

# Male Reproductive Success and Multiple Paternity in Wild, Low-Density Populations of the Adder (*Vipera berus*)

SYLVAIN URSENBACHER, CECILE ERNY, AND LUCA FUMAGALLI

From the Laboratoire de Biologie de la Conservation, Département d'Ecologie et Evolution, Biophore, Université de Lausanne, Switzerland (Ursenbacher, Erny, and Fumagalli); and the School of Biological Sciences, University of Wales—Bangor, Bangor LL57 2UW, UK (Ursenbacher).

Address correspondence to S. Ursenbacher, Department of Environmental Sciences, Section of Conservation Biology, University of Basel, St Johannis-Vorstadt 10, CH-4056 Basel, Switzerland, or e-mail: [s.ursenbacher@unibas.ch](mailto:s.ursenbacher@unibas.ch).

We studied for the first time the occurrence of multiple paternity, male reproductive success, and neonate survival in wild, low-density adder (*Vipera berus*) populations using 13 microsatellite loci. Paternity was assigned for 15 clutches, collected during 3 years. Our data demonstrated that multiple paternity can occur at a high level (69%) in natural populations of *V. berus*, even if the density of adults is low. The high proportion of multiple sired clutches was comparable to the proportion observed in captive populations. Male reproductive success significantly increased with body length, and only the largest males successfully sired entire clutches. Finally, no relationship was detected between the number of fathers per clutch and neonate survival. These results suggest that multiple matings could be beneficial in populations with high level of inbreeding or low male fecundity.

**Key words:** Male reproductive success, multiple paternity, reptile, *Vipera berus*

Multiple paternity has been documented in numerous invertebrate and vertebrate taxa (Birkhead and Møller 1998). This occurs even though a single copulation is usually sufficient to fertilize an entire clutch (Orsetti and Rutowski 2003; but see Madsen et al. 1992). In the absence of direct benefits for females, such as paternal care, indirect genetic advantages might outweigh the costs associated with polyandry (e.g., Simmons 2005). It has been hypothesized that the genetic benefits of polyandry in such species is the acquisition of more good genes as well as an increase in genetic diversity within clutches (for a review, see Yasui 1998). For males, the benefit of multiple copulations with different females is generally an increase in fecundity, especially in species without postcopulatory investments (Garner et al. 2002). However, the search for females is time and energy consuming, and population density could have

an impact on mating opportunities. Consequently, low-density populations show sometimes reduced levels of multiple paternity (Garner et al. 2002; but see also Gromko and Gerhart 1984; Ochando et al. 1996). In addition, male reproductive success is often related to different morphological characters, such as body size as observed in mammals (e.g., Le Boeuf and Reiter 1988) or bird species (e.g., Bryant 1988; Grant PR and Grant BR 2000).

Multiple paternity studies and male reproductive success are numerous for birds and mammals (see Reynolds 1996; Birkhead and Møller 1998), whereas snakes have been less studied (Höggren and Tegelström 1995; Garner et al. 2002; Prosser et al. 2002).

The adder (*Vipera berus*) is a viviparous snake widely distributed throughout northern Eurasia. The female reproductive cycle is bi- or triennial, therefore the operational sex ratio is biased toward males (Madsen and Shine 1993). Males actively court females 3–4 weeks per year. During this limited period, vigorous combats for access to receptive females occur and are generally won by the largest males (Madsen 1988; Madsen and Shine 1992; Luiselli 1993b). Short-term postcopulatory mate guarding is common (Andrén 1986); however, multiple matings (with the same or different male) do occur (Andrén and Nilson 1987; Madsen et al. 1992; Luiselli 1993a), mainly due to sneaky males (males which copulate with females while the other are fighting). Although the presence of a “copulatory plug” preventing further mating has been suggested by Nilson and Andrén (1982), multiple paternity has been observed in captivity (Stille et al. 1986; Höggren and Tegelström 1995, 2002). Based on molecular techniques, Höggren and Tegelström (2002) have shown that the first male to mate with a female sired the highest number of her offspring. Contrary to several other snake species, long-term sperm storage in adders is unlikely (Luiselli 1993a; Höggren and Tegelström 1996). Moreover, newborn viability in wild

populations seems to be related to the number of copulations with different mates (Madsen et al. 1992). Madsen et al. argued that multiple copulations increase the genetic quality of the offspring and that sperm competition improves the quality of the newborns. However, direct relationship between the number of copulation and offspring viability in adder is debated (Capula and Luiselli 1994).

In order to investigate the occurrence of multiple paternity in the wild, we used hypervariable microsatellite markers to determine paternity and tested the influence of male body length on their reproductive success. Father assignment and breeding success in *V. berus* have only been documented in captive, high-density conditions. In natural, low-density populations, we rather expected a limited level of multiple paternity due to a presumed difficulty of finding mates. In addition, we investigated whether the number of fathers per clutch impacted neonate survival rate.

## Methods

### Sampling

Snakes were caught by hand, measured (snout–vent length [SVL]), weighed, marked with a microchip (DataMars SA, Bedano, Switzerland) and tissue samples collected in the field (50  $\mu$ l of blood) over a 3-year period (2000–2002) from 2 distinct populations in the Swiss Jura mountains (Populations CH1 and CH2 in Ursenbacher et al., [forthcoming]). These 2 populations (separated by about 1.5 km) are considered to be isolated from each other because no migration was observed between 1997 and 2004 and significant genetic differentiation was detected ( $F_{ST} = 0.041$ ,  $P < 0.005$ ). Moreover, the estimated population densities are very low (about 1 adult/ha, Ursenbacher and Monney 2003), whereas populations in the Alps usually harbor 3–11 adults/ha (see Neumeier 1987; Luiselli 1993b). To obtain clutches, all pregnant females captured just before parturition were maintained in individual field terrariums (60  $\times$  70  $\times$  50 cm). After birth, offspring were measured and tissue samples were collected. The neonate survival rate was estimated for each clutch within 24 h after birth in order to compare our results with the study of Madsen et al. (1996). All neonates and mothers were then released at their exact capture point. Only one clutch per female was analyzed.

### Microsatellite Amplification and Parentage Analyses

DNA extraction and amplification of 13 microsatellite loci have been conducted following Ursenbacher et al. (forthcoming). Parentage assignments were performed by a likelihood approach using the program CERVUS 2.0 (Marshall et al. 1998), which is particularly sensitive to the proportion of candidate adults sampled (Jones and Ardren 2003). This proportion was consequently estimated using the module CAPTURE (Otis et al. 1978) from the software MARK 3.2 (White and Burnham 1999). The proportion of candidate adults sampled was estimated for each population and each year between 1997 and 2002. The mean percentage of

captured males was estimated to 82% using the model M[t] in CAPTURE. Estimation of genotyping errors was performed by repeating the genotyping procedure 2–5 times for samples showing doubtful results, by counting incompatibilities between the genotypes of the mother and her offspring, and by CERVUS which calculates the ratio of the number of mismatches to the number compared, scaled by the average probability of detecting a mismatch. Because the 2 former approaches probably give an over- and underestimation, respectively, genotyping error rate of 1.6% estimated by the latter approach was used in the CERVUS simulations to assess assignment reliability. Finally, the occurrence of null (nonamplifying) alleles, which might appear in some of the 13 microsatellite markers (see Ursenbacher et al. forthcoming), was evaluated by CERVUS. Testing for the presence of null alleles is important because they can cause false exclusions in the assignments (see Jones and Ardren 2003). Although CERVUS can estimate their frequency for each locus and accommodate for their occurrence, Jones and Ardren (2003) suggest considering all homozygous genotypes as heterozygotes possessing the null allele, in order to prevent false exclusions. Paternity assignments were performed with and without this adjustment and results were compared.

### Reproductive Success of Males

The relationship between SVL of males and their annual reproductive success was tested via regression. The SVL was also compared between fathers and males without assigned offspring as well as between fathers detected in monoparental and in multiple paternity clutches (both using *t*-tests or Mann–Whitney *U* tests depending on the normality of the data). Because some males were not captured during the year, they succeeded in mating; SVL was evaluated for all fathers a posteriori, using the following approach: The SVL was corrected by 0.01 cm per day between the mean date of the mating and the nearest capture (when the capture was done the year before, only 210 days were added due to hibernation). For juvenile males, a growth rate of 0.05 cm per day was used up to the maturity (37 cm). These growth rates were based on capture–recapture data obtained in the 2 populations (Ursenbacher S, unpublished data). We also tested whether the number of fathers for a clutch had an impact on number of offspring or neonate survival rate via linear regression with an angular transformation. In order to avoid pseudoreplication (multiple presence of a male in a test), each year was tested separately or only the first detected paternity was taken into account.

## Results

### Parentage Assignments

Fifteen pregnant females (7 in the first population, 8 in the second) were captured and gave birth to 115 offspring (Table 1); 43 males (25 in the first population, 18 in the second) were considered as potential fathers. Between 2

**Table 1.** Reproductive females, number of offspring, estimated number of fathers within the clutch, number of offspring not assigned, and morphometric measurements of the female before laying

Population	Female code	Number of offspring	Number of fathers in the clutch ( $P \geq 80\%$ )	Number of offspring not assigned ( $P < 80\%$ )	Female SVL (cm)	Female weight (g)	Number of dead offspring
1	16	2	2 (2)	0 (0)	56.0	128	1
1	17	8	3 (2)	0 (1)	58.4	202	7
1	32	9	2 (2)	0 (0)	52.2	155	3
1	<i>48</i>	<i>10</i>	<i>5 (4)</i>	<i>5 (4)</i>	<i>50.9</i>	<i>124</i>	<i>4</i>
1	59	8	2 (2)	0 (0)	53.9	144	3
1	<i>66</i>	<i>13</i>	<i>8 (5)</i>	<i>5 (5)</i>	<i>50.0</i>	<i>137</i>	<i>0</i>
1	85	4	3 (2)	0 (0)	51.5	120	0
2	21	7	4 (4)	1 (1)	52.0	119	1
2	38	7	1 (1)	0 (0)	53.5	108	0
2	41	8	3 (3)	0 (0)	51.9	143	4
2	70	7	3 (4)	2 (1)	49.0	121	1
2	421	10	1 (1)	0 (0)	48.9	114	10
2	442	7	1 (1)	0 (0)	50.0	93	0
2	448	8	3 (3)	0 (0)	51.3	104	0
2	453	7	1 (1)	0 (0)	47.4	98	0
	Mean	$7.7 \pm 2.5$	$2.8 \pm 1.9$ ( $2.2 \pm 1.0$ ) <sup>a</sup>	$0.87$ ( $0.23$ ) <sup>a</sup>	$51.8 \pm 2.83$	$127.3 \pm 27.1$	

In brackets, the results obtained without the correction suggested by Jones and Ardren (2003). In clutches of female 48 and female 66 (italic), less than 70% of the neonates were assigned to a father, and both clutches were consequently not used in the analyses of multiple paternity and male behavior.

<sup>a</sup> Mean without females 48 and 66. Standard deviations for each means are indicated.

(D12) and 13 (D6) alleles were observed. CERVUS detected null alleles in 5 of the 13 loci (Vb-A8, Vb-B1, Vb-B18, Vb-D6, and Vb-D'13; Ursenbacher et al. forthcoming). Discrepancies in paternity assignment with and without the correction suggested by Jones and Ardren (2003) occurred for 9 neonates only. Paternity was assigned to 102 (99 without correction) of the 115 offspring sampled with 80% or 95% confidence, the majority (56%; 45% without correction) at the 95% level (Table 1). Ten (11 without the correction) of the 13 (16) unassigned offspring belonged to 2 clutches from females living at the edge of the first population. Parental assignment in these 2 clutches was low (50% and 62%, respectively), and thus, they were excluded from the behavioral analyses below.

### Male Reproductive Success and Multiple Paternity

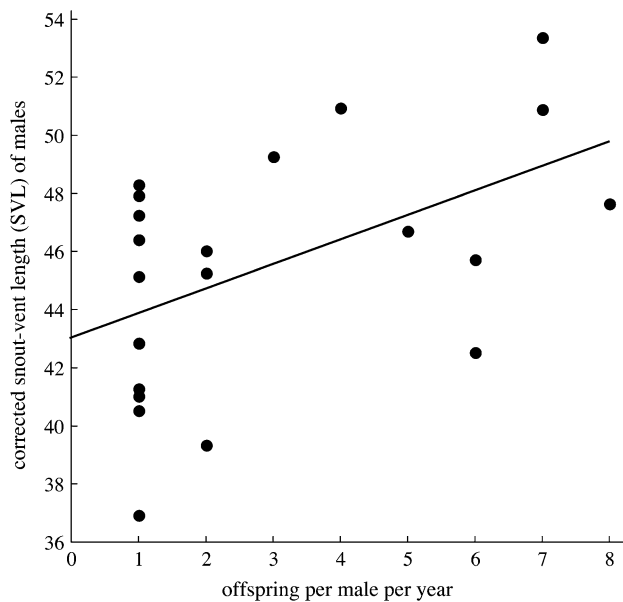
Only 4 clutches (31%) were fathered by a single male, and all belonged to the second population. For remaining clutches, 3 were fertilized by 2 males, 5 clutches by 3 males, and 1 clutch by 4 different males (Table 1). The number of fathers per clutch was not significantly different between the 2 studied populations ( $\text{mean}_{\text{pop1}} = 2.4$ ;  $\text{mean}_{\text{pop2}} = 2.1$ ;  $t$ -test:  $t_{11} = 0.460$ ,  $P = 0.65$ ). Twenty-one different fathers (18 without correction) were detected in the 13 clutches, 5 (5) of them in 2 different clutches and 1 (2) in 3 clutches (sometimes across different years). Fathers that sired offspring in more than one clutch were significantly larger (SVL corrected) than other fathers ( $\text{mean}_{\text{one clutch}} = 43.9$ ;  $\text{mean}_{\text{more than one clutch}} = 49.4$ ;  $U = 12$ ,  $N_1 = 14$ ,  $N_2 = 7$ ,  $P = 0.006$ ;  $P < 0.001$  without correction). Furthermore, we found that monopaternal clutches were fathered by males larger than other fathers (data of the 3 years pooled:  $\text{mean}_{\text{monopaternal fathers}} = 51.4$ ;  $\text{mean}_{\text{multipaternal fathers}} =$

46.1;  $U = 67$ ;  $N_1 = 17$ ;  $N_2 = 4$ ;  $P < 0.001$ ;  $P < 0.001$  without correction). In addition, the annual number of offspring sired by a male was correlated with paternal body length (SVL corrected; Spearman rank correlation:  $r_s = 0.252$ ,  $N = 21$ ,  $P = 0.020$ ; see Figure 1;  $P = 0.005$  without correction). Moreover, the number of fathers of a clutch had no impact either on the number of offspring ( $r_s = -0.174$ ,  $N = 13$ ,  $P = 0.549$ ;  $P = 0.882$  without correction) or on the neonate survival rate ( $r_s = 0.176$ ,  $N = 13$ ,  $P = 0.954$ ;  $P = 0.524$  without correction).

## Discussion

### Multiple Paternity

Our study demonstrates for the first time that multiple paternity occurs in natural populations of *V. berus*, even if the density of adults is low. The proportion of multiple paternity (69% of the clutches were fathered by more than one male) observed in our study is comparable to that reported by Höggren and Tegelström (2002) for captive populations. The high level of multiple paternity observed in these natural populations questions the copulatory plug hypothesis (Nilson and Andrén 1982). These authors described male renal sex-segment secretions causing a contraction of the female uterus after copulation and suggested that this phenomena could prevent fertilization by subsequent copulating males. This copulatory plug hypothesis has already been questioned by Stille et al. (1986) and Höggren and Tegelström (1995). Although male renal secretions do not seem to prevent further fertilization by other males, they could reduce the fecundity from further matings within a season as observed by Höggren and Tegelström (2002).



**Figure 1.** Correlation between the corrected SVL of the father and its number of offspring per year (Spearman rank correlation:  $r_s = 0.252$ ,  $N = 21$ ,  $P = 0.02$ ).

### Density and Level of Multiple Paternity

The relationship between density and levels of multiple paternity is a currently debated subject, particularly in birds. Indeed, several studies demonstrated a positive correlation between density and levels of multiple paternity (e.g., Bjornstad and Lifjeld 1997), whereas others demonstrated a lack of correlation (e.g., Veiga and Boto 2000). In a review, Kokko and Rankin (2006) suggested that density has an impact on female mating strategies, which can explain the observed discrepancy. However, these assumptions have been obtained for species that mainly live in couples. For species with limited interactions between sexes, only few studies have been published to our knowledge. For instance, in the ovoviparous fish *Heterandria formosa*, Soucy and Travis (2003) found a positive relationship between density and multiple paternity, which is probably related to a limited contact between males and females. No such correlations were observed in *Drosophila* species (Gromko and Gerhart 1984; Ochando et al. 1996). Comparing 2 populations of garter snake (*Thamnophis sirtalis*), Garner et al. (2002) suggested that the level of multiple paternity could be related to the density of males. The results stemming from our study do not confirm this hypothesis because the values observed in the adder are similar between our study (69% of multiple paternity; 100% in the first and 50% in the second population, although density is similar) and data from captive, high-density populations (84% and 75% in Höggren and Tegelström 1995 and Höggren and Tegelström 2002, respectively). Therefore, the lack of a consistent pattern in species with no parental care, such as most snakes, highlights the need for a meta-analysis in order to determine

general trends concerning the relationship between density and multiple paternity levels.

### Impact of Body Size (SVL) on the Number of Offspring

Combats between males occur for access to reproductive females, and the largest males win most of the time (Andrén 1986; Madsen 1988; Luiselli 1993b) fathering more offspring (this study). Sneaky males do, however, mate and fertilize some ova. Our data show that, even though small males successfully fathered offspring, the biggest males fathered the majority of offspring or were the single fathers in a clutch. Only in rare circumstances were smaller males the dominant fathers of a clutch. This may occur when they are the first males to mate with a female, before being dislodged by bigger males during combats. Because smaller males are unlikely to father a large number of offspring and because they cannot outcompete bigger males for mating opportunities, their best option to enhance their reproductive success may be through multiple copulations.

### Neonate Survival Rate and Multiple Paternity

Madsen et al. (1992) observed a positive relationship between the number of copulations and the neonate survival rate, suggesting that sperm competition in the female's reproductive tract occurred and that some males have a very low fecundity. We found no relationship between the number of fathers in a clutch and the survival of the neonates (confirming the observation of a lack of influence on survival rate between multiple and single mate copulations conducted by Capula and Luiselli (1994). The low neonate survival rate observed by Madsen et al. (1992) may be related to the low genetic diversity, and hence possibly low male fertility, in that particular population (Madsen et al. 1996; Madsen et al. 1999). In contrast, the Jura Mountain populations do not suffer from reduced genetic variability (Ursenbacher et al. forthcoming). Theoretically, when males have a low fertility, multiple copulations with different males have a particularly positive effect on offspring survival (see Loman et al. 1988) by avoiding deleterious effects of inbreeding and ensuring against male infertility. Consequently, multiple mating (with different males) should have a beneficial impact on the neonate survival rate particularly in populations characterized by a low genetic diversity and/or a low male fertility (see also Shine 2003; Blouin-Demers et al. 2005). Effects of multiple paternity should, however, not only be tested with respect to neonate survival but also in relation to offspring survival from birth to adult.

This study has demonstrated that in *V. berus*, the largest males father the majority of offspring. Moreover, multiple paternity appears to be common in low-density populations of this species. The influence of female sexual behavior has probably been underestimated in previous reptile studies. Further investigations focusing on female mate choice should thus be conducted to refine our knowledge of the reproductive strategies in *V. berus* and other snake species.

## Funding

Swiss National Foundation (3100-059132.99/1 and a Fellowship for prospective researchers); Société Académique Vaudoise.

## Acknowledgments

We thank V. Helfer, C. König, and S. Sachot for field assistance. N. Duvoisin and N. di Marco helped us in the laboratory. We thank P. Christe, J. Cosendai, T. Madsen, J.-C. Monney, J. Parker, C. E. Pook, P. Taberlet, J. Yearsley, and 3 anonymous reviewers for helpful comments on earlier versions of the manuscript.

## References

- Andrén C. 1986. Courtship, mating and agonistic behaviour in a free-living population of adders, *Vipera berus*. *Amphib Reptil.* 7:353–383.
- Andrén C, Nilson G. 1987. The copulatory plug of the adder, *Vipera berus*—does it keep sperm in or out. *Oikos.* 49:230–232.
- Birkhead TR, Møller AP. 1998. Sperm competition and sexual selection. London: Academic Press.
- Bjornstad G, Lifjeld JT. 1997. High frequency of extra-pair paternity in a dense and synchronous population of Willow Warblers *Phylloscopus trochilus*. *J Avian Biol.* 28:319–324.
- Blouin-Demers G, Gibbs HL, Weatherhead PJ. 2005. Genetic evidence for sexual selection in black ratsnakes, *Elaphe obsoleta*. *Anim Behav.* 69: 225–234.
- Bryant DM. 1988. Lifetime reproductive success of house martins. In: Clutton-Brock TH, editor. Reproductive success. Studies of individual variation in contrasting breeding systems. Chicago: University of Chicago Press. p. 173–188.
- Capula M, Luiselli L. 1994. Can female adders multiply. *Nature* 369:528.
- Garner TWJ, Gregory PT, McCracken GF, Burghardt GM, Koop BF, McLain SE, Nelson RJ. 2002. Geographic variation of multiple paternity in the common garter snake (*Thamnophis sirtalis*). *Copeia* 15–23.
- Grant PR, Grant BR. 2000. Non-random fitness variation in two population of Darwin's finches. *Proc R Soc Lond B Biol Sci.* 267: 131–138.
- Gromko MH, Gerhart PD. 1984. Increased density does not increase remating frequency in laboratory populations of *Drosophila melanogaster*. *Evolution* 38:451–455.
- Höggren M, Tegelström H. 1995. DNA-fingerprinting shows within-season multiple paternity in the adder (*Vipera berus*). *Copeia* 1995:271–277.
- Höggren M, Tegelström H. 1996. Does long-term storage of spermatozoa occur in the adder (*Vipera berus*)? *J Zool.* 240:501–510.
- Höggren M, Tegelström H. 2002. Genetic evidence for first-male mating advantage in the adder (*Vipera berus*). In: Schuett GW, Höggren MH, Douglas ME, Greene HW, editors. Biology of the Viper. Sandy (UT): Eagle Mountain Publishing.
- Jones AG, Ardren WR. 2003. Methods of parentage analysis in natural populations. *Mol Ecol.* 12:2511–2523.
- Kokko H, Rankin DJ. 2006. Lonely hearts or sex in the city? Density-dependent effects in mating systems. *Philos Trans R Soc B Biol Sci.* 361:319–334.
- Le Boeuf BJ, Reiter J. 1988. Lifetime reproductive success in northern elephant seals. In: Clutton-Brock TH, editor. Reproductive success. Studies of individual variation in contrasting breeding systems. Chicago: University of Chicago Press. p. 173–188.
- Loman J, Madsen T, Hakansson T. 1988. Increased fitness from multiple matings, and genetic heterogeneity: a model of a possible mechanism. *Oikos.* 52:69–72.
- Luiselli L. 1993a. High philopatry can produce strong sexual competition in male adders, *Vipera berus*. *Amphib Reptil.* 14:310–311.
- Luiselli L. 1993b. Are sperm storage and within season multiple mating important components of the adder reproductive biology. *Acta Oecol-Int J Ecol.* 14:705–710.
- Madsen T. 1988. Reproductive success, mortality and sexual size dimorphism in the adder, *Vipera berus*. *Holarctic Ecol.* 11:77–80.
- Madsen T, Shine R. 1992. A rapid, sexually selected shift in mean body size in a population of snakes. *Evolution* 46:1220–1224.
- Madsen T, Shine R. 1993. Temporal variability in sexual selection acting on reproductive tactics and body size in male snakes. *Am Nat.* 141:167–171.
- Madsen T, Shine R, Loman J, Hakansson T. 1992. Why do female adders copulate so frequently? *Nature* 355:440–441.
- Madsen T, Shine R, Olsson M, Wittzell H. 1999. Conservation biology—restoration of an inbred adder population. *Nature* 402:34–35.
- Madsen T, Stille B, Shine R. 1996. Inbreeding depression in an isolated population of adders *Vipera berus*. *Biol Conserv.* 75:113–118.
- Marshall TC, Slate J, Kruuk LEB, Pemberton JM. 1998. Statistical confidence for likelihood-based paternity inference in natural populations. *Mol Ecol.* 7:639–655.
- Neumeyer R. 1987. Density and seasonal movements of the adder (*Vipera berus* L.) on a subalpine environment. *Amphib Reptil.* 2:63–82.
- Nilson G, Andrén C. 1982. Function of renal sex secretion and male hierarchy in the adder, *Vipera berus*, during reproduction. *Horm Behav.* 16:404–413.
- Ochando MD, Reyes A, Ayala FJ. 1996. Multiple paternity in two natural populations (orchard and vineyard) of *Drosophila*. *Proc Natl Acad Sci USA.* 93:11769–11773.
- Orsetti DM, Rutowski RL. 2003. No material benefits, and a fertilization cost, for multiple mating by female leaf beetles. *Anim Behav.* 66:477–484.
- Otis DL, Burnham KP, White GC, Anderson DR. 1978. Statistical inference from capture data on closed animal population. *Wildl. Monogr.* 62:7–135.
- Prosser MR, Weatherhead PJ, Gibbs HL, Brown GP. 2002. Genetic analysis of the mating system and opportunity for sexual selection in northern water snakes (*Nerodia sipedon*). *Behav Ecol.* 13:800–807.
- Reynolds JD. 1996. Animal breeding systems. *Trends Ecol Evol.* 11:68–72.
- Shine R. 2003. Reproductive strategies in snakes. *Proc R Soc Lond B Biol Sci.* 270:995–1004.
- Simmons LW. 2005. The evolution of polyandry: sperm competition, sperm selection, and offspring viability. *Annu Rev Ecol Evol Syst.* 36:125–146.
- Soucy S, Travis J. 2003. Multiple paternity and population genetic structure in natural populations of the poeciliid fish, *Heterandria formosa*. *J Evol Biol.* 16:1328–1336.
- Stille B, Madsen T, Niklasson M. 1986. Multiple paternity in the adder, *Vipera berus*. *Oikos.* 47:173–175.
- Ursenbacher S, Monney JC. 2003. Résultats de 5 années de suivi d'une population de Vipère péliade (*Vipera berus*) dans le Jura Suisse: estimation des effectifs et discussion des méthodes d'estimation. *Bull Soc Herp Fr.* 107:15–25.
- Ursenbacher S, Monney J-C, Fumagalli L. Forthcoming. Limited genetic diversity and high differentiation among the remnant adder (*Vipera berus*) populations in the Swiss and French Jura Mountains. *Conserv Genet.* doi: 10.1007/s10592-008-9580-7.
- Veiga JP, Boto L. 2000. Low frequency of extra-pair fertilisations in House Sparrows breeding at high density. *J Avian Biol.* 31:237–244.

White GC, Burnham KP. 1999. Program MARK: survival estimation from population of marked animals. *Bird Stud.* 46:120–139.

Yasui Y. 1998. The 'genetic benefits' of female multiple mating reconsidered. *Trends Ecol Evol.* 13:246–250.

**Received April 18, 2008; Revised September 18, 2008;  
Accepted November 4, 2008**

**Corresponding Editor: William Modi**