).

327
328 Models were fit using the 'lme4' package (Bates *et al.* 2015). Fixed effects and random
329 effects were analysed using 'anova()', 'ranova()', 'Anova()' or by backwards model selection
330 (Kuznetsova *et al.* 2017, Fox & Weisberg 2019). Post-hoc comparisons between cross identities
331 were performed using the 'emmeans' package (Lenth 2022).

RESULTS

Female Mate Choice Assay

Overall, ACO females showed a significantly higher proportion of homotypic (0.57 ± 0.04 , mean $\pm 1.96 \times \text{se}$) compared to heterotypic matings (0.42 ± 0.04 , $P = 0.01$, Figure 1a), with no significant heterogeneity among replicate population pairs ($P = 0.66$, Table S1). The ACO₅ population showed a significant deviation from the null hypothesis, with more homotypic matings compared to heterotypic matings ($P = 0.04$; Figure S1a). Females from the ACO₃ population showed a similar trend towards more homotypic matings with borderline significance ($P = 0.05$). ACO₁ females showed no significant difference between mating types ($P = 0.26$). When data was pooled across trials within each replicate population, both ACO₃ and ACO₅ populations, but not the ACO₁ population, showed a significant deviation from a random mating ratio ($P_1 = 0.21$, $P_3 = 0.02$, $P_5 = 0.02$), with no significant heterogeneity between individual trials for any of the 3 replicate populations ($P_1 = 0.30$, $P_3 = 0.28$, $P_5 = 0.21$).

Overall, CO females showed a significantly higher proportion of homotypic (0.60 ± 0.05) compared to heterotypic matings (0.40 ± 0.05 , $P = < 0.0001$, Figure 1a), with no significant heterogeneity among replicates ($P = 0.65$, Table S1). Two CO populations, CO₁ and CO₃, showed an overall significant deviation from the null hypothesis, with more homotypic copulations ($P_1 = < 0.01$, $P_3 = 0.01$, Figure S1a). CO₅ females showed a trend towards more homotypic matings, but this was not significant ($P = 0.08$). When data was pooled across trials, all three CO populations showed a significant deviation from the null hypothesis of random mating ($P_1 = < 0.001$, $P_3 = 0.01$, $P_5 = 0.04$), with no significant heterogeneity among trials for any of the replicate populations ($P_1 = 0.13$, $P_3 = 0.15$, $P_5 = 0.28$).

Only mated male identity had a significant effect on mating latency ($P = < 0.00001$, Table S2). The crossed random effects of the replicate population and trial were significant ($P = < 0.0001$), but the cumulative proportional variance attributed to the random effects of replicate population and trial was minimal (0.05). Both ACO and CO females displayed significantly shorter latencies in mating with CO males (11.8 ± 1.2 min, 11.6 ± 1.0 min respectively), compared to ACO males (15.3 ± 1.3 min, 15.5 ± 1.5 min respectively, Figure 2a). There was no significant effect of female identity or interaction between male and female identity.

Female identity and interaction between male and female identity had significant effects on mating duration ($P = < 0.00001$, Table S2). The crossed random effects of the replicate population and trial were significant ($P = < 0.0001$), but cumulative proportional variance attributed to the random effects of replicate population and trial was minimal (0.04). In general, ACO females (15.0 ± 3.8 min) displayed shorter mating durations than CO females (16.7 ± 4.2 min), and both females mated longer with homotypic partners (ACO♀-ACO♂ 15.4 ± 0.4

min, ACO♀-CO♂ 14.3 +/- 0.5 min, CO♀-ACO♂ 16.1 +/-0.6 min, CO♀-CO♂ 17.1 +/- 0.4 min). All pairwise contrasts were significant (Figure 2c) with the exception of ACO females mated with ACO males vs CO females mated with ACO males. There was no significant effect of mated male identity.

Male Mate Choice Assay

Overall, ACO males showed a significantly higher proportion of matings with ACO females (0.64 +/- 0.05) compared to CO females (0.36 +/- 0.05, $P = < 0.00001$, Figure 1b), with no significant heterogeneity between populations ($P = 0.81$, Table S3). All three A populations showed an overall significant deviation from the null hypothesis, displaying more homotypic compared to heterotypic matings ($P_1 = < 0.0001$, $P_3 = 0.01$, $P_5 = < 0.001$). There was significant heterogeneity observed among trials within the ACO₁ population ($P = 0.01$), but not the ACO₃ ($P = 0.70$) or ACO₅ ($P = 0.38$) populations.

Overall, CO males displayed a nearly equal proportion of matings with ACO (0.51 +/- 0.02) and CO females (0.49 +/- 0.02, $P = 0.95$, Figure 1b), with no heterogeneity among replicate populations ($P = 0.93$, Table S3). All three CO populations showed no significant deviation from the null hypothesis of random mating ($P_1 = 0.99$, $P_3 = 0.65$, $P_5 = 0.96$). There was also no significant heterogeneity among trials within any of the replicate populations ($P_1 = 0.97$, $P_3 = 0.54$, $P_5 = 0.91$).

None of male identity, female identity or interaction between male & female identity influenced mating latency (Table S2). The crossed random effects of the replicate population and trial were significant ($P = 0.01$), but cumulative proportional variance attributed to the random effects of replicate population and trial was minimal (0.02). Females from the CO populations displayed a slightly higher mating latency with both ACO and CO males (18.0+/-1.7 min, 18.1+/-1.6 min respectively), compared to the ACO females (16.6+/-1.1 min, 16.9+/-1.4 min respectively) but these differences were non-significant (Figure 2b).

Only mated female identity had a significant effect on mating duration ($P = < 0.00001$, Table S2). The crossed random effects of the replicate population and trial were significant ($P = < 0.0001$), but cumulative proportional variance attributed to the random effects of replicate population and trial ID was minimal (0.04). Males from the both populations displayed a slightly higher mating latency with CO females (16.3+/-0.7 min, 17.3+/-0.6 min respectively), compared to the ACO females (14.8+/-0.5 min, 15.0+/-0.7 min respectively) (Figure 2d). There was no significant effect of male identity or interaction between male and female identity.

Group Mate Choice Assay

Overall, ACO females showed a significantly higher proportion of matings with CO males (0.62 +/- 0.05) compared to ACO males (0.38 +/- 0.05, $P = <0.00001$, Figure 1c), with significant heterogeneity being observed between replicate populations ($P = 0.01$, Table S4). ACO₃ and ACO₅ females showed an overall significant deviation from the null hypothesis, displaying more heterotypic compared to homotypic matings ($P_3 = 0.02$, $P_5 = <0.0001$). ACO₁ females showed no deviation from the null expectation ($P = 0.78$). There was no significant heterogeneity observed among trials within any of the three replicate ACO populations ($P_1 = 0.49$, $P_3 = 0.45$, $P_5 = 0.50$).

Overall, CO females showed significantly more homotypic (0.61 +/- 0.04) compared to heterotypic (0.39 +/- 0.04) copulations ($P = <0.0001$, Figure 1c), with no heterogeneity between replicate populations ($P = 0.12$, Table S4). CO₅ females showed an overall significant deviation from the null hypothesis, with more homotypic compared to heterotypic copulations ($P = <0.001$). CO₁ and CO₃ populations showed no overall deviation from a random mating ratio of 1:1 ($P_1 = 0.10$, $P_5 = 0.07$). When data was pooled across trials, both CO₃ and CO₅ populations showed significant deviation from a random mating ratio ($P_3 = 0.02$, $P_5 = <0.0001$). There was no significant heterogeneity among trials within any of the populations ($P_1 = 0.10$, $P_3 = 0.58$, $P_5 = 0.96$).

Post-mating prezygotic reproductive isolation

Hatchability

Cross identity had a significant effect on egg hatchability ($P = <0.01$, Table S5). The random effect of replicate population was not significant ($P = 0.89$). The only significant differences in hatch rate was the decline of CO_i♀ × A_i♂ F2 hybrids compared to ACO and CO parental lines (Figure 3a). Overall, hatchability was high with average values between 0.96 +/- 0.01 (CO_i♀ × A_i♂ F2 hybrids) and 0.98 +/- 0.01 (ACO parentals).

Postzygotic reproductive isolation

Larvae to adult viability

There was also a significant effect of cross identity ($P = <0.0001$, Table S5), but no significant random effect of replicate population ($P = 0.31$) on the viability of flies. On average, larvae to adult viability was lowest for ACO parental (0.78 +/- 0.03) and highest for CO_i♀ × ACO_i♂ F2 flies (0.93 +/- 0.02). Both CO_i♀ × ACO_i♂ hybrids and ACO_i♀ × CO_i♂ F2 hybrids demonstrated significantly higher viability than ACO, CO and ACO_i♀ × CO_i♂ F1 flies (Figure 3b).

Development time

There was a significant effect of cross identity ($P = <0.0001$, Table S6), but no significant random effect of replicate population ($P = 0.51$) on the development time of flies. All four hybrid crosses demonstrated development times that were significantly longer than ACO parental flies (175.76 +/- 0.5 hours) and significantly shorter than CO parental flies (218.32 +/- 1.2 hours). There were no significant differences in development times among the four hybrid crosses (Figure 3d).

Body size

Cross identity also had a significant effect on the body size of flies ($P = <0.0001$, Table S6). The random effect of the replicate population was not significant ($P = 0.23$). All four hybrid lines showed average individual weights that were significantly higher than parental ACO lines (0.21 +/- 0.01 g), and significantly lower than parental CO lines (0.31 +/- 0.01 g). Hybrid $ACO_i\text{♀} \times CO_i\text{♂}$ F1 flies were also significantly heavier than $ACO_i\text{♀} \times CO_i\text{♂}$ F2 flies (Figure 3c).

Reproductive fitness

Cross identity had a significant effect on female reproductive fitness ($P = <0.0001$, Table S7). Parental CO lines showed the highest average fitness (0.24 +/- 0.03), parental ACO lines the lowest (0.12 +/- 0.02) and hybrids displayed intermediate fitness between parental lines (Figure 4). All cross identities produced significantly more offspring than parental ACO females and flies from all cross identities, except for $CO_i\text{♀} \times ACO_i\text{♂}$ F1 females, produced significantly fewer offspring than parental CO flies. In both cases, second generation (F2) females displayed lower reproductive fitness than first generation (F1) hybrids, but this effect was only significant for the $CO_i\text{♀} \times ACO_i\text{♂}$ crosses. There was no significant effect of population replicate on female fertility ($P = 1$).

There was a significant fixed effect of cross identity ($P = <0.01$, Table S7) but not replicate population ($P = 1$) on the fertility of male flies. $CO_i\text{♀} \times ACO_i\text{♂}$ F1 male hybrids showed the highest average fitness (0.13 +/- 0.04) and ACO males the lowest (0.05 +/- 0.02). CO and $CO_i\text{♀} \times ACO_i\text{♂}$ F1 males demonstrated a significantly higher fitness than ACO males (Figure 4). Similar to that observed in the female assay, male F2 hybrids displayed lower fitness than F1 hybrids, but these differences were not significant. While female flies displayed fitness values around the expected value, males of both parental and hybrid cross identities produced a lower proportion of offspring than expected when in competition with four competitors.

DISCUSSION

Our experimental assays show the striking differentiation between the ACO and CO populations after over three decades and 1500 generations of divergent selection on demography. The ACO population adults eclose nearly two days earlier than the CO population adults do. The clear tradeoff between speed and size is evident, as A adults are about 33% lighter than CO adults, and this is likely related to the strong differences in adult fitness we observed between the two selection treatments. Interestingly, the tradeoff between developmental speed and juvenile viability (Chippindale et al. 1997) is not apparent here or in other recent unpublished data on these populations, suggesting compensatory evolution over time. Given these differences in morphology and life history, we asked if we could detect the early stages of the speciation process for these populations in allopatry under different ecological circumstances.

To summarise our results, we found evidence for the evolution of premating barriers to gene flow between these diverged populations. Mate-choice assays generally reflected positive assortative mating patterns driven by female mate-choice. Females from both focal populations showed strong homotypic mate choice in individual tests, but this pattern was less clear in group tests, particularly as the smaller ACO females tended towards heterotypic mates. Under conditions of female choice, mating durations of homotypic pairs were consistently longer than heterotypic pairs. Despite this (and homotypic mate choice), the larger CO males showed lower latencies to mating with both females. In contrast, under conditions of male choice (no male-male competition), ACO and CO males showed no differences in mating latency. Further, under male choice conditions, both males mated for longer durations with the larger CO females, suggesting sperm catering based on body size rather than homotypic preference. We show that post-mating prezygotic barriers have not evolved to the point of detectability between these populations in the form of gamete incompatibility, measured through egg hatchability. We also find no evidence of postzygotic reproductive isolation in the form of survival to reproductive maturity, and measurements of development time and body size show that hybrids display intermediate trait values relative to the parental treatments. We observed only suggestive trends for postzygotic reproductive isolation in the form of hybrid breakdown in competitive reproductive fitness assays. F_1 hybrids display intermediate fitness to that of the two parentals, however, there is a tendency for F_2 hybrid offspring to display lower fitness than the F_1 hybrids; while potentially an indicator of evolving Dobzhansky-Muller incompatibilities, our data are merely suggestive at this juncture, meriting further investigation.

It has been suggested that prezygotic isolating barriers will typically evolve before postzygotic barriers, and *Drosophila* seems to adhere to this rule (Coyne & Orr 2004, Arthur & Dyer 2015). Within the category of prezygotic barriers, premating RI is expected to evolve before postmating-prezygotic RI (Turissini et al. 2018). Our data are generally in accord with

these predictions, as our diverged population pairs show marked premating reproductive isolation, at least under individual choice tests, but clear evidence for later stage RI is lacking.

Strong divergence in body size exists between ACO and CO animals (Xiao et al. 2019; data herein) and are presumed to correlate with differences in persistence/resistance capabilities between the selection treatments. This size difference led us to expect the presence of some premating reproductive isolation. However, we predicted premating isolation in one direction, on the basis of greater resistance from larger (CO) females, greater persistence of CO males, and a possible preference for large mates. That directional RI was noted by Ghosh & Joshi (2012) in a study similar to ours: bigger mates were preferred by both large/slow and small/fast flies. Contrary to that precedent, females of both regimes displayed strong homotypic mate choice and longer mating durations with their homotypic partners in individual tests. This result is especially intriguing for the ACO females, as they displayed a preference for smaller ACO males even though *D. melanogaster* females are expected to prefer larger males (Pitnick 1991, Jagadeeshan et al. 2015, but see Prasad et al. 2007).

Male mate choice tests may present trickier interpretations. ACO selected males largely mated with homotypic females, but male Cos mated nearly randomly. While ACO males appear to be showing positive assortative mating preferences, both types of males were observed making repeated courtship attempts with both homotypic and heterotypic females (unquantified observations) suggesting that the skewed mating pattern could be the result of the increased capability of CO females to resist unwanted mating attempts, particularly from the smaller A males. In alignment with this hypothesis, we observed a non-significant trend of longer mating latencies with CO females. In populations of flies evolved under a similar pair of life history selection regimes, Ghosh and Joshi (2012) found that small males attempted to court large females, but were unsuccessful, and, as a result, mated more frequently with small females. We therefore cannot easily ascribe mating assortment to the choice of the male, who may be indiscriminating under the test conditions, particularly with no competition for mates. Concurrent with this, we note no differences in the mating latencies of the two male types in the absence of male-male competition. ACO-type females are also under tremendous pressure to mate, as they are selected to lay eggs about 24h after becoming sexually mature. In the absence of competition, many males may have simply taken the most receptive mate first; for the CO males this may have counterbalanced a homotypic preference, creating the ambivalent outcome observed. The results of the mate choice assays taken together suggest that premating isolation in this system is most strongly driven by female choice.

Body size is a likely trait involved in the evolution of reproductive isolation between the population pairs tested. However, it remains possible that the patterns of assortative mating stem from traits that are yet to be characterised. Some possibilities include courtship behaviours (mating call, wing movements) and cuticular hydrocarbon profiles that may have diverged under

the divergent selection regimes. Besides female preference, body size differences also play a clear role in the differential capacities displayed by females to resist mating attempts by males, which creates a further reproductive barrier between large females and small males. Despite large body size differences between the selection treatments, it seems unlikely that the patterns of positive assortative mating can be explained by mechanical incompatibilities in copulation alone; some earlier anecdotal observations suggested they may exist, but we saw no clear evidence in these trials.

In the group mate choice assays, female CO flies continued to show homotypic mating preference, but A females displayed reversed preferences, mating more frequently with heterotypic males than homotypic males. The existence of heterotypic mate choice in groups, in spite of homotypic mate choice in individual assays suggests that the nature of the assay plays a role in the mating patterns observed. In the group mating assays, vials were more crowded (34 individuals vs. 3) - potentially creating circumstances where resisting forcible mating-attempts is more difficult for females. This is a likely explanation given that only the small A females displayed reversed mate choice preferences, while the larger CO females largely reproduced the same mate choice patterns as displayed in individual assays. It is worth testing if similar mate choice assays conducted in larger arenas (such as 90mm petri dishes) would reverse this preference to homotypic choice. At the time these experiments were conducted (since changed) there was an important difference in the mating environment of the ACO and CO selection treatments: whilst the early-reproducing ACO's primarily mated in vials, with only a brief phase in cages for egg-laying, the CO's, with 14 days in cages probably do all mating relevant to fitness in the cage environment. With these differences in the "natural" mating environment of our LTEE, the mate choice patterns found in individual assays seem to show us that A-selected females have a preference for homotypic mating but are unable to exercise it in a group mating setting with larger males.

Asynchronous development time alone can contribute to reproductive isolation, as seen in the experimental work of Miyatake and Shimizu (1999) on the melon fly. Because the mating times diverged in lines selected for development time, significant RI arose that was evidenced in experimental mate choice trials. Similarly, the extreme early reproduction regime of the A populations means that neither CO individuals nor A/CO hybrids could hybridize, as the latter groups would not even be sexually mature before a generation is completed. This scenario is, of course, artificial because natural populations generally do not grow on synchronised, discrete generations. they are not directly responsible for the assortative mating patterns observed in the controlled mate-choice assays.

Post-mating, we report no evidence of reproductive isolation deriving from gamete incompatibility, as would be evident in reduced egg hatch rates. Similarly, our measurements of larva to adult viability data show that there are no hybrid treatments with lower levels of viability

than the purebred parental populations. In fact, we observed trends of hybrid vigour, with some of the hybrid treatments displaying higher larva to adult viability than the parental treatments. Interestingly, the larva to adult viabilities in F2 treatments were significantly higher than that of the parental selection treatments and one F1 treatment. While the increased viability in hybrids could suggest the existence of homozygous deleterious variation in developmental loci in our selection regimes, this should be consistently seen in both F1 and F2 generations, instead of being more common in the F2 generation. Development time and body size displayed no suggestion of abnormalities indicative of developmental instability, with intermediate hybrid trait values relative to the parental treatments in both sexes.

The competitive reproductive fitness of hybrids was generally intermediate to the parental treatments, and we found no evidence of hybrid sterility. Given that both assays subjected a focal fly to 4 competitors of the same sex, we expect focal flies to sire/dam $\frac{1}{5}$ th of the offspring produced, if all else was equal. In female reproductive fitness assays, our populations did cluster around the predicted threshold, with hybrids scoring intermediate fitness levels relative to the low fitness ACOs and high fitness COs. In male reproductive fitness assays though, fitness fell well below the predicted threshold in all treatments. We rule out the possibility of this being a case of Haldane's rule, because male fitness falls well below predicted levels in parental males as well as hybrids. We believe that this is a consequence of the usage of *IVb* flies as competitors/mates in these assays. Selected as common ancestors to both selection regimes, the *IVb*s have evolved in allopatry with both the populations, with a slightly different maintenance regime. We believe that the reduced male fitness estimates may be a consequence of homotypic mate choice by *IVb* females in male fitness assays. While we do see different levels of competitive reproductive output for the different treatments in the female assay, it is not possible to disentangle male mate choice from innate fecundity and harm resistance differences in the females of different selection regimes. Overall, in spite of the depressed male fitness output relative to predictions, hybrids in both sexes still consistently showed intermediacy in fitness, suggesting no reproductive sterility and a link to an additive trait such as body size.

The comparison between F1 and F2 reproductive fitness revealed an interesting trend: F1 fitness was consistently higher than that of F2 offspring, however, this difference was statistically significant in only 1 out of 4 comparisons (2 reciprocal crosses * 2 sexes). We interpret this with caution as a possible trend of hybrid breakdown as a consequence of incompatible gene complexes evolving in the two selection regimes.

We generally found high repeatability across our three independent replicates, which suggests that the characters measured and pre-mating RI observed stem from consistent treatment differences between the CO and ACO populations. Internal random processes of divergence in allopatry, such as the coevolution of arbitrary signaller-receiver relationships in allopatry may also drive the evolution of RI. A system like ours opens the door to a potential test

647 for such divergence if RI were investigated between replicate populations *within* selection
648 treatment, rather than *between* treatments. Although our three-decade experiment seems too
649 young to see more than hints of post-mating RI, more sensitive queries of characters like
650 developmental stability (e.g., as measured by fluctuating asymmetry) may be of interest. Further
651 investigation of the sources of premating RI and its environmental sensitivity are warranted.

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FIGURE CAPTIONS

Figure 1. The proportion of homotypic compared to heterotypic matings recorded during the female (a), male (b) and group (c) mate choice assay. Female (a,c) and male population identity is shown on the x-axis. An asterisk represents a significant deviation from a random mating ratio as indicated by a repeated G-test for goodness of fit ($p < 0.05$).

Figure 2. The latency (a,b) and duration (c,d) of matings in the choice assays. Female (a,c) and male (b,d) population identity is shown on the x-axis. Letters indicate significant differences between mating types.

Figure 3. The hatchability (a), egg to adult viability (b), body weights (c) and average development time (d) of flies from parental and hybrid crosses. Letters indicate significant differences between cross identities.

880 **Figure 4.** The average proportion of red eyed offspring produced by flies from parental and
881 hybrid crosses, in competition with brown eyed competitors. Average proportions were
882 determined for each replicate population and plotted. The dashed line indicates the expected
883 proportion of $\frac{1}{5}$ of the offspring. Letters indicate significant differences between cross identities.
884

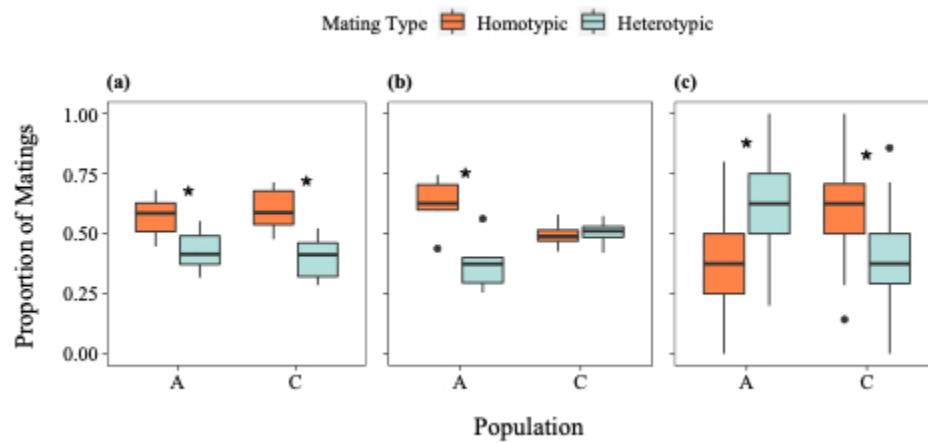


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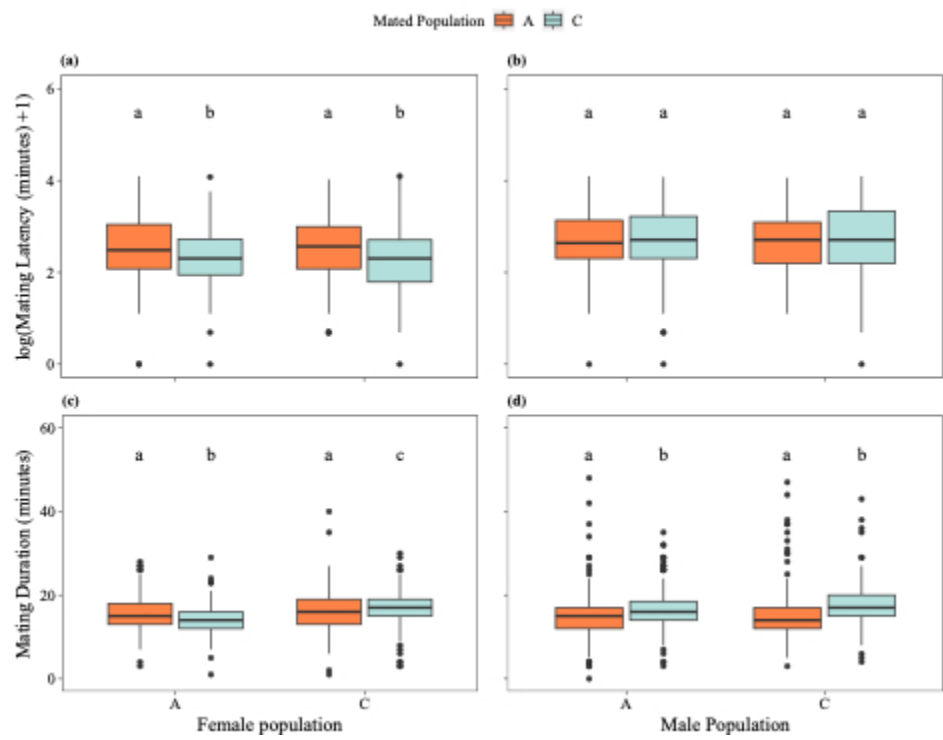


Figure 2. The latency (a,b) and duration (c,d) of matings in the choice assays. Female (a,c) and male (b,d) population identity is shown on the x-axis. Letters indicate significant differences between mating types.

165x131mm (72 x 72 DPI)

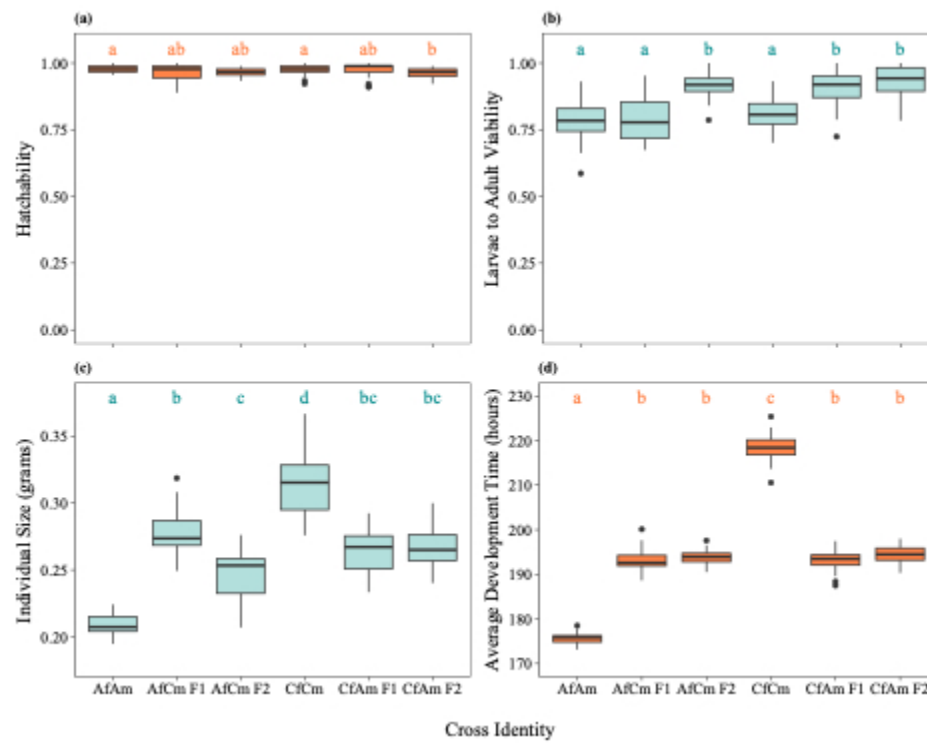


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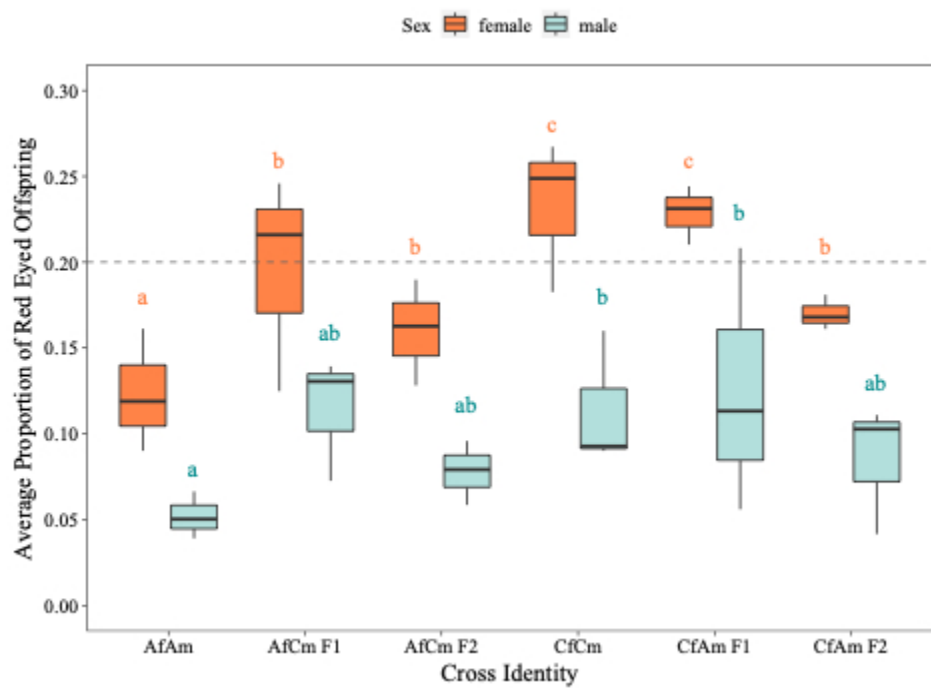


Figure 4. The average proportion of red eyed offspring produced by flies from parental and hybrid crosses, in competition with brown eyed competitors. Average proportions were determined for each replicate population and plotted. The dashed line indicates the expected proportion of $\frac{1}{2}$ of the offspring. Letters indicate significant differences between cross identities.

165x123mm (72 x 72 DPI)