#### **ORIGINAL ARTICLE**



# Benefit of polyandry in a highly monandrous species when females mate with already mated males

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### Abstract

Female mating frequency varies among animal taxa. A benefit to females of remating has usually been found, but almost all tests have been with polyandrous species. A species being monandrous does not guarantee that mating only once benefits the female, instead the monandry may result from sexual conflict, where her failure to remate benefits her mate, but not her. The parasitoid wasp *Spalangia endius* (Hymenoptera: Pteromalidae) is highly monandrous. Females do not benefit from either immediate or delayed remating when their first mate is virgin. However, some females are likely to mate with already mated males because sex ratios are female-biased. Here, the effect of experimentally induced polyandry on female fitness was examined for females whose first mate had already mated four times, i.e., for fifth females. Fifth female *S. endius* produce significantly fewer daughters than first females. Production of daughters, but not sons, requires sperm in hymenopterans. Fifth females were experimentally induced to mate with a second male, by preventing such females' first mate from providing postcopulatory courtship. The proportion of female offspring produced by these polyandrous fifth females was greater than by monandrous fifth females and not significantly different than by monandrous first females. Total number of offspring did not differ among the three treatments. These results show that there are conditions under which females benefit from polyandry in this highly monandrous species and that the benefit is through effects on offspring sex ratio, not fecundity.

#### Significance statement

Mating frequency varies widely among animals. In most of the more than 100 past studies of insect species in which females mate multiply, females benefit from remating. However, this same question has been addressed in few species in which females mate just once. In the tiny parasitic wasp studied here, females were tricked into remating. Females benefited from remating if their first mate had previously mated multiple times. The benefit was not producing more offspring, but rather being able to produce a greater proportion of daughters. Sons can be produced without sperm in wasps and wasp relatives. Explanations for why females do not normally remate include that they have been manipulated by their first mate and that waiting for a male means sacrificing searching for hosts.

Keywords Monandry · Remating · Polyandry · Mating history · Sex ratio · Sexual conflict

# Introduction

Animals vary widely in their number of mating partners (Krasnec et al. 2012). Given the opportunity, males are usually polygynous, rarely monogynous (Schneider and Fromhage

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B. H. King bking@niu.edu 2010). In contrast, there are many species in which females mate once (monandry) and many in which females mate multiply (polyandry) (Taylor et al. 2014). Among insects, monandry is less common than polyandry (Ridley 1988), and the adaptiveness of monandry remains poorly understood (Arnqvist and Nilsson 2000; Monceau and van Baaren 2012). Monandry may benefit a female and/or for her first mate, but overcoming monandry may benefit her subsequent suitors (Arnqvist and Rowe 2005).

Most experimental tests of the effects of mating frequency on female fitness have been with polyandrous species (Ridley 1988; Arnqvist and Nilsson 2000; Okada et al. 2017). A meta-

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analysis of 122 studies shows a fecundity increase from multiple mating, even in species without nuptial feeding, although mating too many times decreases fitness (Arnqvist and Nilsson 2000). No hymenopterans were included.

Hymenopterans are of special interest because, in many, sperm are necessary for the production of daughters; sperm are not necessary for the production of sons. Thus, even unmated females can produce sons. Whether unmated females produce as many total offspring as mated females varies among species (e.g., Sousa and Spence 2000; King 2002a; Cheng et al. 2004 and references therein; Metzger et al. 2008). Offspring sex ratios of parasitoid wasps tend to be female-biased (Heimpel and Lundgren 2000), which appears to be adaptive, e.g., as described by local mate competition theory (Hamilton 1967; Hardy 1994; King 2002b). If a female fails to mate often enough or mates with male(s) with insufficient sperm, she may be constrained to produce a sex ratio that is more male-biased than is optimal (Charnov 1982; Steiner and Ruther 2009; West 2009; Boulton et al. 2015).

In the present study, the fitness consequence of experimentally induced polyandry was examined in *Spalangia endius*, a parasitic hymenopteran that is highly monandrous. Less than 5% of females copulate with a second male (King et al. 2005). Females not only avoid remating immediately after mating is complete, but also after a dramatic drop in daughter production after 4 days of oviposition (King and Bressac 2010). Sex ratios of *S. endius* appear to be adaptive in conformance with local mate competition theory (King 2002b).

Fitness of females that mate once versus multiply has been examined in only a few highly monandrous insects (Baer and Schmid-Hempel 2005; Arnqvist and Andres 2006; King and Bressac 2010; Jiao et al. 2011). If females of monandrous species benefit from multiple mating like polyandrous species do (Arnqvist and Nilsson 2000), this would suggest that monandry may have evolved because males benefit more by preventing their mates from remating than females do from multiple mating and than males do from mating already-mated females.

The few studies that have examined how a female's fitness is affected by being monandrous versus polyandrous have done so by experimentally manipulating females to remate. Artificial insemination with sperm from more than one male has a negative effect on queen fitness in the bumble bee *Bombus terrestris* (Baer and Schmid-Hempel 2005). Remating has no detectable effect on female fitness in *S. endius*, in the house fly, or in a wolf spider, at least when the males are virgin (King and Bressac 2010; Arnqvist and Andres 2006; Jiao et al. 2011). However, polyandry seems most likely to be beneficial when a female's first mate is sperm-depleted. Thus, the present study of *S. endius* tested whether a female benefits from remating when her first mate has already mated multiple times.

Some *S. endius* females are likely to mate with alreadymated males because there are more females than males (King and Fischer 2010). Female *S. endius* whose mate already mated four times previously, i.e., fifth females, produce the same number of total offspring but fewer daughters than first females do. The present study tests whether such females benefit from remating, either through greater offspring production or through more female-biased sex ratios, and how any benefit compares to having mated with a virgin male initially.

## Methods

#### **Biology of** Spalangia endius

*Spalangia endius* develops within fly pupae that live among decaying organic matter (Rueda and Axtell 1985b). Hosts occur singly or in multiples, and generally one *S. endius* emerges per host; two offspring are produced per host only about 3% of the time (Broski and King 2017). Males begin emerging before females and wait multiple days for females to emerge from nearby hosts (King 2006). After mating, females sometimes have to disperse in order to find fresh hosts, and sometimes fresh hosts will be available nearby (King, personal observation). A *Spalangia* male emerges from his host with his lifetime supply of sperm (Gerling and Legner 1968; King and King 1994). Males provide no nuptial gifts or parental care (King 2002a).

Mated females release an antiaphrodisiac when approached (Mowles et al. 2013), with the result that females that have already mated tend not to be mounted. In addition, if mounted, mated females rarely open their genital orifice (King et al. 2005; King and Fischer 2010). Opening is necessary for copulation. Failure to open appears to result from some aspect of postcopulatory courtship (King and Fischer 2005). Females can be manipulated into a second mating by preventing the first male from providing postcopulatory courtship (King and Bressac 2010). Females that have not received postcopulatory courtship are still attractive and receptive to subsequent males (King and Fischer 2005).

#### **Production of** Spalangia endius

The *S. endius* was a strain originally collected from Zephyr Hills, FL, USA. They were reared at approximately 25 °C with a photoperiod of 12-h light:12-h dark. The host *Musca domestica* was reared so as to produce uniform size host pupae, thereby minimizing variation in body size of the *S. endius* developing in them (King et al. 2014). Parasitized host pupae were individually isolated in glass test tubes ( $12 \times 75$  mm) to obtain male and female *S. endius* that were virgin and newly emerged (1-day-old or less). Matings were observed. Each mating was in a clean container, a plastic dish (35-mm diameter × 10-mm height) that was about two-thirds full of dampened sand to keep humidity high, which reduces static.

#### Effects of remating

I tested whether remating by fifth females allowed them to produce a greater proportion of daughters than not remating and whether remating allowed them to produce as many or more daughters as if they had mated with a virgin male initially. Fifth females were used to match a previous study of the same parasitoid species and to be within the range of how many times a male is likely to mate in nature (King and Fischer 2010). There were three treatments (N = 16 per treatment): monandrous fifth females, polyandrous fifth females, and monandrous first females.

- 1) In the monandrous first treatment, females were each presented with a single virgin male, and mating was observed.
- 2) In the monandrous fifth treatment, females were each presented with a male that had already mated with four other virgin females, and mating was observed.
- 3) In the polyandrous fifth treatment, females were each presented with a male that had already mated with four other virgin females, and mating was observed. However, this mating was interrupted: the male was removed immediately after he withdrew his aedeagus from the female, i.e., just before postcopulatory court-ship would normally begin. Then, the female was presented with a second male, this time a virgin male, and the second mating was allowed to complete naturally.

About 10 min after the last mating, each female was presented daily for 7 days with 20 hosts and with honey ad libitum, a total of 140 hosts per female. This is less than the mean of about 80 offspring (maximum of about 120) that a female produced in her lifetime in a previous experiment (unpublished analysis of first experiment in King and Bressac 2010). King (2002a) reported that no female produced offspring from all 20 hosts in a day.

The hosts had pupated within the last 2 days when they were presented to females. One week of hosts was chosen as being sufficient to show meaningful effects on daughter production if they exist. Offspring sex ratio in the first 7 d of oviposition differs significantly between fifth females and first females, with fifth females producing more sons and fewer daughters (unpublished analysis of King and Fischer 2010). Offspring production drops dramatically within the first week for both types of females, as a result of a drop in daughter production (King and Bressac 2010; King and Fischer 2010). Adult offspring were allowed to complete development, emerge from their hosts, and die. Then, because some adults crawl back into a host, the hosts were dissected, and the number of adult sons and the number of adult daughters were counted. The person counting the wasps was blind to what the treatments were.

#### Statistical analyses

Analyses were with SPSS (IBM Corp 2016). Total number of sons and total number of daughters were each summed across all 7 days for each mother. From those, total offspring production and the proportion of daughters were calculated for each mother. Residuals of a one-way ANOVA (analysis of variance) on offspring production were highly kurtotic and skewed as assessed by a Shapiro-Wilk test; the same was true for the proportion of daughters. Thus, Krukal-Wallis tests, which are nonparametric, were used instead. Post hoc pairwise comparisons of treatments were by an ANOVA when residuals were normally distributed, and by a Mann-Whitney *U* test otherwise. Conclusions would have been unaffected if a Bonferroni correction had been applied to control for multiple comparisons (Rice 1989).

Data availability The data generated and analyzed during the current study are available in the Harvard Dataverse repository, King BH, 2017, "KingBH\_FL\_Spalangia\_ endius\_1stf\_5thf\_5th&1stf.xlsx", doi:https://doi.org/10. 7910/DVN/NL1EM3, Harvard Dataverse, V1, UNF:6: URbK01VzI+rrr1T/IGxZuA==.

# Results

The number of offspring produced per day in the present study was 0 to 25, and the lifetime number was 11–123. There was no significant effect of treatment on offspring production (Fig. 1; Kruskal-Wallis  $\chi^2 = 3.82$ , df = 2, P = 0.15), but there was an effect of treatment on sex ratio (Fig. 2; Kruskal-Wallis  $\chi^2 = 12.20$ , df = 2, P = 0.002). Sex ratio did not differ between monandrous first females and polyandrous fifth females (F = 0.35, df = 1, 30, P = 0.56), but the proportion of daughters produced by those two treatments combined was more female-biased than that of monandrous fifth females (Mann-Whitney U = 98.0, n = 32, n = 16, P = 0.001). How offspring production and sex ratio changed over time in each treatment is shown in Figs. 3 and 4, respectively.

# Discussion

Unlike the lack of a benefit to polyandry when *S. endius* females have first mated with a virgin male (King and Bressac 2010), polyandry was beneficial for females that had mated with a male that had previously mated multiple times. By remating, fifth females increased their proportion of daughters so much that their offspring sex ratios became as female-biased as those of females that had mated only once but with a virgin male (Figs. 2 and 4). These results are consistent with previous suggestions that the cost of monandry



Fig. 1 Total number of offspring added across all 7 days of oviposition for first monandrous, fifth monandrous, and fifth polyandrous mothers

and benefit of polyandry is most likely to be in females whose first mate has mated multiply before her (Jones 2001; Montrose et al. 2004). In contrast to the effects of remating on sex ratio, but as expected, offspring production was not significantly affected by female mating status, specifically, by whether a female was a monandrous first female, a monandrous fifth female, or a polyandrous fifth female (Figs. 1 and 3).

Given the apparent advantage to a female from remating, three non-mutually exclusive hypotheses may explain why *S*. *endius* females do not normally remate. One explanation is



**Fig. 2** Total number of sons (white circle) and total number of daughters (black circle) added across all 7 days of oviposition for first monandrous, fifth monandrous, and fifth polyandrous mothers



**Fig. 3** Total number of offspring among first monandrous (increment), fifth monandrous (white circle), and fifth polyandrous (black circle) mothers during each of the 7 days of oviposition

that remating would reduce offspring production by interfering with females beginning to burrow in search of hosts (Rueda and Axtell 1985a; Geden 2002). To remate, a female would have to instead stay on the surface, because that is where males are (King 2002a).

A second explanation is interparty conflict and cooperation. Whether a female remates can be influenced by the behavior of at least three parties: the female, her first mate, and subsequent suitors (Chapman et al. 2003; King and Bressac 2010). The three parties' fitness-interests may or may not all coincide (Parker 1979; Arnqvist and Rowe 2005). The present study suggests that, provided the female's first mate has recently mated multiply, she will benefit from remating, through more daughters allowing her to produce adaptive sex ratios (Hamilton 1967; King 2002b). How the fitness of her first and second mate is affected will depend on patterns of sperm usage in this species. The pattern of sperm precedence is context-dependent in confamilials of S. endius (van den Assem and Feuth-DeBruijn 1977; Khanh et al. 2005) and is unknown in S. endius. The second S. endius mate's expected benefit in our experiment was likely at least as great as the difference in the number of daughters that fifth polyandrous females produce minus the number of daughters that fifth monandrous females produce. That females do not typically remate may indicate that during the species' recent evolutionary history, the benefits of remating to the female and to her subsequent suitors have not exceeded the benefit of monandry to the first male.

A third explanation is that the frequency at which females encounter and mate with males that have already mated four times may be low. Given the sex ratio in this species, males are expected to mate one and a half to three times on average (King and Fischer 2010), and reproductive success is highly variable among males even in the laboratory (King and Napoleon 2006). Fig. 4 Number of sons (white circle) and number of daughters (black circle) among first monandrous, fifth monandrous, and fifth polyandrous mothers during each of the 7 days of oviposition



Thus, some males will have more success than average. The risk of mating with a multiply-mated male will be lower in *S. endius* when virgin males are also present. This is because virgin males are quicker than mated males to approach females, with the result that females usually mate with virgin males when they are present (King and Fischer 2010).

The lack of increased fecundity from multiple mating in the present study of *S. endius* is consistent with the other studies of highly monandrous species (Baer and Schmid-Hempel 2005; Arnqvist and Andres 2006; Jiao et al. 2011), including King and Bressac's (2010) study of *S. endius*, which used only virgin male. In contrast, in *S. endius*, a sex ratio benefit from remating is seen only when a female's first mate has mated multiply (the present study), not when he is virgin (King and Bressac 2010).

The number of polyandrous parasitoid wasps that have been studied is small, but most do not show a fecundity benefit from multiple mating (Chevrier and Bressac 2002; Cheng et al. 2004; Jacob and Boivin 2005; Khanh et al. 2005; but see mixed results in Boulton and Shuker 2015). This contrasts with a fecundity benefit from moderate multiple mating by females in many other animals, even species without nuptial feeding (Arnqvist and Nilsson 2000). So far, only one of five polyandrous parasitoid wasps show a sex ratio benefit from multiple mating (Chevrier and Bressac 2002; but see Cheng et al. 2004; Jacob and Boivin 2005; Khanh et al. 2005; Boulton and Shuker 2015). The two species where longevity effects have been examined show a positive effect in some conditions (Jacob and Boivin 2005; Boulton and Shuker 2015). There is not yet evidence of indirect genetic benefits in parasitoid wasps (Boulton and Shuker 2015), but indirect benefits have been seen in some other animals (Slatyer et al., 2012). It is important to keep in mind that effects of a given mating system on fitness may not be the same as when that mating system first evolved or first reevolved in a population. This is because when mating frequency changes, male and female reproductive traits evolve (Katvala et al. 2008; Kraaijeveld et al. 2009). Some of these traits could in turn affect the current benefit of different mating frequencies, e.g., sperm storage, male ejaculation quality and quantity, and timing of dispersal or timing of the switch from attraction to sex pheromones to attraction to host pheromones.

To better understand the evolution of female mating frequency, more studies of highly monandrous species are needed, including tests of effects of indirect benefits. Together, the two studies of effects of multiple mating on female fitness in *S. endius* demonstrate the importance of examining fitness effects on sex ratio, not just fecundity, and demonstrate the importance of examining effects when males have already mated, not just when they are virgin (the present study; King and Bressac 2010).

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## Compliance with ethical standards

**Ethical approval** All procedures were in accordance with the ethical standards of the institution at which the studies were conducted.

**Conflict of interest** The author declares that she has no conflict of interest.

**Informed consent** Informed consent was not required as no human participants were involved.

## References

- Arnqvist G, Andres JA (2006) The effects of experimentally induced polyandry on female reproduction in a monandrous mating system. Ethology 112:748–756
- Arnqvist G, Nilsson T (2000) The evolution of polyandry: multiple mating and female fitness in insects. Anim Behav 60:145–164
- Amqvist G, Rowe L (2005) Sexual conflict. Princeton University Press, Princeton, NJ
- Baer B, Schmid-Hempel P (2005) Sperm influences female hibernation success, survival and fitness in the bumble-bee *Bombus terrestris*. Proc Roy Soc Lond B Bio 272:319–323
- Boulton RA, Shuker DM (2015) The costs and benefits of multiple mating in a mostly monandrous wasp. Evolution 69:939–949
- Boulton RA, Collins LA, Shuker DM (2015) Beyond sex allocation: the role of mating systems in sexual selection in parasitoid wasps. Biol Rev 90:599–627
- Broski SA, King BH (2017) Effects of size and age of the host *Musca domestica* (Diptera: Muscidae) on production of the parasitoid wasp *Spalangia endius* (Hymenoptera: Pteromalidae). J Econ Entomol 110:282–287
- Chapman T, Arnqvist G, Bangham T, Rowe L (2003) Sexual conflict. Trends Ecol Evol 18:41–47
- Charnov EL (1982) The theory of sex allocation. Princeton University Press, Princeton, NJ
- Cheng LI, Howard RW, Campbell JF, Charlton RE, Nechols JR, Ramaswamy SB (2004) Mating behavior of *Cephalonomia tarsalis* (Ashmead) (Hymenoptera: Bethylidae) and the effect of female mating frequency on offspring production. J Insect Behav 17:227–245
- Chevrier C, Bressac C (2002) Sperm storage and use after multiple mating in *Dinarmus basalis* (Hymenoptera: Pteromalidae). J Insect Behav 15:385–398
- Geden CJ (2002) Effect of habitat depth on host location by five species of parasitoids (Hymenoptera: Pteromalidae, Chalcididae) of house flies (Diptera: Muscidae) in three types of substrates. Environ Entomol 31:411–417
- Gerling D, Legner EF (1968) Developmental history and reproduction of *Spalangia cameroni*, parasite of synanthropic flies. Ann Entomol Soc Am 61:1436–1443
- Hamilton WD (1967) Extraordinary sex ratios. Science 156:477-488
- Hardy ICW (1994) Sex ratio and mating structure in the parasitoid Hymenoptera. Oikos 69:3–20
- Heimpel GE, Lundgren JG (2000) Sex ratios of commercially reared biological control agents. Biol Control 19:77–93
- IBM Corp. Released (2016) IBM SPSS statistics for Windows, Version 24.0. IBM Corp, Armonk, NY
- Jacob S, Boivin G (2005) Costs and benefits of polyandry in the egg parasitoid *Trichogramma evanescens* Westwood (Hymenoptera: Trichogrammatidae). Biol Control 32:311–318
- Jiao XG, Chen ZQ, Wu J, Du HY, Liu FX, Chen JA, Li DQ (2011) Male remating and female fitness in the wolf spider *Pardosa astrigera*: the role of male mating history. Behav Ecol Sociobiol 65:325–332
- Jones TM (2001) A potential cost of monandry in the lekking sandfly, Lutzomyia longipalpis. J Insect Behav 14:385–399

- Khanh HDT, Bressac C, Chevrier C (2005) Male sperm donation consequences in single and double matings in Anisopteromalus calandrae. Physiol Entomol 30:29–35
- King BH (2002a) Breeding strategies in females of the parasitoid wasp Spalangia endius: effects of mating status and body size. J Insect Behav 15:181–193
- King BH (2002b) Sex ratio response to conspecifics in a parasitoid wasp: test of a prediction of local mate competition theory and alternative hypotheses. Behav Ecol Sociobiol 52:17–24
- King BH (2006) Mate location and the onset of sexual responsiveness in the parasitoid wasp *Spalangia endius* (Hymenoptera: Pteromalidae). Environ Entomol 35:1390–1395
- King BH, Bressac C (2010) No fitness consequence of experimentally induced polyandry in a monandrous wasp. Behavior 147:85–102
- King BH, Fischer CR (2005) Males mate guard in absentia through extended effects of postcopulatory courtship in the parasitoid wasp *Spalangia endius*. J Insect Physiol 51:1340–1345
- King BH, Fischer CR (2010) Male mating history: effects on female sexual responsiveness and reproductive success in the parasitoid wasp *Spalangia endius*. Behav Ecol Sociobiol 64:607–615. https:// doi.org/10.1007/s00265-009-0878-3
- King BH, King RB (1994) Sex ratio manipulation in response to host size in the parasitoid wasp *Spalangia cameroni*—is it adaptive? Behav Ecol 5:448–454
- King BH, Napoleon ME (2006) Using effects of parasitoid size on fitness to test a host quality model assumption with the parasitoid wasp *Spalangia endius*. Can J Zool 84:1–5
- King BH, Saporito KB, Ellison JH, Bratzke RM (2005) Unattractiveness of mated females to males in the parasitoid wasp *Spalangia endius*. Behav Ecol Sociobiol 57:350–356
- King BH, Colyott KL, Chesney AR (2014) Livestock bedding effects on two species of parasitoid wasps of filth flies. J Insect Sci 14:185
- Kraaijeveld K, Franco P, Reumer BM, van Alphen JJM (2009) Effects of parthenogenesis and geographic isolation on female sexual traits in a parasitoid wasp. Evolution 63:3085–3096
- Krasnec MO, Cook CN, Breed MD (2012) Mating systems in sexual animals. Nat Edu Knowl 3:72 https://www.nature.com/scitable/ knowledge/library/mating-systems-in-sexual-animals-83033427. Accessed 14 January 2018
- Metzger M, Bernstein C, Desouhant E (2008) Does constrained oviposition influence offspring sex ratio in the solitary parasitoid wasp *Venturia canescens*? Ecol Entomol 33:167–174
- Monceau K, van Baaren J (2012) Female teneral mating in a monandrous species. Ecol Evol 2:1426–1436
- Montrose VT, Harris WE, Moore PJ (2004) Sexual conflict and cooperation under naturally occurring male enforced monogamy. J Evol Biol 17:443–452
- Mowles SL, King BH, Linforth RST, Hardy ICW (2013) A female-emitted pheromone component is associated with reduced male courtship in the parasitoid wasp *Spalangia endius*. PLoS One 8:e82010
- Okada K, Suzaki Y, Sasaki R, Katsuki M (2017) Fitness costs of polyandry to female cigarette beetle *Lasioderma serricorne*. Behav Ecol Sociobiol 71. https://doi.org/10.1007/s00265-017-2316-2
- Parker GA (1979) Sexual selection and sexual conflict. In: Blum MS, Blum NA (eds) Sexual selection and reproductive competition in insects. Academic Press, London, pp 123–166
- Rice WR (1989) Analyzing tables of statistical tests. Evolution 43:223-225
- Ridley M (1988) Mating frequency and fecundity in insects. Biol Rev 63: 509–549
- Rueda LM, Axtell RC (1985a) Effect of depth of house fly pupae in poultry manure on parasitism by six species of Pteromalidae (Hymenoptera). J Entomol Sci 20:444–449

- Rueda LM, Axtell RC (1985b). Guide to common species of pupal parasites (Hymenoptera: Pteromalidae) of the house fly and other muscoid flies associated with poultry and livestock manure, Technical Bulletin 278. North Carolina Agricultural Research Service, North Carolina State University
- Schneider J, Fromhage L (2010) Monogynous mating strategies in spiders. In: Kappeler P (ed) Animal behaviour: evolution and mechanisms. Springer, Berlin, Heidelberg
- Slatyer RA, Mautz BS, Backwell PR, Jennions MD (2012) Estimating genetic benefits of polyandry from experimental studies: a metaanalysis. Biol Rev Camb Philos Soc 87:1–33
- Sousa JM, Spence JR (2000) Effects of mating status and parasitoid density on superparasitism and offspring fitness in *Tiphodytes*

gerriphagus (Hymenoptera: Scelionidae). Ann Entomol Soc Am 93:548-553

- Steiner S, Ruther J (2009) How important is sex for females of a haplodiploid species under local mate competition? Behav Ecol 20:570–574
- Taylor ML, Price TA, Wedell N (2014) Polyandry in nature: a global analysis. Trends Ecol Evol 29:376–383
- van den Assem J, Feuth-De Bruijn E (1977) Second matings and their effect on the sex ratio of the offspring in *Nasonia vitripennis* (Hymenoptera: Pteromalidae). Entomol Exp Appl 21:23–28
- West SA (2009) Sex allocation. Princeton University Press, Princeton, NJ