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1 Home range and dispersal of the field vole Microtus agrestis in an organic agro-ecosystem 2 3 Thomas Secher Jensen, Kent Olsen, Christina Vedel-Smith & Tine Sussi Hansen 4 Natural History Museum Aarhus, Wilhelm Meyers Allé 210, DK-8000 Aarhus C, Denmark 5 6 Corresponding author: Tel.: +45 86129777; fax +45 86130882. 7 E-mail address: tsj@nathist.dk (T. S. Jensen). 8 9 Word count: 4.285 10 11 12 **Abstract** 13 During 2008-2011 field voles Microtus agrestis were studied in a Danish agro-ecosystem in connection with 14 a programme elucidating the role of organic farming as refuge for biodiversity, due to the observed losses 15 of biodiversity in conventional farmland. The aim of the study was to quantify home ranges and dispersal of 16 voles in order to test a hypothesis that vole populations living in organic farms could colonize unoccupied 17 areas in conventional farms. Voles were radio-collared and tracked until death or disappearance. Results 18 showed that vole home ranges were larger during the onset of the breeding season than during the height 19 of the breeding season and the non-breeding period. Males had larger home ranges than females. Voles 20 occupying isolated small biotopes had restricted home range and only few individuals crossed fields from 21 one small biotope to another. Although crossings do occur, the magnitude of dispersal seems restricted in 22 fragmented agro-ecosystems.

Keywords: Microtus agrestis, home range, dispersal, telemetry, organic farming

## Introduction

In North European lowlands such as Denmark, southern Sweden, northern Germany, northern Poland, the Netherlands, Belgium and northern France, most of the non-urban areas are used for agriculture, leaving few and often highly fragmented small biotopes in between. Conventional farming is often intensive with a minimal crop rotation, an abundant use of pesticides and artificial fertilisers, and large mono-cultural fields surrounding small fragmented biotope patches. In contrast, organic farming avoids pesticides and artificial fertilisers, and uses more mechanical weed control and organic manure. Organic farms are normally less intensively managed, and the number and the size of small biotopes are higher.

Generally, biodiversity in agro-ecosystems is associated with the surrounding small biotopes rather than the fields in rotation and the crops. On a farm level, small mammal biodiversity is low, although on a larger geographical scale several species can be found (Jensen and Hansen, 2003). Field vole *Microtus agrestis*, is one of the most common mammalian species in open grassy habitats and comprises one of the most important prey items for a number of specialist predators, e.g. weasel *Mustela nivalis* and stoat *Mustela erminea*, as well as generalist predators, e.g. fox *Vulpes vulpes*, badger *Meles meles*, buzzard *Buteo buteo*, kestrel *Falco tinnunculus* and a number of owl species. The voles are most often found in grassy habitats like banks, hedge rows, buffer zones, pond surroundings, moist meadows and set-aside fields (Jensen and Hansen, 2001). These agro-ecosystem components are often scattered and fragmented and can thus be considered as small habitat islands, where populations of voles are subjected to extinction and may rely on re-colonisation.

The population dynamics and dispersal of *M. agrestis* has been well described at northern latitudes in connection with the cyclicity of the species (Hansson, 1971; Myllymäki, 1977; Stenseth et al., 1977; Lambin et al., 2000) and more to the south in contrasting non-cyclic populations (Erlinge et al., 1990; Sandell et al., 1991). Also damage by the species to forest saplings has led to several studies (e.g. Larsson, 1975). In contrast, studies of the species in agro-ecosystems are few (Jensen and Hansen, 2003; Huitu et al., 2008; Yletyinen and Norrdahl, 2008).

In this study we employed radio telemetry to obtain information on home range and short time dispersal of *M. agrestis* within and between small agro-ecosystem biotopes. The home range data are necessary prerequisites for further development of agent-based models, i.e. ALMaSS (Topping et al., 2001) and genetic models (Marchi et al., submitted).

## **Materials and methods**

Field work was conducted in 2008-2011 in Eastern Jutland, Denmark, with seven sites selected within the property of the Kalø Estate (56 17 N, 10 29 E). This estate is managed by means of organic farming principles, i.e. the crops are not treated with pesticides, and artificial fertilisers are not used. The estate is surrounded by a number of smaller private farms managed in conventional practice. Originally the project intended to study differences between conventional and organic farming systems; however, due to a very low number of voles in the conventional farms, all telemetry studies were performed within organic farms.

Crop rotations within the estate are dominated by spring barley, spring wheat, triticale, and a leguminous crop and intercropping (or mixed cropping) with undersowing of clover and ryegrass.

Generally the size of the fields is large and the soil is mostly clay. As organic farming practices are employed, several wildlife friendly small biotopes are created, e.g. set-aside strips along field margins, beetle banks within larger fields, and smaller set-aside fields. The matrix surrounding the cultivated fields consists of small habitat patches of different biotopes, e.g. hedgerows, grassy banks, ponds and their immediate surroundings, and wildlife plantings. Two larger forests, Hestehave Forest and Ringelmose Forest, are located at the western and eastern border of the estate, respectively.

The seven study sites (table 1) included: 1) a 17-year-old set-aside field surrounded by mixed forest on one side, a hedge row on the opposite side and fields in rotation along the two remaining sides; 2) a 17-year-old grassy bank along an old hedgerow and running between fields in rotation and connected to site 1 in its southern end; 3) a 16-year-old grassy bank surrounding a rainwater reservoir, 4) a cattle grazed field surrounded by mixed forest, hedge rows and a field in rotation; 5) a 5-year-old wildlife friendly set-

aside stripe with beetle bank running parallel with a grassy bank within a field in rotation; 6) a 6-year-old rectangular shaped set-aside field encircling a dry depression and surrounded by fields in rotation; and 7) a 15-year-old set-aside field surrounded by hedge row and coniferous forests.

#### **INSERT TABLE 1**

Live trapping for field voles was performed with Ugglan lemming and Ugglan special traps,

Grahnab AB, Sweden. Voles were tagged with a Passive Integrated Transponder tag (BSA International AG,

Sweden) and applied with a 0.9 g cable-tie radio tag from Biotrack (TW-4 and PIP3 transmitter types).

Radio-collared voles were tracked by means of a Stabo radiotracking receiver carrying a Flexi Yagi antenna.

Consecutive fixes were spaced by at least 45-minute intervals to avoid auto-correlations (de Solla et al.,

1999).

The positions of the tracked voles were marked in the field and subsequently GPS positioned and transferred to MapInfo (Insight, 2011) to visualise activity area in relation to updated Ortophotos. All individuals were either adults or sub-adults weighing more than 20 g, i.e. transmitter weight was less than 5% of the vole weight.

The tracking of voles continued until the signal disappeared, or until the position remained fixed for several tracking sessions. In the latter case the transmitter was subsequently located and, if possible, the final status of the vole determined. If a vole was predated or it lost its collar before sufficient fixes were obtained, the tag was applied to a new animal and tracking sessions commenced. Three days before the tags were timed out due to the internal battery capacity the voles were recaptured and freed from their collar.

Home range data was analysed in Ranges 8 (Kenward et al., 2008) and included 99-5% convex polygon cores, incremental plots, and kernel contour core analysis. All statistical test were performed using the SAS Enterprise Guide 4.3 (SAS, 2006) and statistical significance was recognized at P <

0.05 and prior to pooling any predictor variables for more than one season or site, differences between years and sites were tested, even though all such results are not presented in the paper. Differences among sample means were examined using one way analyses of variance (ANOVA) when comparing more than one sample, and reported as the distance between individual distributions (F) with degrees of freedom and sample size in subscript and probability (*P*).

#### Results

In all periods loss of transmitters was high, and new individuals had to be caught in order to maintain a recordable population. Losses could mainly be ascribed to probable predation (33%), when remains of the animal was found attached to a collar. In further 31% the signal was lost as we were unable to locate the animal, probably because the collar was moved too far away by a predator, and finally 17% of the collars were found again without trace of the animal, partly because of a too loose collar or partly due to predation (table 2). Main predators suspected were weasel *Mustela nivalis*, when transmitters were found in the runways with parts of the animal remaining, and common buzzard *Buteo buteo* and kestrel *Falco tinnunculus*, when transmitters were found in or below trees. Total disappearance of transmitter signal was ascribed to long distance removal of the whole carcass by e.g. birds of prey or foxes *Vulpes vulpes*.

## **INSERT TABLE 2**

## **INSERT FIGURE 1**

In 2008 the studies were performed during late autumn, e.g. at a time of year when breeding in general had stopped and the tagged individuals were either post-reproductive adults or sub-adults who had not bred during the present breeding season. Only one set-aside field was used in contrast to the consecutive seasons and except for one individual all tagged animals were males. Mean 95% polygon core

home range estimate was 262 m<sup>2</sup>, range 11-900 m<sup>2</sup> (table 3). There was no significant difference in home range estimates between 95% polygon core, the 95% kernel contour and the minimum convex polygon analysis methods ( $F_{2,30} = 0.43$ ; P = 0.6535; fig. 1).

In 2009 studies were performed during autumn at six different sites (table 1). The results showed a mean 95% polygon core home range estimate for males of 421 m<sup>2</sup>, range 10-1290 m<sup>2</sup>, and for females of 176 m<sup>2</sup>, range 72-443 m<sup>2</sup> (table 3). Again, there was no significant difference in home range estimates between the three different analysis methods ( $F_{2,54} = 0.80$ ; P = 0.4565; fig. 1).

In July 2010, at the height of the breeding season, the mean 95% polygon core home range estimate tended to be larger than in autumn 2008 and 2009 (fig. 1). The mean 95% polygon core home range estimate was found to be 511 m<sup>2</sup> for males and 443 m<sup>2</sup> for females though considerable variation was present with a male range of 12-1312 m<sup>2</sup> and female range of 7-1704 m<sup>2</sup> (table 3). The mean 95% kernel contour home range estimate was somewhat larger than 95% polygon core and minimum convex polygon ranges, however, not significantly so  $(F_{2,66} = 1.85; P = 0.1657;$  fig. 1).

In April-June 2011, at the onset of the breeding season and when field crops started growing, home range estimates generally increased, especially so for sexually active vole males. The overall mean 95% polygon core home range estimate was 603 m<sup>2</sup> with a range of 12-2482 m<sup>2</sup>, whereas mean male estimates was 746 m<sup>2</sup> with a range of 12-2482 m<sup>2</sup> and for females 423 m<sup>2</sup> with a range of 37-666 m<sup>2</sup> (table 3). There was no significant difference between the three calculation methods ( $F_{2,54} = 1.76$ ; P = 0.1828; fig. 1).

## **INSERT TABLE 3**

# **INSERT FIGURE 2**

Generally, voles did not move much between consecutive telemetry tracking as indicated by the overall mean inter-location line-of-sight dispersal distance being only 6 meters (fig. 2). However, maximum distance was 122 meters (SD = 27) with a mean of 31 meters (table 4). The largest inter-location distances found belonged to a few reproductive males at the onset of the breeding season; however, also a few reproductive females had large inter-location distances. The mean inter-location dispersal distance did not differ between voles in large versus small habitats (female:  $F_{1,32} = 0.02$ ; P = 0.8960; male:  $F_{1,36} = 0.04$ ; P = 0.8492 and both sexes:  $F_{1,68} = 0.67$ ; P = 0.4158).

#### **INSERT TABLE 4**

## **INSERT FIGURE 3**

The overall relation between the inter-location dispersal distance and the size of the 100% MCP area estimate was found to be positive and highly significant ( $F_{1,68} = 38.74$ ; P < 0.0001; fig. 3a). However, when data was separated between sex and habitat size only three of the four combinations were significant (females in large habitats:  $F_{1,22} = 34.28$ ; P < 0.0001; females in small habitats:  $F_{1,10} = 3.36$ ; P = 0.1040; males in large habitats:  $F_{1,13} = 10.16$ ; P = 0.0087; and males in small habitats:  $F_{1,23} = 18.81$ ; P = 0.0003). Alternatively, there was no overall positive relation between mean inter-location measure and the number of fixes obtained ( $F_{1,68} = 1.46$ ; P = 0.2309; fig. 3b).

The incremental plots in figure 4 illustrate the relationship between number of telemetry fixes and the estimated home range area for each individual and show at which point the final home range area estimate only increases little in size when adding new fixes. The 100% minimum convex polygon area estimate was generally reached within a relative short time, e.g. 20-30 fixes, indicating stable home ranges. However, even though there was a gradual increase in 100% MCP over the entire tracking period for some voles several individuals showed sudden abrupt increases indicating either a shift in core area or an

increase in home range size. In small habitats (fig. 4a) there was a higher tendency for voles to make sudden increase in home range compared to the large habitat where home ranges increased more gradually (fig. 4b) without sudden jumps. This was most commonly seen in the April-June sessions (fig. 4f).

## **INSERT FIGURE 4**

In the linear habitats male voles stayed within the habitat and did not disperse beyond habitat boundaries into the surrounding fields and, accordingly, they had very small home ranges with a mean 95% polygon core estimate of 296 m², range 12-845 m². In linear habitats along hedge rows home ranges were restricted to the habitat strip itself and in the wildlife friendly strip habitat (site 5) in particular the 95% polygon core home range estimates were severely restricted in size with a range of only 12-288 m². The individual voles all stayed within the linear habitat completely without crossing the boundary into the surrounding cultivated field. Voles in non-linear habitats such as all the large habitat sites and a few small habitat sites stayed also mainly within the set-aside habitat, however, their home ranges were much larger on average with a mean 95% polygon core estimate of 484 m², range 10-2482 m². The female vole home range estimates were smaller than that of males, and were higher in spring than in autumn, with a mean 95% polygon core of 436 m², range 64-1704 m².

#### Discussion

Predation rates were high overall. If we consider all disappeared voles as predated, the overall rate was around 81% in total (2009: 89%, 2010: 77%, and 2011: 78%). This figure is high, however, when voles in agro-ecosystems are concentrated in few and rather small biotope units, predators can concentrate in the same areas and hence impact must be expected to be high. However, Korpimäki et al. (1996) raised the question whether predation rates in telemetry studies are overestimated due to e.g. an inhibitory effect on vole dispersal and a decreased ability to escape the predators when carrying a radio collar. In contrast,

Webster and Brooks (1980) found no effect of radio-transmitters on the survival of the meadow vole Microtus pennsylvanicus.

Home range estimates of *M. agrestis* have mainly been obtained by means of live-trapping, thus restricting the potential home range area to that determined by the trap design. Furthermore, in most calculations the minimum convex polygon method has been applied, whereby the outermost points of trappings have been connected and the area inside calculated, generally leading to an inclusion of areas where the voles have not been recorded (Nilsen et al., 2008). We consider that the minimum convex polygon method generally overestimates the actual home range, as it includes large areas with no occurrence of voles. In contrast, especially the 50% kernel cores only include areas strictly around the main number of actual records, excluding areas with known records, thus leading to obvious underestimates of home ranges.

More recently, telemetry is employed, giving more precise data on vole locations. Loughran (2006, 2007) found by the use of telemetry no significant variation in male home range size during the breeding season or between years. However, there was significant variation in core area between years but not during the breeding season. In contrast, the present results indicate larger mean home ranges during the breeding season. This could be due to differences in population density, as in low density populations, individuals may disperse larger distances and occupy larger territories without competition (Sandell et al., 1991).

Christensen (1999) found by using telemetry on voles in a small set-aside field that mean home ranges and core areas of female voles were considerably larger in April-May than in other seasons, consistent with our findings. This applied both for minimum polygon core calculations and 95% cluster analysis. The incremental plot analysis used in this study highlights that individual vole home range size is a highly flexible entity, which is most difficult to assess. The large spring home ranges are most likely due to an increased sexual and feeding activity at the onset of the breeding season. In contrast to feeding and

mate availability, presence of predators is an important factor which might suppress the size of the home range (Carlsen et al., 2000) by altering microhabitat preferences.

We found that voles living in narrow linear habitats generally stayed within the habitat and only rarely dispersed moved beyond habitat boundaries into the surrounding cultivated fields, and especially voles in the wildlife friendly strip habitat (site 5) all stayed within the linear habitat completely without crossing the habitat boundary. Presumably, they were more or less stranded on a habitat island in a sea of hostile environment, i.e. between ploughed soils. Accordingly, voles in linear habitats had very small home ranges in comparison with vole in non-linear habitats.

In contrast, several voles from the non-linear study sites occasionally entered the surrounding cultivated field matrix; however, they always returned to the core area in their starting habitat. In one case, however, a female vole made a temporally change in home range and dispersed from a set-aside field across a cultivated field and into a grassy glade within an old forest; though shortly after she moved back into the old core area in the set-aside field. Most other voles changed core area within the same overall habitat as they were caught and when they occasionally reached the field boundaries they moved back into the field without crossing into the other habitat type. Thus we found little evidence of voles moving from grassy areas across crop fields to other small, grassy biotopes. As field voles are positively recorded in isolated habitat islands like small biotopes in fragmented open farmland such crossings do occur (Jensen and Hansen, 2001, 2003). Christensen (1999) found that out of 19 small biotopes investigated, voles were found in 70% during autumn and 40% during spring. She also found that frequent extinction and re-colonization of voles in small biotopes was a general phenomenon.

The studies on *Microtus agrestis* clearly reveal that vole individuals follow linear biotopes and stay within small biotopes when dispersing and normally avoid open fields, and that the area and quality of these biotopes set the limit for vole population numbers. Organic farms might contain higher densities of vole due to a higher density of small biotopes, which also entails the basis for an increase in predator numbers. Given that a reasonably fine meshed network of small biotopes is found, numbers of

250 voles and their predators can be increased by increasing these habitats, thus contributing to a higher 251 biodiversity in the agro-ecosystems. 252 253 **Acknowledgements** 254 The project was part of REFUGIA - The role of organic farms as refuge for biodiversity - under the 255 programme DARCOF III 2005-2010, Research in Organic Food and Farming, and funded by the Ministry of 256 Food, Agriculture and Fisheries under the finance and appropriation Act, Sections 24.33.02.10. 257 258 References 259 Carlsen, M., Lodal, J., Leirs, H., and Jensen, T.S. 2000. Effects of predation on temporary autumn 260 populations of subadult Clethrionomys glareolus in forest clearings. Zeitschrift für Säugetierkunde 65: 100-261 109. 262 263 Christensen, S.B. 1999. Småpattedyr i agerlandets småbiotoper. Master thesis. Aarhus Universitet, Aarhus. 264 265 de Solla, S.R., Bonduriansky, R., and Brooks, R.J. 1999. Eliminating autocorrelation reduces biological 266 relevance of home range estimates. Journal of Animal Ecology 68: 221-234. 267 268 Erlinge, S., Hoogenboom, J., Nelson, J., and Sandell, M. 1990. Density-related home-range size and overlap 269 in adult field voles (Microtus agrestis) in southern Sweden. Journal of Mammalogy 71: 597-603. 270 271 Hansson, L. 1971. Small rodent food, feeding and population dynamics. A comparison between granivorous 272 and herbivorous species in Scandinavia. OIKOS 22: 183-198. 273 274 Huitu, O., Laaksonen, J., Klemola, T., and Korpimäki, E. 2008. Spatial dynamics of Microtus vole populations 275 in contuinuous and fragmented agricultural landscapes. Oecologia 155: 53-61. 276 277 Insight, P.B.B. 2011. MapInfo Professional 11.0. Troy, New York, USA: Pitney Bowes Software Inc. 278 279 Jensen, T.S., and Hansen, T.S. 2001. Effekten af husdyrsgræsning på småpattedyr. In: Pedersen L.B., 280 Buttenschøn R.M., and Jensen T.S. (Eds.). Græsning på ekstensivt drevne naturarealer - effekter på 281 stofkredsløb og naturindhold. Park- og Landskabsserien nr. 34, Skov & Landskab, Hørsholm, pp. 107-121. 282 283 Jensen, T.S., and Hansen, T.S. 2003. Biodiversitet og biotopfordeling hos småpattedyr i det åbne land. Flora 284 og fauna 109: 8-21.

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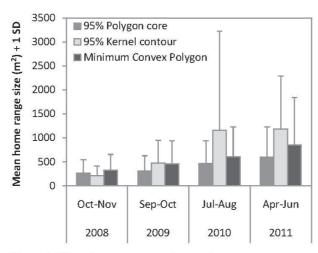


Figure 1: Mean home range estimates in square meters (m²) for all animals at all seven sites according to the three different analysis methods: 95% Polygon Core, 95% Kernel Contour and Minimum Convex Polygon (MCP). Number of sites: 2008 (n=1), 2009 (n=6), 2010 (n=4) and 2011 (n=5).

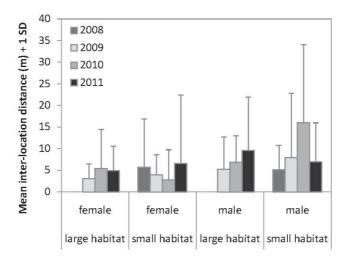


Figure 2: Mean line-of-sight dispersal distance between telemetry fixes divided into sex and habitat groups for each year.

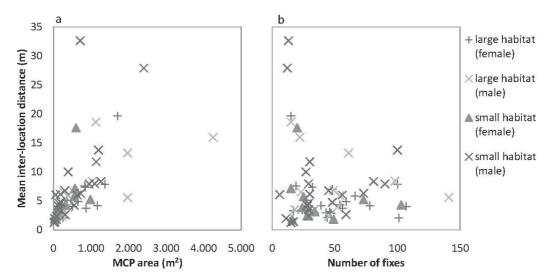


Figure 3: Mean inter-location line-of-sight dispersal distance as a function of minimum convex polygon (MCP) estimates for home ranges (a) and as a function of number of telemetry fixes (b). Illustrates differences between sex and habitat size.

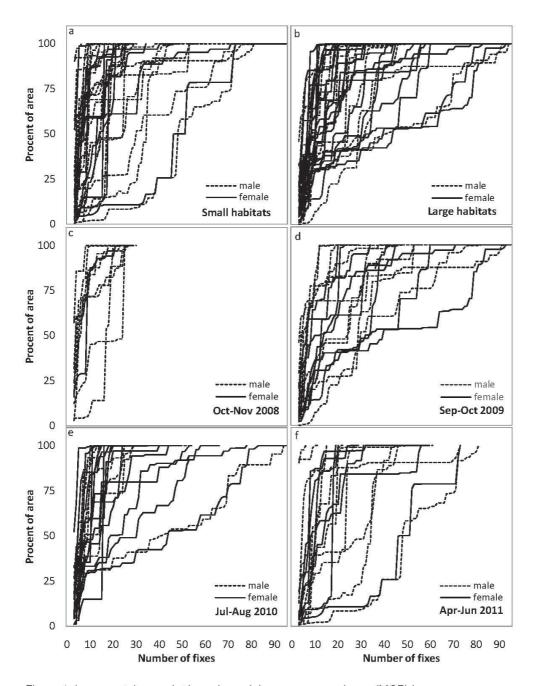


Figure 4. Incremental area plot based on minimum convex polygon (MCP) home range estimates with 100% of telemetry fixes. Illustrates differences between sex (all figures), habitat size (small (a) and large (b) habitats), and season (October-November (c), September-October (d), July-August (e) and April-June (f)).

Table 1: Characteristics of the sites used in the telemetry studies.								
Site	Name	Habitat type	Habitat size	Area (m <sup>2</sup> )	Perimeter (m)	Year of fieldwork		
1	Stegeloekke Field	Set-aside field	Large habitat	20,982	581	2009, 2010, 2011		
2	Stegeloekke Stripe	Set-aside stripe	Small habitat	1,838	316	2009, 2010		
3	Flintbjerg Parking	Habitat patch	Small habitat	3,389	547	2009, 2011		
4	Flintbjerg Meadow	Grazed meadow	Large habitat	38,352	1,040	2011		
5	Flintbjerg Stripe	Set-aside stripe	Small habitat	2,102	482	2009, 2011		
6	Vikaer West Field	Set-aside field	Small habitat	3,208	249	2008, 2009, 2010, 2011		
7	Vikaer East Field	Set-aside field	Large habitat	60,807	1,114	2009, 2010		

Table 2: Final status of tracked *Microtus agrestis. Alive*: Animals tracked until their radio transmitter was removed; *Predated*: Transmitter found together with the remnants of a predated animal or located underneath a tree; *Signal lost*: Signal lost, either due to defect transmitter or mostly likely due to predation; *Disappeared*: Transmitter relocated without the animal.

Year	Alive	Predated	Signal lost	Disappeared	
2009	11 %	17 %	39 %	33 %	
2010	23 %	50 %	18 %	9 %	
2011	22 %	28 %	39 %	11 %	
Total	19 %	33 %	31 %	17 %	

Table 3: Female and male *Microtus agrestis* home range estimates in square meters (m²) according to the three different analysis methods: 95% Polygon Core, 95% Kernel Contour and Minimum Convex Polygon (MCP).

		2008		2009		2010		2011	
			Oct-Nov	Sep-Oct Jul-Aug		Apr-Jun			
Sex	Analysis	n	Mean ± SD	n	Mean ± SD	n	Mean ± SD	n	Mean ± SD
	95% Polygon Core		$262 \pm 282$		$312 \pm 319$		$464 \pm 476$		$603 \pm 631$
All	95% Kernel Contour	10	$212 \pm 207$	18	475 ± 477	22	$1,157 \pm 2,067$	18	$1,185 \pm 1,104$
	MCP		$327 \pm 330$		$463 \pm 478$		$608 \pm 625$		$854 \pm 988$
	95% Polygon Core		483		176 ± 116		$443 \pm 506$		$423 \pm 268$
Female	95% Kernel Contour	1	216	8	209 ± 122	15	$995 \pm 2,046$	8	890 ± 880
	MCP		549		$233 \pm 132$		$500 \pm 518$		$638 \pm 358$
	95% Polygon Core		$237 \pm 287$		$421 \pm 390$		$511 \pm 439$		$746 \pm 802$
Male	95% Kernel Contour	9	$212 \pm 220$	10	$688 \pm 552$	7	$1,505 \pm 2,230$	10	1,421 ± 1,249
	MCP		$303 \pm 340$		$647 \pm 577$		$839 \pm 806$		$1,027 \pm 1,293$

Table 4: Maximum inter-location line-of-sight dispersal distance in meters (m) divided into sex and habitat groups for each year.

	Sm	all habitats	Lar	ge habitats	All habitats		
	n	Distance	n	Distance	n	Distance	
All	33	114	35	122	68	122	
Female	10	114	22	122	32	122	
2008	1	57	-	-	1	57 36	
2009	3	36	5	16	8		
2010	3	52	12	122	15	122	
2011	3	114	5	28	8	114	
Male	23	109	13	81	36	109	
2008	9	51	-	-	9	51	
2009	5	109	5	81	10 109		
2010	4	79	3	35	7	79	
2011	011 5 38 5 80		10	80			