

Polyandry in the ectoparasitic copepod *Lepeophtheirus salmonis* despite complex precopulatory and postcopulatory mate-guarding

Christopher D. Todd^{1,*}, Rebecca J. Stevenson¹, Helena Reinardy¹, Michael G. Ritchie²

¹School of Biology, Gatty Marine Laboratory, University of St Andrews, St Andrews, Fife KY16 8LB, UK

²School of Biology, Sir Harold Mitchell Building, University of St Andrews, St Andrews, Fife KY16 9TH, UK

ABSTRACT: *Lepeophtheirus salmonis* (Krøyer) is an economically important pest on cultured salmonids in the North Atlantic, and has been implicated in declines of some wild salmonid populations. Males inseminate newly-moulted adult females following the cementing of a pair of spermatophores to the female's genital complex. Females can produce multiple pairs of eggstrings over a period of months, but this species is reported to be monogamous as a result of blockage of the female's copulatory ducts by the spermatophore tubules. On wild and farmed Atlantic salmon *Salmo salar* L., respectively, 88 and 78% adult females sea lice bore the typical pair of spermatophores, while 11 and 19% lacked spermatophores. A very few individuals (1% wild, 3% farmed) bore 3 or 4 spermatophores, showing that apparently successful multiple mating is possible. Multiple paternity was confirmed by dual-locus microsatellite typing of offspring for 3 of 7 females carrying 4 spermatophores, but also for 2 of 3 females carrying a single pair of spermatophores. Probably most females on wild fish lose their initial spermatophores and are polygamous during their extended ovigerous lifetime, although effective blockage of the copulatory ducts by the first male almost certainly assures single paternity of the first few pairs of eggstrings. The total level of polyandry or sperm competition faced by males may be relatively low. The ecological implications of multiple paternity are discussed within the context of integrated pest management as a strategy of ameliorating *L. salmonis* infestations impacting both wild and farmed salmonids.

KEY WORDS: Paternity · Microsatellite · Spermatophore · Sea lice · Aquaculture · Salmonid · Integrated pest management

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INTRODUCTION

The specialist ectoparasitic copepod *Lepeophtheirus salmonis* (Krøyer) infests all species of wild salmonids in the North Pacific (Nagasawa 2001) and occurs with 100% prevalence on wild adult salmon *Salmo salar* L. in the North Atlantic (Jacobsen & Gaard 1997, Todd et al. 2000). This ectoparasite species, and to a lesser extent the generalist *Caligus elongatus* Nordmann, remains a major problem to the Atlantic salmon aquaculture industry. At sufficient intensities *L. salmonis* can debilitate or kill the host fish, and the circumstantial evidence that sea lice are at least a major contributory factor to the recent crashes of wild sea trout *Salmo*

trutta L. stocks in parts of Ireland, Scotland and Norway is now considerable (e.g. Butler 2002, Todd et al. 2004). There is good evidence that *L. salmonis* may be detrimental to wild Atlantic salmon smolts migrating through certain Norwegian fjords (Finstad et al. 2000). Within the broader framework of integrated pest management (IPM), control of sea lice in Scottish aquaculture has focused largely on chemotherapeutants (e.g. organophosphates, pyrethroids, avermectins, hydrogen peroxide) in concert with periodic fallowing of farm sites, single year-class culture within sea lochs and co-ordinated sea lice treatments amongst some farms in voluntary area management agreements (Grant 2002, Lindsay & Rae 2003).

*Email: cdt@st-and.ac.uk

Despite these measures, the most recent global estimate of the total economic cost of sea lice to the aquaculture industry exceeds 100 million US \$ per annum (Johnson et al. 2004), and for the Scottish industry alone is at least £25 million per annum (Rae 2002). Effective pest-species control and management demands detailed knowledge of the parasites's life history and population biology, and *Lepeophtheirus salmonis* has been intensely studied (e.g. Boxshall & Defaye 1993, Pike & Wadsworth 1999, Tully & Nolan 2002), especially in the farm environment (e.g. Revie et al. 2002). The typical pattern for *L. salmonis* populations infesting wild Atlantic salmon is one of adult (~90%) and female (~70%) predominance (Jacobsen & Gaard 1997, Todd et al. 2000), because of an extended life span of ovigerous females. Adult female caligids store sperm in a seminal receptacle (Huys & Boxshall 1991), have internal fertilisation, and produce multiple pairs of uniseriate eggstrings over their reproductive life. The female retains each eggstring pair and the hatching stage is a free-swimming nauplius larva. After 2 larval moults during an obligatory planktonic phase of 2 to 9 d (Johnson & Albright 1991), the infective planktonic copepodid stage is attained and colonisation of the host fish results in the first of 4 sessile chalimus stages. The fourth chalimus moults to the first of 2 mobile preadult stages, and both genders undergo a final moult to the mature adult. Adult male caligids actively locate and attach to Preadult II females for hours or days, and await the definitive adult female moult prior to copulation (Boxshall 1990, Ritchie et al. 1996a). Such precopulatory 'proximate mate-guarding' (e.g. Simmons & Siva-Jothy 1998) appears to be a defensive behaviour by the male. At copulation, the male cements a pair of spermatophores onto the posterior ventral surface of the female's genital complex and remains clasped postcopula for some hours after spermatophore deposition (Ritchie et al. 1996a). Males take up to 4 d to synthesise spermatophores following copulation (Hull et al. 1998). Tubules extend from the attached spermatophores and cross the midline to enter the female copulatory pores on opposite sides (see Huys & Boxshall 1991, p. 274, for *L. pectoralis*). Emptying of the spermatophore occurs within hours and it is reported that the combination of inner secretions of the spermatophores, the spermatophore tubules themselves, and the male's cement all effectively block the female copulatory ducts and prevent subsequent mating (Ritchie et al. 1996b). Simmons & Siva-Jothy (1998) term such behaviour 'remote post-copulatory mate-guarding'.

Ritchie et al. (1996a) deduced that *Lepeophtheirus salmonis* is monogamous, and Pike & Wadsworth (1999) concluded that although singly-mated female *L. salmonis* could produce multiple pairs of fertile egg-

strings, there was no information to date about female *L. salmonis* showing multiple mating. Given the importance of a detailed understanding of the mating system for any commercially or ecologically important pest species, and in view of the prevalence of multiple mating throughout the animal kingdom (e.g. Birkhead & Møller 1998, Neff & Pitcher 2002) and probable benefits of multiple mating to females (Arnqvist & Nilsson 2000, Tregenza & Wedell 2000), we sought to empirically assess whether or not *L. salmonis* is truly monogamous. Ritchie et al. (1996b) reported that non-virgin female *L. salmonis* (from farmed salmon) without spermatophores are very rarely found, and also that gravid females were never observed with an empty seminal receptacle. There is no observational evidence, but it is likely that female *L. salmonis* are incapable of removing the blocking empty spermatophores and cement, and will therefore presumably remain effectively monogamous for at least as long as these remain on the genital complex.

The typical and maximum life spans of female *Lepeophtheirus salmonis* either on farmed or wild Atlantic salmon remain unknown, but almost certainly extend to several months (Pike & Wadsworth 1999, Tully & Nolan 2002), during which time perhaps up to 11 pairs of eggstrings may be produced (Heuch et al. 2000). Adult female *L. pectoralis* (parasitising plaice *Pleuronectes platessa*) can survive for up to 10 mo (Boxshall 1974). Anstensrud (1990) showed that spermatophores of *L. pectoralis* remained cemented to the female for a mean of 26 d (range 2 to 43 d), and that females can be copulated more than once, but only after the spermatophores have been lost. The deduction was that male *L. pectoralis* can only temporarily seal the female copulatory ducts as a result of spermatophore deposition. Female *L. salmonis* may differ markedly from *L. pectoralis* in perhaps retaining their spermatophores throughout reproductive life and remaining monogamous; but assuming that spermatophore retention/loss is similar in *L. salmonis* and *L. pectoralis*, monogamy of *L. salmonis* evidently still is possible if (1) the female copulatory ducts remain permanently blocked following first insemination, (2) all potential eggstrings can be fertilised by sperm stored in the seminal receptacle from the one mating, and (3) if old adult females show an increasing tendency to have lost the spermatophores. Because the last is unlikely to apply (Ritchie et al. 1996a, see last paragraph) this implies that multiple mating of *L. salmonis* may occur commonly, despite laboratory observations of a strong mating preference of males for virgin adult females (Hull et al. 1998). There has been a perception in the Scottish industry that aquaculture sites are liable to self-reinfestation by *L. salmonis* (e.g. Bron et al. 1993, Costello 1993, see also

Heuch et al. 2003). If this were indeed a significant feature of the infection dynamics of *L. salmonis* on farms then confirmation of obligate female monogamy would have potentially important implications for the possible inclusion of interference of male fertility within a wider IPM strategy.

Our objectives were 2-fold. First, to compare spermatophore occurrence on adult female *Lepeophtheirus salmonis* from both wild and farmed Atlantic salmon, and second, to ascertain paternity of embryo clutches for individual females from wild hosts. The first few pairs of eggstrings clearly will be fertilised by the initial male but, given the extended life span of adult female *L. salmonis*, and the observations of spermatophore loss in *L. pectoralis*, it was considered likely that *L. salmonis* females can lose their spermatophores and be multiply mated during their ovigerous lifetime. However, whether any such subsequent matings result in successful fertilisation by sperm of a second male (P_2 fertilisation, e.g. Simmons & Siva-Jothy 1998) remains unclear. Because of their commonly high levels of polymorphism, single-locus DNA microsatellites have long been the method of choice in parentage analyses (e.g. Strassmann et al. 1996). Herein we applied dual-locus genotyping of samples of embryos from single eggstrings for 10 female *L. salmonis*, using specifically developed microsatellites (Todd et al. 2004).

MATERIALS AND METHODS

Spermatophore numbers and frequency. Ethanol-preserved adult female *Lepeophtheirus salmonis* from wild and farmed Atlantic salmon were examined for spermatophores. Bulk samples (not separated for each host fish) were obtained from 2 Scottish salmon farms. One farm sample (June 1998; Farm 3 of Todd et al. 2004) was from the mainland coastline of Wester Ross, and the other from the Western Isles (September 1997; Farm 5 of Todd et al. 2004). Complete collections of *L. salmonis* from individual wild Atlantic salmon were obtained from the commercial coastal nets at Strathy Point, north Scotland (58° 06' N, 04° 00' W) in July 1999, 2002, 2003 and 2004.

Female *Lepeophtheirus salmonis* from wild salmon were distinguished as 'young' and 'old' adults, according to the degree of expansion of the genital complex

(see Ritchie et al. 1996a,b), which proceeds in newly moulted adult females irrespective of copulation. The Preadult I and II female stages were distinguished for the Strathy Point 2002 to 2004 samples, but in 1999 only total preadult females were recorded. All adult females were scored as bearing 0 to 4 spermatophores. Females carrying the typical single pair of spermatophores (see Huys & Boxshall 1991, p. 279) are referred to here as 'paired spermatophore' individuals. Individuals bearing an additional 1 or 2 spermatophores generally had these cemented along the outer edge of the initial pair of spermatophores; these we refer to as 'multiple spermatophore' females.

Microsatellite typing of females and their families. Multiple spermatophore females clearly had been multiply mated, and 7 such females were specifically chosen for microsatellite genotyping. Our primary requirements were to assess whether or not (1) specifically chosen females that had unequivocally mated multiply, and (2) randomly chosen females that carried the typical pair of spermatophores, were monogamous or producing polyandrous clutches of offspring. In 3 instances of multiply mated females (Females 4, 5, 6), we randomly sampled from the same respective fish another ovigerous, paired-spermatophore female (Females 1, 2, 3; Table 1).

For microsatellite PCR, a variable number of offspring embryos (5 to 10) from both the proximal (attached) and distal (free) ends of 1 of the 2 available eggstrings were isolated for each of 10 females. Of the 7 multiple spermatophore females, 4 (Nos. 4, 5, 6, 9) had broken eggstrings: for these the individual embryos were isolated from both the proximal and the broken end, but were known to not have been maxi-

Table 1. *Lepeophtheirus salmonis*. The 10 females and single eggstrings used for paternity analysis. All were sampled from returning wild adult Atlantic salmon *Salmo salar* netted at Strathy Point, North Scotland. For host fish, year of sampling, fish identifier and host sea-age (1SW, 2SW = 1/2 sea-winter[s]) are given. For female parasites, condition of sampled eggstring, egg complement (intact eggstrings only; na = not available), number of spermatophores borne and numbers of offspring genotyped (positions within eggstring) are shown

Female sea louse	Year/ Fish no./ Sea-age	Egg-string	Total egg no.	No. of spermatophores	No. offspring typed (position)
1	2001/02/2SW	Intact	202	2	10 (5 proximal + 5 distal)
2	2001/39/1SW	Intact	445	2	10 (5 proximal + 5 distal)
3	2001/28/1SW	Intact	528	2	10 (5 proximal + 5 distal)
4	2001/02/2SW	Broken	na	4	10
5	2001/39/1SW	Broken	na	4	10
6	2001/28/1SW	Broken	na	4	10
7	1999/33/1SW	Intact	261	4	16 (8 proximal + 8 distal)
8	2001/31/1SW	Intact	429	4	20 (10 proximal + 10 distal)
9	2002/59/1SW	Broken	na	4	20
10	2002/63/1SW	Intact	325	4	20 (10 proximal + 10 distal)

mally separated within the original eggstring. For Females 1–3 (2 spermatophores) and 7, 8 and 10 (multiple spermatophores), the sampled offspring were contiguous and equally balanced from the 2 ends of the intact eggstring (Table 1) to provide offspring genotypes for the first and last fertilised embryos within those eggstrings.

PCR was undertaken for genomic DNA of individual embryos from other females in a preliminary study, with extraction by means of either a NucleoSpin (Abgene) kit or by the 'single fly' preparation (Gloor & Engels 1992). The latter methodology consistently resulted in the clearer PCR products and was subsequently adopted throughout. Female parent tissue for PCR genotyping was obtained by excising an appropriately small piece of the anterior cephalothorax and also by removal of a maxilliped. These were extracted with the same protocol as the individual embryos and the single limb was found to consistently provide the clearer PCR products. PCR for all families included primers for 2 highly polymorphic *Lepeophtheirus salmonis* microsatellites (see Todd et al. 2004 for primer sequences and details of PCR amplifications). *LsalSTA3* (GenBank Accession No. AY509256) and *LsalSTA5* (AY509258), respectively, had shown 38 and 43 alleles in a previous analysis of population structuring of *L. salmonis* from throughout the North Atlantic. PCR products were visualised by PAGE silver-staining and alleles were sized against 10 bp ladders and against a previously genotyped individual as a positive control.

Statistical analyses. Spermatophore frequencies for the separate years and for 'wild' and 'farmed' *Lepeophtheirus salmonis* were analysed for heterogeneity by the *G*-test. For families with apparently single paternity, statistical power was assessed by means of the Monte Carlo simulation model of Neff & Pitcher (2002). The probability of detecting multiple paternity within a clutch (Neff & Pitcher's *PrDM*) is a function of the numbers of (1) loci used, (2) alleles in the population, (3) offspring genotyped, and (4) effective males. Moreover, males may have differential fertilisation success. Neff & Pitcher's programme is dimension-limited to 30 alleles per locus. For present purposes we deleted alleles with the lowest frequencies from our original Atlantic frequency distributions for the 2 loci (Todd et al. 2004), and thereby reduced population diversity from 38 to 26 alleles (for *LsalSTA3*) and 43 to 27 alleles (for *LsalSTA5*),

respectively. Population allele frequencies were recalculated for the 2 truncated distributions and included in the model. Each simulation included the known maternal genotype for the 2 loci and 2 (genetically unknown) males were presumed for each set of offspring analysed; the probability of detecting dual paternity was computed assuming both equal (0.5:0.5) and highly skewed (0.9:0.1) fertilisation success of 2 males. The tabulated probabilities are the averages for 10 runs of the model.

RESULTS

Spermatophore numbers and frequency

Most adult female *Lepeophtheirus salmonis* from wild and farmed host fish bore a pair of spermatophores (Table 2), cemented in the typical posterior-ventral position on the genital complex. It is generally impossible to visually assess whether females carrying the typical paired spermatophores were (1) singly mated, or (2) had possibly lost their initial spermatophores plus cement—perhaps some weeks after copulation (see Anstensrud 1990 for *L. pectoralis*)—and been multiply copulated. Some individuals evidently had lost one of the pair, as indicated by traces of remaining male cement. Few females lacked spermatophores, but many of these also showed traces of cement, indicating that they had been inseminated at least once.

A very low percentage of females from both wild and farmed salmon did, however, carry multiple (3 or 4) spermatophores, and almost certainly had been mated by at least 2 separate males. There was significant heterogeneity amongst the 4 yr for wild salmon (Table 2; $G = 15.09$, 6 df, $p = 0.020$): 2002 to 2004 did not differ significantly from one another, but 1999 differed significantly from all 3, due to fewer than expected individuals lacking spermatophores in 1999. There was no significant heterogeneity between the 2 farm samples

Table 2. *Lepeophtheirus salmonis*. Adult females from wild and farmed Atlantic salmon *Salmo salar*, showing % (frequency) bearing 0 to 4 spermatophores

Spermatophores	Strathy Point (wild fish)					Farm (fish)		
	1999 (n = 42)	2002 (n = 53)	2003 (n = 64)	2004 (n = 62)	Pooled (n = 221)	5 (n = 14)	3 (n = 11)	Pooled (n = 25)
0	8.0 (60)	13.0 (81)	11.8 (107)	10.3 (112)	10.7 (360)	22.4 (26)	12.0 (6)	19.3 (32)
1 or 2 (‘paired’)	90.5 (677)	86.4 (538)	87.2 (788)	89.1 (971)	88.4 (2974)	74.1 (86)	86.0 (43)	77.7 (129)
3 or 4 (‘multiple’)	1.5 (11)	0.6 (4)	1.0 (9)	0.6 (6)	0.9 (30)	3.4 (4)	2.0 (1)	3.0 (5)

($G = 3.04$, 2 df, $p = 0.219$), although the sample sizes are small. Comparison of the pooled frequencies for parasites sampled from wild versus farmed hosts showed significant heterogeneity ($G = 15.56$, 2 df, $p < 0.001$), with farm lice showing more than expected females lacking spermatophores or carrying multiple spermatophores.

Multiple spermatophore individuals almost invariably were old adult females (fully expanded genital complex), although 1 young adult female (small genital complex) was recorded in this category for 2002. Distinguishing the wild adult females into 'young' and 'old' (Table 3) showed that whilst approximately one-third of young adult females remained virgin, only 8% of old females were lacking spermatophores. The latter had not, however, all recently undergone the definitive moult from the Preadult II stage because, for example, 13% of the 87 old adult females lacking spermatophores in 2003 bore both cement and copulation marks from the male appendages on their genital complex. Together, these features strongly indicate that these 'old' females had been previously mated but had lost their spermatophores.

Microsatellite typing and analysis of paternity

For *LsalSTA3* and *LsalSTA5*, respectively, 15 and 18 different alleles were detected amongst the 10 females and their families (Table 4): 3 other alleles at *LsalSTA5* were recorded as maternal. Multiple paternity was confirmed among the families for 2 of the 3 females bearing the typical pair of spermatophores.

Table 4. *Lepeophtheirus salmonis*. Family samples for 10 females genotyped at each locus and for both loci. *PrDM* (Neff & Pitcher 2002) calculated for families for which single paternity was apparent: values are average probabilities (10 model simulations) of detecting dual paternity, assuming males with either equal (0.5:0.5) or strongly biased (0.9:0.1) mating success. nd: no data (failed PCR). No. typed (no. n-m alleles): no of offspring typed (no. of non-maternal alleles detected); *PrDM*: probability of detecting multiple paternity (modelled male paternity ratio)

Female no.	No. typed (no. n-m alleles)			No. of spermatophores	Inferred paternity	<i>PrDM</i>	
	<i>LsalSA3</i>	<i>LsalSA5</i>	Both loci			(0.5:0.5)	(0.9:0.1)
1	6 (2)	10 (3)	6	2	Dual	–	–
2	8 (3)	9 (2)	8	2	Dual	–	–
3	10 (2)	nd	nd	2	Single	0.906	0.524
4	8 (3)	9 (4)	8	4	Dual	–	–
5	9 (2)	9 (2)	9	4	Single	0.987	0.589
6	10 (2)	10 (2)	10	4	Single	0.991	0.628
7	8 (3)	15 (3)	8	4	Dual	–	–
8	18 (2)	18 (2)	18	4	Single	0.999	0.832
9	16 (4)	16 (3)	16	4	Dual	–	–
10	20 (2)	20 (2)	20	4	Single	0.999	0.862
Total	113 (15)	116 (18)					

Table 3. *Lepeophtheirus salmonis*. Adult females from wild Atlantic salmon *Salmo salar* netted at Straithy Point over 4 yr period, showing % (frequency) of 'young' and 'old' lacking (without) or bearing (with) spermatophores

Year	– Percent young (n) –		— Percent old (n) —	
	without	with	without	with
1999	25.6 (11)	74.4 (32)	7.0 (49)	93.0 (656)
2002	38.9 (35)	61.1 (55)	8.6 (46)	91.4 (487)
2003	30.3 (20)	69.7 (46)	10.4 (87)	89.6 (751)
2004	24.3 (43)	75.7 (134)	7.6 (69)	92.4 (843)
Pooled	29.0 (109)	71.0 (267)	8.4 (251)	91.6 (2737)

The test of paternity for the third family in this category (Female 3) was, however, weak due to the lack of data for *LsalSTA5* (failed PCR). The single inferred male for Family 8 was homozygous at *LsalSTA3* and shared an allele at that locus with Female 8.

For Females 1 and 2 (Table 4) single paternity was implied for 1 of the 2 loci, and yet dual paternity was confirmed by the alternative locus, showing that the males again shared alleles at 1 locus despite the potentially high allele numbers. Neff & Pitcher's *PrDM* for Female 3 was high (>90%), but only when assuming equal mating success of 2 putative fathers. For the strongly skewed simulation, the probability of detecting multiple paternity for these 10 offspring fell to 52%. For Females 8 and 10, confidence in single paternity was consistently high for both simulations because of the larger families screened.

The relative frequencies of the male alleles (both loci) for each family of offspring are shown in Fig. 1. In all 5 cases of multiple paternity—2 of 3 paired spermatophore females (Nos. 1, 2) and 3 of 7 multiple spermatophore (Nos. 4, 7, 9) individuals—all offspring could be genetically explained by just 2 males. Although the family samples are small it is important to note that, for the multiply mated females, the least successful paternal allele always had a very low frequency (~0.08 for both loci; Fig. 1). If these females had mated only twice (no families showed >4 paternal alleles), and males had equal fertilisation success, this frequency would be expected to approximate 0.25. Hence, male fertilisation success probably is not equal in this species, but there are numerous processes (variation in sperm depletion or quality, insemination rates, sperm competition, cryptic female choice) which could be responsible for this pattern.

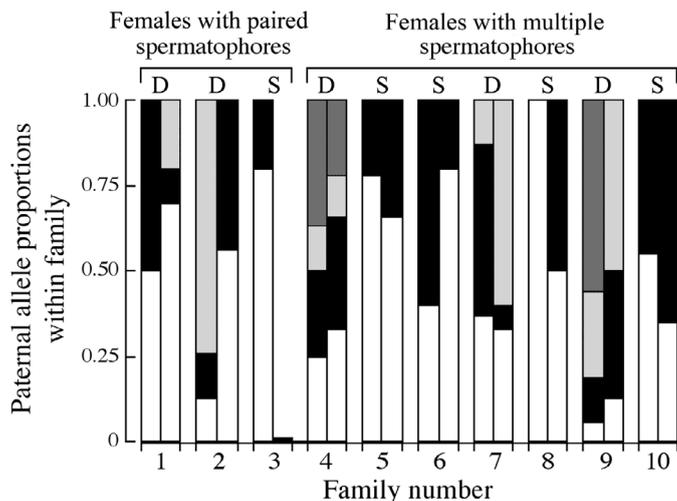


Fig. 1. *Lepeophtheirus salmonis*. Proportional frequencies of male alleles for *LsaSTA3* (left-hand bar) and *LsaSTA5* (right-hand bar) for offspring families of Females 1 to 10. Shading shows number and proportions of male alleles for each family; similar shading does not imply identical alleles across families; 1 or 2 non-maternal alleles = single paternity, S (Females 3, 5, 6, 8, 10); 3 or 4 non-maternal alleles = dual paternity, D (Females 1, 2, 4, 7, 9). Females 1 to 3 bore single pair of spermatophores; Females 4 to 10 bore 4 spermatophores. Data for *LsaSTA5* for Family 3 missing due to failed PCR

DISCUSSION

Multiple mating and paternity

If the cemented spermatophores of *Lepeophtheirus salmonis* are successful in preventing subsequent male insemination, the simplest explanation of the paternity data is as follows: (1) Of the 7 multiply mated females, 4 (Females 5, 6, 8, 10) still bore their original spermatophores and a subsequent pair. Their copulatory pores remained blocked by the original pair, and despite having been multiply mated they were effectively still monogamous. (2) For the remaining 3 multiply mated females (Females 4, 7, 9), the spermatophores borne were the second and third pairs, and dual paternity of offspring was attributable to the first (lost) and second pairs of spermatophores. The second pair did, however, still block the copulatory pores from the most recently acquired third pair. (3) Of those bearing the typical single pair of spermatophores, Female 3 had been mated only once, whereas Females 1 and 2 bore a replacement second pair, having lost the initial pair of spermatophores and were producing offspring which had been successfully fertilised by the second male (P_2 fertilisation).

The higher percentage of adult females from farmed hosts carrying no spermatophores (19%; Table 2) was unexpected, given observations of the often very

marked predominance of males on farmed salmon (e.g. Bron et al. 1993) in contrast to female predominance on wild salmon (e.g. Todd et al. 2000), and the presumed reduced life span of parasites on farmed fishes. Despite 29% of young adult females on wild salmon being unmated (Table 3), it is very probable that within the 8 d (at 12°C; Ritchie et al. 1996a) required for full expansion of the genital complex and the transformation of an adult female from 'young' to 'old', all females on wild fish will become inseminated. The likelihood is, therefore, that old adult females lacking spermatophores are seldom, if ever, virgin, but will be previously inseminated individuals that have lost both the spermatophores and their cement, and may well already be polyandrous.

Numerical predominance of females over males appears widespread in populations of free-living calanoid copepods (e.g. Cuoc et al. 1997, Bathélémy et al. 1998). Because the mobile preadult and adult stages of *Lepeophtheirus salmonis* almost certainly do not transfer between wild host fish (but see Ritchie 1997 for hosts in farm pens), the individual fish is the effective demographic 'unit' for these ectoparasites. It clearly is possible that on a given fish there may be insufficient adult males to pair with all the reproductively 'available' female complement (Preadult II females, plus unfertilised young adult females and 'old' adult females lacking spermatophores): this could therefore lead to incidences of old virgin adult females. Every wild fish sampled at Strathy Point in 2002 to 2004 bore 1 or more adult male sea lice (range 1 to 25), and always adult males on a given fish numerically exceeded Preadult II females (data not shown). Similarly, the number of adult males almost always exceeded the complement of females reproductively available to those males on a given fish (Fig. 2). Predictions of male:male competition, and of female-mating frequency, in both free-living and parasitic copepods may, therefore, be erroneous if such expectations are based on population proportions of total males and females. For *L. salmonis* specifically, the general surfeit of adult males over available females on individual fish, and the consistent occurrence of a large proportion of unfertilised young adult females (Table 3), shows that successful fertilisation of adult females does not always occur as soon as they undergo the final moult from the Preadult II stage. The rare occurrence of multiple spermatophores (Table 2) does, however, confirm that adult females already bearing spermatophores may very occasionally be multiply mated. Taken together, these data, and the low frequency of old adult females lacking spermatophores (Tables 2 & 3), all strongly indicate that multiple mating is a common feature in *L. salmonis*, despite clear laboratory behavioural preferences by adult males for the younger female stages (Hull et al. 1998).

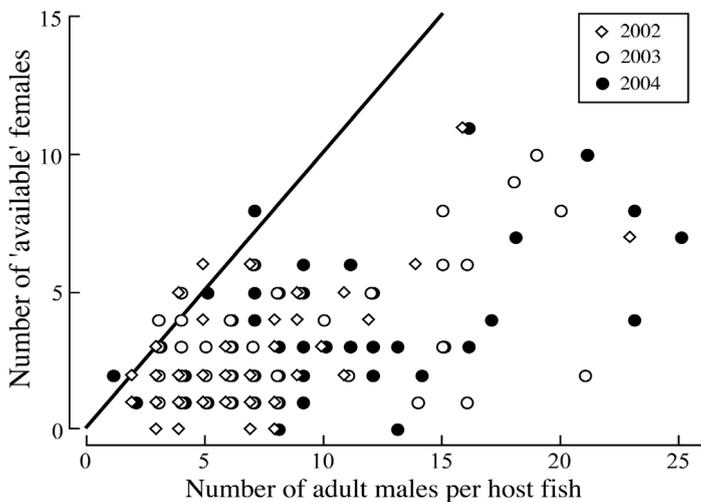


Fig. 2. *Lepeophtheirus salmonis* on wild Atlantic salmon *Salmo salar* for period 2002 to 2004 inclusive ($n = 179$ fish; Table 2). Scatterplot of 'available' females (= Preadult II females + 'young' adult females [0 spermatophores] + 'old' adult females [0 spermatophores]) on adult males per host fish. Fitted line indicates equality of numbers of males and 'available' females. Abscissa values for Years 2002 and 2004 offset for clarity

Further data for both wild and farmed host fish are required, but if the observed two-thirds of paired spermatophore females (Nos. 1 and 2) showing dual paternity is representative, it is likely that most females on wild and farmed salmon will be polyandrous. Probably only those females that fail to survive beyond loss of the initial spermatophores and their cement will be effectively monogamous. In the farm environment, therefore, routine chemotherapy and its consequential restriction of the adult life span might result in effective monogamy of females. Spermatophores on a female are visible evidence of the outcome of precopulatory competition amongst various males on a host fish; but the finding of multiple paternity of offspring does not, in itself, necessarily reflect postcopulatory male competition, sperm competition, or even cryptic female choice (e.g. Eberhard 1998). Only detailed manipulative studies of the copulation and fertilisation history of individual females, and of changes in individual male availability through their ovigerous lifetimes, will reveal the nature and extent of such interactions. It will, for example, be of especial importance to ascertain the extent of initial sperm depletion prior to loss of the first pair of spermatophores in relation to possible sperm displacement and competition, male precedence, and the mean and variance of second male paternity (P_2 fertilisation, e.g. Simmons & Siva-Jothy 1998, Simmons 2001).

The temporal and energetic investment by male caligid in proximate mate-guarding of the Preadult II

female, and the subsequently inseminated young adult female, is relatively small—a matter of hours, or perhaps days (Boxshall 1990, Ritchie et al. 1996a). Conversely, the durability of the attached spermatophores and cement provides the individual male with extended, cheap and effective remote postcopulatory mate-guarding (cf. Hull et al. 1998), perhaps for some weeks, during which the (initially monogamous) female may produce several pairs of eggstrings. Insect mating plugs do generally represent effective short-term barriers to re-insemination (Simmons 2001), but for the majority of insects (which have a single female genital opening), long-term barriers might also prevent the female from ovipositing and thereby be detrimental to both male and female reproductive success. In sharp contrast to most insects, however, female caligid copepods have separate paired copulatory ducts and gonopores through which the fertilised eggstrings are extruded. Effective first-male blockage of the female does not, therefore, compromise either first male or female fitness in *Lepeophtheirus salmonis*. Nonetheless, loss of the blockage does allow a degree of polygamy.

Monogamy and pest control

IPM approaches to pest control are deliberately multifaceted and, amongst other benefits, aim to reduce reliance on insecticides and the evolution of treatment resistance by the target organism. The sterile insect technique (SIT) typically involves the rearing and sterilisation of males by irradiation before their subsequent release to the environment (e.g. Curtis 1985); SIT has proved to be a successful means of controlling certain terrestrial pest species (e.g. Ferguson et al. 2005), especially when applied within a broader IPM strategy (e.g. Dargie 2000, Twohey et al. 2003). An effective contribution to pest control by sterile male release (SMR) in the aquatic environment is, however, restricted to attempts to control sea lampreys *Petromyzon marinus* parasitising lake trout and lake whitefish in the Laurentian Great Lakes (e.g. Twohey et al. 2003). Migrating male lampreys are trapped, chemically sterilised by individual injection and then released. An attractive element of the SMR approach is its potentially minimal wider environmental effect (Siefkes et al. 2003). Of the licensed sea louse chemotherapeutants available to the Scottish industry, only 2 (cypermethrin, emamectin—both of which non-specifically interfere with the neural membrane function) are widely used. Teflubenzuron, a chitin synthesis inhibitor, which is accordingly ineffective against the (non-moulting) adult stage, is in use only intermittently (Grant 2002). It is therefore understandable that

the industry has concerns over possible self-reinfestation of farm sites and development of resistance to the few compounds available to them. Whilst the direct experimental evidence of resistance of sea lice to chemical treatments (e.g. dichlorvos, Jones et al. 1992; deltamethrin, Sevatdal et al. 2005) is equivocal, there is a perception within the industry that this can be a problem. An IPM strategy that ameliorates larval sea lice production, and that minimises dependence on agents to which resistance may evolve, could be advantageous to the aquaculture industry.

Monogamy of a pest species renders it especially vulnerable to SMR, but it can have an adverse consequence in genetic control if already-mated females immigrate from without the treatment area (Curtis 1985), or if females begin to avoid sterile males (Ferguson et al. 2005). Unlike many insect pests, which may be highly vagile, adult *Lepeophtheirus salmonis* are effectively sedentary (and thereby manageable and readily treatable) on their host fish in captivity, because it is only the planktonic larval stages that emigrate and immigrate. For polyandrous caligid copepods, SMR, or deployment of male sterilitant treatments, might yet offer a plausible addition to current sea lice IPM by the aquaculture industry if extremely high sterile:wild ratios of males could be attained on fish within the treatment area. It will never be feasible to rear (on live fish) and release the required numbers of manipulated males for SMR approaches; and, because of their poor abilities to reattach to host fish (Pike & Wadsworth 1999), released adult males almost certainly will not establish themselves on farms in suitable numbers. Because populations of *L. salmonis* on salmon farms are demographically open (Todd et al. 2004), it will never be possible to eradicate this pest at aquaculture sites. However, it may be possible to develop a specific, biologically or photochemically degradable male sterilitant. The wider environmental impact of such an agent could perhaps be reduced by its targeted application as an in-feed additive, but application would have to be regular in order to continually sterilise males as they develop on the fish from the immigrating infective copepodid larval stages.

Our previous study of Atlantic and Pacific *Lepeophtheirus salmonis* (Todd et al. 2004) revealed that levels of gene flow and cross-infection amongst the different host species and between wild and farmed salmonids are sufficiently high throughout the North Atlantic to prevent significant population genetic differentiation on an ocean-wide scale. Planktonic larvae of *L. salmonis* exported by currents from a given farm site may well infect both adjacent farms and local populations of wild salmonids, in addition to perhaps ultimately being re-imported and re-infesting the

natal farm site. High levels of gene flow (= larval colonisation) between wild and farmed salmonids, together with the considerable numerical imbalance between wild and cultured salmonids in the coastal waters of Scotland (e.g. Butler 2002), lead to the conclusion that the bulk of the larval flux of *L. salmonis* is probably from farmed to wild hosts. Notwithstanding the logistic and practical difficulties in achieving SMR in the aquatic environment, the incorporation of male sterilisation of *L. salmonis* within a commercially beneficial and environmentally benign IPM strategy—including co-ordinated area management agreements amongst farm sites—may therefore have significant mutual benefits for the welfare of both wild and farmed salmonid stocks. The effectiveness of the contribution to control by sterilising male parasites as they develop on the host fish would also be enhanced if chemotherapeutants could be used to ensure that ovigerous females generally did not survive beyond loss of their initial spermatophores.

From an ecological perspective, only experimental manipulation and long-term maintenance of individual copepods on isolated host fish will allow a full understanding of the possible roles of sperm depletion, competition, displacement or stratification, and patterns of male sperm precedence or cryptic female choice in this economically and environmentally important pest species. The key finding—that polyandry occurs and probably is widespread amongst female *Lepeophtheirus salmonis*—is important to the modelling of the dynamics and the management of caligid infestations on cultured salmonids (e.g. Revie et al. 2002), and requires quantification in the farm environment. Polyandrous mating of adult females also has implications for the evolution and possible spread of resistance to chemotherapeutant treatments on an ocean-wide scale. Furthermore, the fact that polyandry is frequent despite post-copulatory mate-guarding in this species should urge caution regarding inferences of single or multiple matings (e.g. Cuoc et al. 1997, Bathélémy et al. 1998) in other free-living copepods based on anatomical observations alone.

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