Mating system affects population performance and extinction risk under environmental challenge

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Failure of organisms to adapt to sudden environmental changes may lead to extinction. The type of mating system, by affecting fertility and the strength of sexual selection, may have a major impact on a population's chances to adapt and survive. Here, we use experimental evolution in bulb mites (Rhizoglyphus robini) to examine the effects of the mating system on population performance under environmental change. We demonstrate that populations in which monogamy was enforced suffered a dramatic fitness decline when evolving at an increased temperature, whereas the negative effects of change in a thermal environment were alleviated in polygamous populations. Strikingly, within 17 generations, all monogamous populations experiencing higher temperature went extinct, whereas all polygamous populations survived. Our results show that the mating system may have dramatic effects on the risk of extinction under environmental change.

Keywords: extinction vortex; environmental stress; sexual selection; fertility; adaptation

1. INTRODUCTION

Rapid environmental changes, including those caused by human activity, can result in a mismatch between an organism's physiology and its new environment, potentially leading to population decline and extinction [1,2]. The chances of avoiding extinction will depend on the ability of a population to adapt. However, adaptation may be difficult if environmental stress has a severe effect on demography. Population bottlenecks, caused by unfavourable environmental conditions, may deplete genetic variation necessary for the adaptation process. Moreover, such bottlenecks increase inbreeding, leading to reduced survival and reproductive capacity of individuals associated with inbreeding depression (reviewed by Keller & Waller [3]), hence amplifying negative effects of environmental stress on demography. These synergistic interactions between environmental and genetic factors of extinction, termed ‘extinction vortex’ [4], are further exacerbated by the tendency for inbreeding depression to increase with environmental stress [5].

The type of mating system may affect a population's chances to survive environmental change in several ways. On the one hand, it will affect the opportunity for sexual selection [6]—a process that may enhance the rate of adaptation, provided that better-adapted males win reproductive competitions over maladapted rivals (reviewed by Candolin & Heuschele [7]). Thus, sexual selection should increase the chances of fixation of favourable alleles [8]. Furthermore, sexual selection may increase the effectiveness of selection against deleterious mutations [9–12], which are the main cause of inbreeding depression [13]. Finally, if environmental change adversely affects male fertility, polyandrous matings may benefit females by assuring egg fertilization [14–16]. In a population facing an environmental challenge, this should increase demographic stability, as most or all females capable of producing progeny will be fertilized. Thus, polyandry might prevent populations from entering an extinction vortex.

On the other hand, polygamy decreases effective population size by increasing variance in male reproductive success [17,18]. In consequence, it may increase inbreeding and the risk of beneficial alleles being lost as a result of genetic drift [19,20]. Furthermore, adaptations to sexual competition may trade off with other fitness components [21–23], possibly hindering ecological adaptation and increasing extinction risk [24] (but see [25,26]). Finally, sexual selection may lead to sexual conflict, which in turn may be detrimental to female fitness [27].

Thus, the net effect of mating system on survival prospects of a population subject to environmental stress is not easy to predict and should be addressed with an experimental approach. Experiments investigating the role of mating systems during adaptation to novel environments have given inconsistent results [28–30]. However, as discussed above, influencing the rate of adaptation is one of a number of ways by which mating systems may affect the risk of population extinction under environmental challenge. Crucially, the effects of mating system on population performance under environmental stress severe enough to potentially cause extinction have not, to our knowledge, been studied.

Here, we investigate the impact of mating system on long-term response to increased temperature in replicate lines of the bulb mite Rhizoglyphus robini (Acari: Acaridae), drawn from a laboratory-adapted base population. The study species is characterized by high promiscuity: both sexes mate many times a day, with average time to remating estimated at 80 min. [31]. Females mated to multiple males produce more fecund daughters than
those mated to a single male, as a result of genetic corre-
lution between female fecundity and male sperm
petitiveness [32]. Apart from sperm competition, sexual selection in this species involves fierce male fights
over access to females [33]. Thus, sexual selection acting
both pre- and post-copulation may improve fitness of
bulb mite populations. Indeed, a polygamous mating
system has been shown to help purge inbreeding depression and
decrease extinction rate, compared with a monog-
amous system, in bottlenecked bulb mite populations
[34]. On the other hand, sexual conflict has been shown
to negatively affect reproductive success of bulb mite
females, and the harmful effects of males on females were
selected against under enforced monogamy [35]. These
harmful effects of polygamy may actually increase the risk
of the population being caught into an extinction vortex.

We established selection lines in which we either
enforced monogamy or allowed promiscuity. We applied
two temperature regimes—lines were maintained either at
the increased temperature or at the temperature to which
our base colony was adapted. This way, we manipulated
the opportunity for sexual selection and conflict (mating
system treatment) both among environmentally challenged
and control lines (temperature treatment). Moreover, owing
to the increased variance in male reproductive success under
polygamy on the one hand, and negative effects of
temperature on mite fertility on the other (see §3b), the
effective population sizes \( N_e \) were likely to differ between
treatments. Specifically, we expect that among the lines
evolving at control temperature, polygamous ones had
lower \( N_e \) owing to increased variance in male reproductive
success [18]. Conversely, temperature-induced infertility
should have a stronger effect on \( N_e \) of monogamous lines,
because infertility of one individual in a pair would automa-
tically prevent reproduction of the other. Such
differences in \( N_e \) will be inherently linked to differences
between mating systems in nature and are likely to be an
important part of the net effect of mating system on the
probability of population extinction. We show that, overall,
a polygamous mating system helps populations to adapt to
increased temperature and prevents their extinction, which
seriously affect monogamous populations.

2. MATERIAL AND METHODS
(a) General procedures
Base populations and larger groups of mites were kept in
plastic containers (2.5 cm diameter and 2 cm high). Individu-
ally isolated mites and pairs were kept in 0.8 cm
diameter glass tubes (2 cm high) with plaster of Paris bases
soaked with water. Humidity of greater than 90 per cent
was maintained and powdered yeast was supplied ad libitum
as food.

(b) Base population
The mites used in the experiment originated from a stock
culture combined of two populations adapted to 24°C, one
for 12 years (approx. 280 generations) and the other for 3
years (approx. 60 generations). Each of these two populations
had been derived from a colony of about 200 individuals
found on onions in a garden near Kraków, Poland, in 1998
and 2008, respectively, and had been maintained in the
laboratory in large numbers (more than 1000 individuals, sub-
divided into periodically mixed sub-populations). The two
populations had been mixed approximately 10 generations
prior to the commencement of our experiment. Such a pro-
cedure is expected to increase additive standing genetic
variation, which is crucial in experimental evolution.

(c) Selection lines
To test whether mating system affects evolution at increased
temperature, we established selection lines in which we either
enforced monogamy (M lines, 20 pairs per line) or allowed
sexual selection (polygamy, P lines, 20 mites of each sex inter-
acting freely). We applied two temperature regimes (‘selection
temperatures’)—lines were maintained either at standard 24°C
(control, C lines) or at 28°C (high temperature, HT lines).
Higher temperatures are rarely encountered by bulb mites in
nature—soil temperature at the depths where bulb mites are
found may occasionally reach 28°C only around midday in
summer (M. Klimek 2012, Field Research Station in Łazy,
personal communication)—and had not been encountered
by our laboratory populations throughout their history of
adaptation to 24°C (see §2b). Our pilot study showed that
stock population females developing at 28°C had drastically
decreased fecundity (mean number of eggs per female ± s.e.:
47.53 ± 5.89) compared with those developing at 24°C
(163.87 ± 15.85; t16 = 8.55, p < 0.001), and their progeny
(also developing at 28°C) had lower embryonic viability
(78 ± 5.8%) than the 24°C controls (98 ± 1.3%; t8 = 3.29,
p = 0.002). Thus, increased temperature has considerable
negative effects on reproductive fitness in the bulb mite.

In the M-HT treatment, we established six replicate lines,
and the remaining three treatments were represented by five
replicates each. In P lines, 20 males and 20 females were
placed into one container for 5 days, so that intra- as well as
inter-sexual selection could operate, whereas in M lines individ-
uals were kept in randomly assigned pairs for 5 days, so that
sexual selection was prevented. After this time, all females
from each line were moved to a common container to lay eggs.
The size of the containers ensured low density of ovipositing
females and developing larvae in all the lines, and food was pro-
vided ad libitum. When tritonymphs (last larval stage) emerged,
about 90 of them were isolated to individual vials. Emerging
adults were then sexed, and 20 virgin males and 20 virgin
females were put together either in pairs (M lines) or in groups
(P lines) in order to obtain the next generation. Such a pro-
cedure enables natural selection on survival and fertility in
both M and P lines, as larvae with either higher survival or
those produced by more fecund females will be over-represented
in the group used to start the next generation.

(d) Line extinction
Whenever it was not possible to obtain 20 adults of each sex
despite isolating all available tritonymphs, all emerged adults
were used to establish the next generation. A line was
considered extinct if (i) the total number of individuals sur-
viving to adulthood was less than 5, (ii) no eggs were
produced by females or (iii) all eggs failed to hatch.

(e) Female fecundity and fertility assay
The assay was performed after 14 generations of selection. At
this stage, only three M-HT lines were available for this
experiment, as the other three had gone extinct at genera-
tions 11, 12 and 14. Fecundity of females from each of the
surviving lines was measured at both temperatures
(24°C and 28°C).

Fifteen previously mated females from each selection line
were placed in a single container for 3 days of oviposition.
After this time, the females were discarded. Half of the eggs were placed at 24°C, and the other half at 28°C (‘test temperatures’). Emerging tritonymphs were isolated to individual vials in order to obtain virgin individuals for the fecundity assay. After reaching maturity, females (10–15 per line per test temperature, 429 in total) were placed individually in vials, and each of them was mated to a randomly assigned male from the same line. After 24 h, the male was discarded and replaced by another male from the same line. The second male was replaced 24 h later with a third male, which stayed with the female for the remaining 72 h until the completion of fecundity assay. The purpose of sequential mating with three males was to ensure that individual male effects on female fecundity were minimized: bulb mites do not lay unfertilized eggs, so male sterility would prevent a female from laying eggs if she did not have a chance to remate (however, mating with one fertile partner is sufficient to ensure egg fertility for about one week).

After 5 days of constant access to males, the number of eggs laid by each female was scored. The number of females that had not laid any eggs was also recorded for each line. Bulb mites females oviposit continuously for about two to three weeks at a nearly constant rate; hence, the number of eggs laid during the first 5 days of oviposition is a good estimate of lifetime fecundity [36].

Fecundity of fertile females was analysed in STATISTICA, using a nested ANOVA with mating system, selection temperature and test temperature as fixed factors, and line as a random factor nested in the mating system × selection temperature interaction.

Proportion of fertile females was analysed in R [37], using a generalized linear model with binomial errors (numbers of fertile and infertile females in each line constituted a response variable, whereas mating system, selection temperature and test temperature were explanatory variables).

(f) Trans-generation effects
Although mites used for the fecundity and fertility assay were reared in common environments since the egg stage, their parents developed and mated at their own selection temperatures. The reason for this was that progressing extinction of M-HT lines prompted us to perform the assays without a delay, to make sure that the three surviving M-HT lines were still available. In order to assess the potential trans-generation effects of temperature and mating system on female fecundity and fertility at the stressful environment, we performed a separate experiment.

We placed groups of 50–100 previously mated stock females in common containers for oviposition. We then assigned each container with developing eggs either to the 24°C or to the 28°C treatment. Emerging tritonymphs were isolated individually to obtain virgin adults for the parental generation. In the polygamy treatment, 10 males and 10 females were placed into a single container, such that sexual selection could operate, whereas in the monogamy treatment, males and females were kept in pairs. In both treatments, mites interacted for 5 days. After this period, females from each treatment were placed in fresh containers to lay eggs. Two days later, all eggs were placed to develop at 28°C. Emerging tritonymphs were isolated individually to obtain virgin adults for the fecundity and fertility assay. In this assay, virgin females were placed individually in vials at 28°C, and each was sequentially mated to two randomly assigned males from the same treatment (males were replaced after 24 h; the second male stayed with the female until the completion of the assay). After 5 days of constant access to males, eggs laid by each female were counted. A proportion of fertile females and their fecundity were analysed using logit model and ANOVA, respectively. Significant influence of parental temperature and mating system conditions on female fecundity and fertility would indicate that trans-generation effects of these environmental variables could have also affected female reproductive performance in our main experiment (see §2c).

(g) Data accessibility
The data (EXCEL) are deposited in DRYAD (doi:10.5061/dryad.3s2q0).

3. RESULTS

(a) Female fecundity
We found significant selection temperature × test temperature and mating system × test temperature interaction effects on the number of eggs laid by fertile females (table 1). Given significant interactions, we further analysed the effects of selection temperature and test temperature separately for both the M and P treatments to look for the evidence of thermal adaptation under different mating systems. M-HT lines were less fecund than M-C lines at both temperatures ($F_{1,76} = 6.8, p = 0.032$), and both types of M lines had decreased fecundity at 28°C ($F_{1,113} = 76.3, p < 0.001$; interaction: $F_{1,113} = 1.4, p = 0.232$; figure 1a). In contrast, in polygamous lines, there was a significant selection temperature × test temperature interaction: compared with C lines, HT lines were more fecund at 28°C but less fecund at 24°C (selection temperature: $F_{1,82} = 0.3, p = 0.591$; test temperature: $F_{1,185} = 55.2, p < 0.001$; interaction: $F_{1,185} = 8.7, p = 0.004$; figure 1b). The significant interaction indicates that adaptation to increased temperature had occurred in polygamous lines.

(b) Female fertility
A number of females assayed failed to lay eggs. There was a significant mating system × selection temperature × test temperature interaction effect on the proportion of fertile females ($p = 0.033$; table 2 and figure 2). Both M-C and M-HT lines had lower fertility rates at 28°C ($p < 0.001$; figure 2a). In polygamous lines, however, there was a significant selection temperature × test temperature interaction: polyandrous control (P-C) lines, but not P-HT lines, showed decreased fertility at 28°C ($p = 0.048$; figure 2b).

(c) Line extinction
By generation 17, three more M-HT lines went extinct (following the three lines extinct prior to the fecundity and fertility assay), resulting in 100 per cent of M-HT lines being extinct, compared with no extinctions in the remaining treatments. Thus, mating system significantly affected extinction probability in HT lines (Fisher’s exact test, $p = 0.002$).

(d) Trans-generation effects
Poor performance and extinction of M-HT lines could have resulted from their genetic deterioration or from the detrimental effects of thermal stress carried over to the next generation via parental effects. In an additional experiment testing for such effects, we found that the proportions of fertile matings did not differ between progenies of parents
reared at different temperatures and mating systems: they were 0.84, 0.85, 0.71 and 0.84 for polygamous 28°C, monogamous 28°C, polygamous 24°C and monogamous 24°C treatments, respectively (parental rearing temperature: Wald's $t = 0.64$, $p = 0.425$; mating system: Wald's $t = 0.85$, $p = 0.355$; interaction: Wald's $t = 0.53$, $p = 0.466$).

Similarly, we have found no significant effect of parents' rearing temperature on daughter fecundity ($F_{1,86} = 0.1$, $p = 0.78$; means ± s.e.: polygamy at 28°C, 56.52 ± 5.29; monogamy at 28°C, 46.00 ± 4.06; polygamy at 24°C, 54.00 ± 7.24; monogamy at 28°C, 45.54 ± 4.78). The interaction of parents' rearing temperature × mating system was also non-significant ($F_{1,86} = 0.04$, $p = 0.849$). The effect of parents' mating system was marginally non-significant ($F_{1,86} = 3.1$, $p = 0.081$).

### 4. DISCUSSION

Our data consistently show that mating system strongly affected population performance under a novel selection pressure. At the beginning of the experiment, both monogamous and polygamous populations in the HT treatment faced a high risk of entering an extinction vortex [4,38] where stress-reduced female fecundity and embryonic viability potentially decrease the effective population size, hampering adaptation and increasing inbreeding, generation after generation. Indeed, our data provide a spectacular example of an extinction vortex in progress: by generation 14, half of the monogamous high-temperature (M-HT) populations were extinct, and over 60 per cent of females in the surviving M-HT lines were sterile (figure 2), which was a much higher percentage compared with 15 per cent at the second generation of monogamy exposed to 28°C, as estimated in our trans-generation effects experiment. An additional experiment (see the electronic supplementary material) showed that both sexes in M-HT lines were equally likely to be sterile.

This decline in fertility can be explained by progressing inbreeding, as predicted by the extinction vortex scenario. Indeed, the next extinctions of M-HT lines followed closely, and none of the M-HT lines survived to generation

<table>
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<th>effect</th>
<th>d.f.</th>
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<td>17.149</td>
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<td>1,18.5</td>
<td>6.707</td>
<td>0.018</td>
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<tr>
<td>test temperature</td>
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<td>120.733</td>
<td>&lt;0.001</td>
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<td>mating system × selection temperature</td>
<td>1,18.7</td>
<td>4.069</td>
<td>0.058</td>
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<td>1.933</td>
<td>0.023</td>
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<tr>
<td>mating system × test temperature</td>
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<td>298</td>
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By contrast, all polygamous high-temperature (P-HT) populations survived, and their higher fecundity at 28°C compared with the P-C lines indicates that they have been adapting to elevated temperature. Neither the higher extinction rate of the M-HT lines, nor their lower fertility and fecundity could be attributed to trans-generation effects of increased temperature. An additional experiment, designed to address this possibility, showed that fertility of progeny reared at 28°C and produced by parents exposed to 28°C was over 80 per cent, irrespective of the mating system. Such low infertility levels could not on their own lead to extinction. Importantly, parental environment effects on fertility and fecundity were not significant and also did not interact with mating system. A marginally non-significantly higher fecundity of daughters of polyandrous females replicated earlier results, which demonstrated that this effect was due to genetic, rather than maternal, effects of polyandry [32,36].

Previous work showed that in very small populations (n = 10 individuals), polygamy helps to purge inbreeding depression, decreasing the chances of extinction due to exposure of deleterious mutations under inbreeding: 27 per cent of polygamous versus 49 per cent of monogamous populations went extinct over eight generations [34]. Here, the populations were larger (N = 40), and hence the expected effective population sizes at the onset of the experiment (i.e. before temperature- and inbreeding-induced sterility began to reduce them) were \( N_e = N = 40 \) for monogamy and \( N_e = 8N/(5.3 + 2 + 4) = 28.3 \) for polygamy (assuming random amount of variance in female reproductive success and variance in male reproductive success equal to 5.3, as estimated by Radwan et al. [39]). Earlier experimental evolution studies have shown that populations with \( N_e \) of similar [40] or even lower size [41] can respond to manipulation of mating system. On the basis of effective population sizes in the present study, the inbreeding coefficient after 10 generations (i.e. just before extinctions occurred) under unchanged environment could be estimated at 0.12 and 0.17 for monogamy and polygamy, respectively. This level of inbreeding was much lower than that at population size \( N = 10 \) [34] (0.5 and 0.7 at generation 10 for monogamy and polygamy, respectively) and did not on its own cause extinctions in either monogamous (M-C) or polygamous (P-C) control populations. However, under environmental challenge, the dramatic effects of mating system on population survival were revealed.
One way by which polygamy could positively affect population survival under stress is by alleviating harmful effects of inbreeding, and thus mitigating the risk of entering extinction vortex. Sexual selection has been demonstrated to increase the efficacy of purging inbreeding depression [34], whereas environmental stress tends to magnify deleterious effects of inbreeding [5]. Therefore, sexual selection could allow more effective selection against alleles’ increasing sensitivity to stress when homozygous. Additionally, stress-induced increase in proportion of sterile individuals will have a smaller impact on the effective population size under polygamy, because such a mating system augments the chances that all fertile individuals actually reproduce. This effect could, to some extent, counterbalance the lower $N_e$ of polygamous populations resulting from their higher variance in male reproductive success.

The difference in extinction probability between $M$ and $P$ high-temperature lines may also be due to the effects of sexual selection [7,12], where males carrying alleles beneficial in the novel environment achieve the highest reproductive success, therefore increasing the rate of adaptation [42] (but see [43]). Our results suggest that in contrast to monogamous lines, polygamous lines adapted to increased temperature, as indicated by significant test temperature × selection temperature interactions (figures 1 and 2). Adaptation must have increased the resilience of polygamous populations to thermal stress, thus preventing them from entering the extinction vortex. To date, the only study that looked at thermal adaptation in the context of sexual selection failed to find any effects of mating system [30]. However, the level of polygamy imposed in that experiment was the lowest possible (two mating partners), as was the number of replicate lines. Furthermore, the level of stress imposed by increased temperature in that study was probably lower than in ours, as no cases of extinction were reported.

In conclusion, we have provided the first (to our knowledge) empirical test of the effects of mating system on population extinction risk in the face of environmental challenge. We show that these effects may indeed be dramatic: in our experiment, all monogamous lines went extinct within only 17 generations of thermal stress, whereas the populations where sexual selection was operating have survived and adapted. These results may have important implications for assessing the vulnerability of animal species endangered by human-induced environmental changes.

We thank Joe Tomkins (University of Western Australia), Łukasz Michalczyk (University of East Anglia), Göran Arnqvist (University of Uppsala), William Rice (University of California, Santa Barbara), Eoin Duffy (Jagiellonian University) and two anonymous reviewers for their comments on earlier versions of the manuscript, Magda Jarzębowska for assistance in laboratory work, and Mariusz Klimek for information on soil temperatures. The project was supported by grant no. N303 529438 from the Polish Ministry of Science and Higher Education, and by the Foundation for Polish Science (professor subsidy no. 9/2008 to J.R).

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