Male sperm storage compromises sperm motility in guppies

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Sperm senescence can have important evolutionary implications due to its deleterious effects on sperm quality and offspring performance. Consequently, it has been argued that polyandry (female multiple mating) may facilitate the selection of younger, and therefore competitively superior, sperm when ejaculates from multiple males compete for fertilization. Surprisingly, however, unequivocal evidence that sperm ageing influences traits that underlie sperm competitiveness is lacking. Here, we used a paired experimental design that compares sperm quality between ‘old’ and ‘young’ ejaculates from individual male guppies (Poecilia reticulata). We show that older sperm exhibit significant reductions in sperm velocity compared with younger sperm from the same males. We found no evidence that the brightness of the male’s orange (carotenoid) spots, which are thought to signal resistance to oxidative stress (and thus age-related declines in sperm fitness), signals a male’s ability to withstand the deleterious effects of sperm ageing. Instead, polyandry may be a more effective strategy for females to minimize the likelihood of being fertilized by aged sperm.

1. Introduction

Ageing can have multiple deleterious effects on fitness, extending from the whole organism to cellular-level processes, including gamete function. Post-meiotic sperm senescence is the gradual accumulation of damage during the lifespan of sperm and is caused by a number of factors, most notably oxidative stress attributable to an imbalance in the accumulation of reactive oxygen species. Sperm cells are particularly vulnerable to oxidative damage because they are rich in mitochondria and polyunsaturated fatty acids, and possess a small cytosol with relatively limited capacity for self repair [1–3].

Recent interest from evolutionary biologists in post-meiotic sperm senescence centres around the deleterious effects of sperm ageing on a range of fitness traits, including fertilization ability [4] and offspring performance [5,6]. In particular, the potential for sexual selection to act on female traits that mitigate the costs of mating with males with impaired (aged) sperm has been explored from a theoretical perspective [2,7]. This work suggests that females may benefit either by basing their mating preferences on traits that reveal information about sperm ageing-related quality (precopulatory selection), or by mating multiply to maximize the likelihood that aged (competitively inferior) sperm are unlikely to fertilize their eggs (postcopulatory selection). Both scenarios generate the prediction that sperm senescence is related to sperm quality, but the precopulatory selection hypothesis generates the additional requirement that males advertise their ability to protect sperm from senescence though the expression of sexually selected traits.

In this paper, we test the twofold predictions (i) that sperm ageing has deleterious effects on sperm quality and (ii) that male sexual ornamentation reflects an individual’s ability to mitigate the effects of sperm ageing on sperm quality. We use guppies (Poecilia reticulata), a highly polyandrous livebearing fish, to test both predictions within a paired experimental framework that...
manipulates sperm age during male sperm storage without confounding sperm age with mating history or male age. To address the first aim, we experimentally manipulated sperm age in vivo through repeated sperm (or sham) extractions in order to test how sperm age influences sperm viability (the proportion of live relative to dead sperm in the ejaculate) and sperm swimming velocity (a proxy for sperm competitiveness in guppies [8] and other species [9,10]). To address our second aim, we determined whether carotenoid-based colour pigments reflect a male’s ability to protect sperm quality (viability and velocity) from the deleterious consequences of sperm ageing, and thus whether male secondary sexual traits can reveal information about sperm age-related quality during mate choice. Carotenoid-based colour spots are used by female guppies to assess male attractiveness [11] and are thought to signal resistance to oxidative stress in sperm [12]. Accordingly, males with bright carotenoid-based coloration should be better equipped to protect their sperm from oxidative stress, and hence from the deleterious effect of sperm ageing, than their drab counterparts.

2. Material and methods

(a) Study species and its maintenance

The guppies used in this experiment were descendants of fish captured from Alligator Creek River in Queensland, Australia. Stocks were maintained in mixed-sex groups (approx. 1:1 sex ratio) at 26 ± 1°C and fed a mix of Artemia nauplii and commercial flake food.

(b) Experimental design

Six-month-old males (n = 32) were individually isolated in 1 l plastic tanks prior to the experiment in order to standardize recent mating history and social environment. After 7 days, all males were stripped to empty their sperm reserves (hereafter ‘strip 0’, see figure 1).

The experiment consisted of a paired design in which sperm quality traits from each male were assessed twice under two conditions: ‘young’, where sperm were collected 3 days after the initial strip (strip 0), and ‘old’, where sperm were collected 12 days post-stripping. To balance the order in which young and old sperm were tested, half of the males were tested in the young treatment first, while the other half experienced the old treatment first. Once the first test was concluded the males were rested and housed for 1 week with non-virgin females (which are generally sexually unresponsive but would have maintained the males’ sexual interest). The females were rotated among males so that any difference in the females’ behaviour or attractiveness to the males was minimized. Finally, to standardize the number of times males underwent the stripping procedure (including anaesthesia), a ‘sham’ stripping procedure was performed on day 9 (i.e. the same time as males in the ‘young’ treatment) for males in the old treatment (see figure 1).

(c) Sperm traits assays

We collected ejaculates and assessed sperm quality using standard procedures [13]. Sperm curvilinear velocity (VCL, μm s−1), which predicts competitive fertilization success in guppies [8], was measured using a CEROS sperm tracker (Hamilton-Thorne Research, Beverly, MA, USA). Sperm viability, the proportion of live and dead sperm in the ejaculate, was assessed using a live/dead viability kit (L-7011, Molecular Probes Inc., Eugene, OR, USA) on a minimum of 100 sperm cells from each male.

(d) Colour pattern analyses

The left side of each male was photographed using a digital SLR camera (Nikon D70s) fitted with a 105 mm macro lens (AF-S f/2.8 VR Micro-Nikkor). We used the same camera settings throughout (aperture: f/13, ISO: 200, shutter speed: 1/4 s). Illumination was provided by two bench top fluorescence lamps (each containing 2 × 9 W Osram Dulux bulbs), angled at 45° to reduce specular reflectance. A simulated Gretag Macbeth colour standard with known reflectance properties was included in each image to provide a reference for the imaging software. Each image was captured in RAW format (.NEF file) and subsequently converted to TIFF format for import into the software package Colour Worker (http://www.chrometrics.com). Colour Worker was used to measure the reflectance spectra of the male’s carotenoid spots (orange/yellow coloration) by selecting guppy-specific orange spectra as references (for details see [14]). We took three measures for each spot, which were used to calculate the mean reflectance for each fish. Spot brightness was then measured by calculating the area under the reflectance spectrum (i.e. integrating over the wavelength range 400–710 nm). We also tested the effectiveness of the Colour Worker program in evaluating the brightness of orange coloration by comparing the spectral output of the program with reflectance data obtained using a spectrometer (see the electronic supplementary material).

(e) Statistical analysis

We used paired t-tests to test for differences in sperm motility between sperm of different ages (‘young’ and ‘old’) obtained from the same male. These tests were run separately for sperm velocity and sperm viability (arc sine square root transformed). A linear mixed-effects model (LMM) was used to test whether the brightness of a male’s orange coloration can reflect his ability to mitigate the effects of ageing on sperm quality. In the model,
tive fertilization success in guppies [8], our findings suggest
interactions. As sperm velocity is a determinant of competi-
tion attributable to mating history and male–female social
ability to disentangle the effects of sperm ageing
use natural matings to manipulate sperm ageing, we were
sperm velocity compared with young sperm. As we did not
traits [13,15], we found that old sperm exhibited impaired
and hence accounting for inter-male variability in sperm
comparing old and young ejaculates within the same male,
Our findings reveal that as sperm age inside the male (during
4. Discussion
that males with aged sperm are likely to be less successful
in sperm competition than those that are able to provide
fresh sperm [8]. In guppies, where sperm competition is
intense [16], any decline in sperm competitiveness is likely
to represent a significant reduction in a male’s reproductive
fitness, and may fuel the evolution of male strategies to
limit such costs (e.g. elevated mating rates). Differences in
sperm velocity due to sperm ageing may also explain the
high variability among ejaculates from the same male often
found in experiments [17].

We also investigated whether the link between sperm
ageing and carotenoid coloration may help explain the
female’s preference for colourful males. Specifically, we
tested for a relationship between the brightness of a male’s
orange spots (which are rich in antioxidants) and the ability
to protect sperm from the deleterious effects of ageing
(thought to be caused by oxidative stress; [12]). Evidence sup-
porting such a scenario comes from two studies of birds [18]
and fish [19] where female choice plays a major role in the
mating system. In guppies, males typically circumvent
female choice by using forced copulations, and therefore
females may have limited capacity to rely on precopulatory
mate choice to discriminate among males. However, we
acknowledge that our findings for this portion of the study
are correlational and demand experimental approaches
(e.g. manipulating the amount of antioxidants available to
males) to discount the precopulatory mate choice hypothesis.
Furthermore, it is possible that female guppies use other
mechanisms to select for males with young sperm at the pre-
copulatory level, for example, by choosing successful males
that have a high turnover of sperm and are therefore more
likely to deliver ejaculates containing fresh sperm [20].

In conclusion, we found that male sperm storage has a
deleterious effect on sperm velocity, which is likely to have
important fitness implications for male guppies [8]. Further-
more, we find no evidence that orange brightness signals a
male’s ability to protect his sperm from the deleterious
effect of ageing. This latter finding suggests that any potential
fitness benefits of mating with males with young sperm are
unlikely to be generated by precopulatory female choice.

3. Results
Within the same male, young sperm swam faster than old
sperm (mean \( \mu m s^{-1} \) ± s.e., young sperm: 115.33 ± 2.31,
old sperm: 104.35 ± 2.35, paired \( t \)-test: \( t_{31} = 3.33, p = 0.001 \),
figure 2). However, we found no differences in sperm viabi-
liity (proportion mean ± s.e., young sperm: 0.71 ± 0.03, old
sperm: 0.70 ± 0.04, paired \( t \)-test: \( t_{28} = 0.040, p = 0.968 \))
between treatments. The order in which sperm were tested
(old or young first) had no effect on either of the sperm
traits (both \( p > 0.27 \)).

We found no effect of orange brightness on the relation-
ship between sperm age and sperm swimming velocity
(LMM: treatment: \( t = 2.344, p = 0.026 \), brightness: \( t = 0.855,
p = 0.396 \), treatment \( \times \) brightness: \( t = -1.454, p = 0.157 \))
or sperm viability (treatment: \( t = 1.281, p = 0.211 \), bright-
ness: \( t = 0.972, p = 0.335 \), treatment \( \times \) brightness: \( t = -1.370,
p = 0.182 \)).

4. Discussion
Our findings reveal that as sperm age inside the male (during
male sperm storage), sperm quality becomes impaired. By
comparing old and young ejaculates within the same male,
and hence accounting for inter-male variability in sperm
traits [13,15], we found that old sperm exhibited impaired
sperm velocity compared with young sperm. As we did not
use natural matings to manipulate sperm ageing, we were
able to disentangle the effects of sperm ageing \( \text{per se} \) from
variance attributable to mating history and male–female social
interactions. As sperm velocity is a determinant of competi-
tive fertilization success in guppies [8], our findings suggest

Figure 2. (a) Mean (± s.e.) of sperm velocity (VCL, \( \mu m s^{-1} \)) of young (3 days) and old (12 days) sperm. (b) Relationship between brightness of orange coloration and decline in sperm velocity due to sperm age (negative values indicate a decline in sperm velocity from young to old sperm). (Online version in colour.)
Instead, by mating multiply, females may exploit postcopulatory mechanisms to favour young sperm in the competition to fertilize eggs.

**Ethics statement.** The study was approved by the University of Western Australia Animal Ethics Committee (permit number: RA/3/100/1050).

**Data accessibility.** Data are available at doi:10.5061/dryad.n687q.

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