First record of multiple paternity in the pygmy rabbit (Brachylagus idahoensis): evidence from analysis of 16 microsatellite loci

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Lagomorphs exhibit polygamy, which is the predominant mating system for mammalian species (Kleiman 1977). Furthermore, promiscuity, in which both sexes mate with multiple partners, is believed to be a common mating system among leporids (Crowan and Bell 1986), although some behavioral evidence suggests polygyny within the genus Lepus (Flux 1981, Rioja et al. 2008) and even monogamy in the whitesided jackrabbit (Lepus callotis; Best and Henry 1993). Traits consistent with promiscuous mating systems in leporids include limited parental care, overlapping home ranges, and multiple mating (Burton 2002). Although extensively described in birds (Griffith et al. 2002) and documented in all reptiles that have been investigated (Uller and Olsson 2008), multiple paternity has only been documented recently in mammals with promiscuous mating systems (Burton 2002, Krajiejeveld-Smit et al. 2002, Radespiel et al. 2002, Hohoff et al. 2003). Multiple paternity occurs when more than one male fathers a single offspring group (i.e., there is more than one sire for a litter or clutch; Sugg and Chesser 1994). In the genus Sylvilagus, which is considered to be a sister taxon of Brachylagus (Halanych and Robinson 1997, Robinson and Matthee 2005), testis size is suggestive of multiple mating in Sylvilagus floridanus (Stockley 2003); however, genetic evidence has not confirmed the occurrence of multiple paternity in any Sylvilagus species. On the other hand, genetic evidence for multiple paternity was detected in 25% of the litters tested in the promiscuous snowshoe hare (Lepus americanus; Burton 2002), and also has been documented in the brown hare (Lepus europaeus; Suchentrunk and Hacklaender 2005).

The pygmy rabbit (Brachylagus idahoensis) is the smallest rabbit in North America. It is a sagebrush-dependent species (Artemisia spp.) that uses areas dominated by big sagebrush (A. tridentata) and relatively deep soils (Orr 1940).
The historic range of the pygmy rabbit includes 8 western states: Idaho, Washington, Nevada, Montana, California, Utah, Wyoming, and Oregon (Green and Flinders 1980). In 2003, the U.S. Fish and Wildlife Service listed the distinct population of pygmy rabbits in the Columbia Basin as endangered, and a petition is currently under consideration for federal listing across the species’ range (Federal Register 2003, 2008).

The mating behavior and parental care of pygmy rabbits was studied in captivity as part of a captive-breeding program to restore the extirpated Columbia Basin population to their historic range in Washington (Elias et al. 2006). Mating behavior consisted of several chases and brief copulations. Unlike females of other lagomorph species in North America, female pygmy rabbits dug natal burrows, usually separated from their residential burrow systems, and covered the entrance of the burrows after the kits were born. In the wild, female pygmy rabbits dug natal burrows at considerable distances from their residential burrow systems (Rachlow et al. 2005).

Few field studies have reported observations of mating behavior of free-ranging pygmy rabbits (Janson 1940, Wilde 1978, Fisher 1979), and no studies have tested for multiple paternity. In captivity, male pygmy rabbits exhibited no parental care, and females provided relatively little care (nursing 1–2 times per day), like many other leporids (Elias et al. 2006, Bautista et al. 2008). In the wild, male pygmy rabbits tend to have larger home ranges and core areas, especially during the breeding season, and more dispersed burrow systems than females, possibly due to mate-search behavior (Burak 2006, Crawford 2008, Sanchez and Rachlow 2008). Switching among burrow systems is a common behavior for both sexes. Also, the overlap of home ranges between individuals of both sexes is common (Burak 2006, Sanchez and Rachlow 2008). Pygmy rabbits breed from February to July, producing up to 3 litters per year with a percentage of mating behavior of free-ranging pygmy rabbits.

To assess whether multiple paternity occurs in *B. idahoensis*, we used 16 polymorphic microsatellite markers and DNA extracted from ear tissue samples obtained from 2 litters collected at 2 sites in east central Idaho. The study sites (Cedar Gulch and Rocky Canyon) are located about 6 km apart in the Lemhi Valley along the Idaho–Montana border. Both sites are dominated by sagebrush vegetation and mounded microtopography. For a detailed description of the sites, see Estes-Zumpf and Rachlow (2009). This analysis represents the first investigation of paternity in free-ranging pygmy rabbits.

We evaluated paternity for 10 rabbits from 2 litters. One litter consisted of 4 near-term kits collected from the carcass of a female at the Cedar Gulch (CG) site. The second litter consisted of 6 kits sampled from a natal burrow found at the Rocky Canyon (RC) site. Because the mother of the second litter was unknown, we also analyzed DNA samples for parentage from potential mothers in the vicinity of the natal burrow. These adult females were trapped during concurrent studies on space use and movement patterns by pygmy rabbits (Sanchez and Rachlow 2008, Estes-Zumpf 2008). All adult females captured within 400 m of the natal burrow were evaluated as potential mothers. This distance was almost double the mean width of female home ranges during the breeding season (250 m; Sanchez and Rachlow 2008).

We extracted DNA using the Qiagen tissue kit, following the Qiagen tissue protocol (Qiagen Inc., Valencia, CA) at the Laboratory for Conservation and Ecological Genetics (University of Idaho, Moscow, ID). Tissue samples were frozen in ethanol prior to DNA extraction. See Estes-Zumpf et al. (2010) for a detailed protocol.

We used the polymerase chain reaction (PCR) to amplify 16 microsatellite loci: 9 microsatellites developed for pygmy rabbits (A2, A10, A121, A124, A133, D103, D118, D121, and D126; Estes-Zumpf et al. 2008) and 7 developed for the European rabbit (*Oryctolagus cuniculus*; Sol08, Sol30, Sat05, Sat07, Sat08, Sat12, and Sol44; Rico et al. 1994, Mougel et al. 1997, Surridge et al. 1997, Warheit 2001). We multiplexed the microsatellite markers following the PCR conditions established by Estes-Zumpf et al. (2008) and Estes-Zumpf (2008, appendixes G and H). The microsatellite genotypes were visualized with an Applied Biosystems 3130xl sequencer (Applied Biosystems, Foster City, CA), and alleles were scored using GeneMapper 3.7 (Applied Biosystems). We conducted a minimum of 2 PCRs per locus per sample and required that each allele was observed ≥2 times before finalizing a genotype. In a few cases, we encountered disagreements among PCRs due to allelic dropouts or false alleles, and genotypes were replicated 3–5 times until an accurate consensus
More than 3 alleles provided evidence of multiple paternity.

We assessed genotypes for 7 adult females that were trapped within 400 m of the natal burrow for the litter sampled at the RC site. None of the females matched the litter across all loci; however, one female was consistent at all loci except for D118. At that locus, the mismatch consisted of a 2 base-pair disagreement with 2 kits. The probability of exclusion for females using this set of loci was 0.999, indicating that the probability of this female not being the mother was <0.001. Thus, we concluded that this female was the mother. D118 is a dinucleotide locus for which we suspect a mutation by replication slippage, which is characterized by the addition or deletion of repeats in the microsatellite (reviewed in Ellegreen 2000).

Our results demonstrated multiple paternity for both litters tested. Evidence of multiple paternity was detected at loci A2, A10, Sat08, Sol44, Sol30, and Sat12 for the CG litter, and loci A133, Sat05, A10, Sol08, Sol44, and D121 for the RC litter (Table 1). For both litters, knowledge of the maternal genotype greatly increased our ability to detect multiple paternity, and in fact, we could not detect multiple paternity for the RC litter without this information (Table 1). For the loci with evidence of multiple paternity, we found 3 and 4 paternal alleles in the CG and RC litters, respectively, which indicates a minimum of 2 sires per litter.

This study is the first to evaluate patterns of paternity for pygmy rabbits and the first to document multiple paternity in this species. Our findings support the expectation that pygmy rabbits, like other lagomorphs, have a promiscuous mating system. Promiscuity has been linked with multiple paternity in other species, including the meadow vole (Microtus pennsylvanicus; Bertaux et al. 1999), agile antechinus (Antechinus agilis; Kraaijeveld-Smit et al. 2002), yellow-toothed cavy (Galea musteloides; Holhoff et al. 2003), gray mouse lemur (Microcebus murinus; Badeispiel et al. 2002), and the snowshoe hare (Lepus americanus; Burton 2002). Our results are also consistent with traits associated with multiple paternity in free-ranging populations. Due to our small sample size, we could not evaluate the prevalence of multiple paternity in our study population. However, we documented multiple paternity in both litters that we tested, which indicates that this reproductive pattern might be relatively common within this species.

### Table 1. Microsatellite analysis of multiple paternity in 2 litters of pygmy rabbits collected in east central Idaho. This table shows the total number of alleles and the number of paternal alleles per litter per locus. Bolded, italicized numbers indicate evidence of multiple paternity. CG = Cedar Gulch litter, and RC = Rocky Canyon litter.

<table>
<thead>
<tr>
<th>Locus</th>
<th>Alleles per litter (CG)</th>
<th>Paternal alleles (CG)</th>
<th>Alleles per litter (RC)</th>
<th>Paternal alleles (RC)</th>
</tr>
</thead>
<tbody>
<tr>
<td>A2</td>
<td>5</td>
<td>3</td>
<td>4</td>
<td>2</td>
</tr>
<tr>
<td>A133</td>
<td>3</td>
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<td>4</td>
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<td>A121</td>
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<td>D103</td>
<td>1</td>
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<td>3</td>
<td>2</td>
</tr>
<tr>
<td>Sat05</td>
<td>2</td>
<td>2</td>
<td>4</td>
<td>4</td>
</tr>
<tr>
<td>A10</td>
<td>4</td>
<td>3</td>
<td>3</td>
<td>3</td>
</tr>
<tr>
<td>Sol08</td>
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<td>Sat08</td>
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<td>D126</td>
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<td>Sol44</td>
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<td>D118</td>
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<td>2</td>
</tr>
<tr>
<td>Sat12</td>
<td>4</td>
<td>3</td>
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</tr>
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<td>Sat07</td>
<td>1</td>
<td>1</td>
<td>3</td>
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</tr>
<tr>
<td>D121</td>
<td>2</td>
<td>2</td>
<td>4</td>
<td>3</td>
</tr>
<tr>
<td>A124</td>
<td>4</td>
<td>2</td>
<td>3</td>
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</tr>
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</table>
Multiple paternity has several ecological and evolutionary implications. Populations that exhibit multiple paternity tend to have larger effective population sizes (Sugg and Chesser 1994), a reduction of inbreeding (Stockley et al. 1993), and a reduction in the costs of reproductive failure resulting from genetic incompatibility (Stockley 2003). Given that pygmy rabbits currently persist in small and often fragmented populations (Rachlow and Svancara 2006, Larrucea and Brussard 2008) and might have done so historically (Grayson 1987), these traits might be important for maintaining genetic diversity within free-ranging populations. Such traits could also assist in meeting the goals of the captive breeding program for the endangered Columbia Basin population.

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**FEDERAL REGISTER.** 2003 [March 5]. Endangered and threatened wildlife and plants; final rule to list the Columbia Basin Distinct population segment of the pygmy rabbit (*Brachylagus idahoensis*) as endangered. Federal Register 68:10388–10409.

**FEDERAL REGISTER.** 2008 [January 8]. Endangered and threatened wildlife and plants; 90-day finding on a petition to list the pygmy rabbit (*Brachylagus idahoensis*) as threatened or endangered. Federal Register 73:13112–13113.


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