Dietary response of the Eurasian badger, Meles meles, to a decline of its main prey in the Doñana National Park

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INTRODUCTION

The Eurasian badger (Meles meles) is considered to be a harvester rather than a hunter (Lruuk, 1989), because its food usually consists of vegetables and small and easily captured animals, such as worms and insects, and it rarely captures mammals (Neal, 1986). However, in the Doñana National Park of Mediterranean Spain the badger apparently behaves as a facultative specialist in the consumption of young rabbits, Oryctolagus cuniculus (Martín, Rodríguez & Delibes, 1995). In 1990, an outbreak of Rabbit Haemorrhagic Disease (RHD) caused high mortality among adult rabbits in most of the Doñana area (Villafuerte et al., 1994), although all sites of Doñana were not affected to the same extent. This situation offered the possibility of comparing the food habits of badgers in the Doñana Biological Reserve (DBR) before (data of Martín et al., 1995) and after the outbreak of RHD, while using another site where rabbit density has changed little over the last few years as a control area.

The study is related to the discussion about whether or not Eurasian badgers forage as food specialists (Lruuk & Parish, 1981; Roper, 1994). The terms ‘specialist’ and ‘generalist’ can be misleading, as they are relative terms and affect different decisions (e.g. before or after to encounter a potential prey). We suggest that the Eurasian badger could be considered a specialized forager where and when it focuses its foraging on one kind of food, which would be preferred over the rest. The theory of feeding specialization requires the selected prey to be profitable and abundant and predicts an increase of dietary diversity when the selected prey becomes scarce (Futuyma & Moreno, 1988). Hence, we test the hypothesis that the dietary diversity of badgers in Doñana is dependent on the relative biomass of rabbits in Doñana, regardless of the remaining consumed foods.

STUDY AREA

The study was carried out in the Doñana National Park (37°10’N, 6°23’W). The Park is situated on the west bank of the mouth of the Guadalquivir River (southwestern Spain). The climate in Doñana is Mediterranean
METHOD

We analysed 145 faeces collected in the DBR in the 2 years following the first known outbreak of RHD (i.e. the period 1990–1992) and 209 faeces collected in MG during 1992–1994. All samples were well distributed over the year. For purposes of comparison, we used the 265 latrine contents collected in DBR before RHD and reported by Martín et al. (1995). Because the availability of young rabbits is higher from December to May (Delibes & Calderón, 1979; Rogers, Arthur & Soriguer, 1994), we considered separately the diets of the so-called wet (Dec.–May) and a dry (June–Nov.) periods at each area.

Contents of faeces were determined using methods described by Reynolds & Aebischer (1991). We excluded infrequent prey (Roper, 1994) that represented less than 0.5% of the ingested biomass. Importance in the diet of each prey category was quantified by 2 methods: 1) frequency of occurrence = number of occurrences of each prey type × 100/number of faecal samples; 2) estimate of ingested fresh biomass by using correction factors (Lockie, 1959); the estimated ingested biomass = fraction of each prey type (estimated visually) × dry weight of the faecal sample × correction factor × 100/Z (dry weight of remains × correction factors). We used the same correction factors as Martín et al. (1995).

Differences in occurrence of particular prey categories among pairs of compared diets were evaluated by chi-square analysis of contingency tables with the Bonferroni confidence intervals (Rice, 1989). Diet diversity was calculated using the Levins Index and diet similarities were calculated using the Renkonen’s Index (Lrebs, 1989). Spearman rank correlation coefficients were used to relate trophic diversity and the importance of different prey categories.

RESULT

Young rabbits were the main prey item consumed by badgers in Matasgordas, during both the wet and the dry period (Table 1). Although frequencies of occurrence of rabbits differed between Matasgordas and the Doñana Biological Reserve before RHD (at least for the wet period; Table 1), the ingested biomass of rabbits was consistently high in both yearly periods and both areas (always above 55%; Table 1). According to the Renkonen indices, dietary similarities were high between MG and DBR before RHD (81.3% and 75.6% in wet and dry season, respectively), low between DBR before and DBR after RHD (60.2% and 46.2%) and intermediate between MG and DBR after RHD (64.0% and 62.3%).

Regarding other food categories, some statistical differences appeared when comparing frequencies of occurrence, probably related to local differences in availability. For example, fruit consumption was higher in MG with regard to DBR before RHD during both seasons (wet: $z^2 = 65.3$, d.f. = 1, $P<0.001$; dry: $z^2 = 24.0$, d.f. = 1, $P<0.001$) and with regard to DBR after RHD during the wet season ($z^2 = 34.4$, d.f. = 1; $P<0.001$).

sub-humid, with an average annual rainfall of 500–600 mm falling mainly from November to April. Two areas were selected within the Park:

1. Doñana Biological Reserve (DBR): described by Martín et al. (1995) and located in the centre of the Park (Fig. 1). Three main biotopes were distinguished within this area: marsh, dunes and scrubland. The scrubland was dominated by Halimium halimifolium shrubs with scattered Quevcur ruvest trees, while tziuyr spp. were dominant in the marsh. As in the study by Martin et al. (1995), we collected faeces in DBR in the ecotone scrubland-marsh. The annual average density of adult rabbits in this area during 1977–78 was not directly measured but assumed to be about 7–9 individuals/ha (Lufner, 1986). After RHD, annual average rabbit abundance in the DBR has remained consistently at levels five or six times below those registered before the disease outbreak (Villafuerte & Moreno, In press).

2. Matasgordas (MG): situated also in the ecotone scrubland-marsh but 15 km north from DBR (Fig. 1). The dominant shrub in this area was Pitaça lentircuz, and Quevcur ruvest and Fraxinus sp. trees were abundant. Though it is situated only 15 km north from the DBR, for unknown reasons the incidence of RHD in this area has been low and average rabbit density has been estimated as 11–18 ind/ha (Palomares et al., 1995).

Fig. 1. Sketch map of the Doñana National Park with the location of the two study areas: 1) Doñana Biological Reserve; 2) Matasgordas. Park limits are shown by a continuous line and other protected areas are indicated by a dashed line. Badgers from each area do not range in the other.
Table 1. Frequency of occurrence and percentage of biomass of each prey type in each study area and period, and diversity (Levins Index values). In brackets sample size. MG = Matasgordas. DBR = Doñana Biological Reserve. RHD = Rabbit Haemorrhagic Disease

<table>
<thead>
<tr>
<th></th>
<th>MG</th>
<th>DBR after RHD</th>
<th>DBR before RHD</th>
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<tr>
<td></td>
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<td>% Biomass</td>
<td>% Occurrence</td>
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<tr>
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<tr>
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<tr>
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Fig. 2. Correlation between dietary diversity (as Levins Index) and the percentage of ingested biomass of rabbit. 1 = Doñana Biological Reserve (DBR) before rabbit haemorrhagic disease (RHD) in wet season. 2 = DBR before RHD in dry season. 3 = DBR after RHD in wet season. 4 = DBR after RHD in dry season. 5 = MG in wet season. 6 = MG in dry season.

while no significant differences were detected with regard to DBR after RHD during the dry season.

Ingested young rabbit biomass declined about 50% during the wet period and 33% during the dry period in the DBR after the outbreak of RHD (Table I). In both wet and dry periods, frequencies of occurrence of rabbit varied significantly with regard to DBR before RHD and to MG (P<0.001). Nevertheless, young rabbits continued to be an important food item, particularly during the wet season (41%). Young rabbits were not replaced by a single prey category in the badger's diet in any season. Alternative foods included insects and birds in the wet season and fruit, insects and birds in the dry season (Table I). Dietary diversity almost doubled during the wet season and increased 1.5 times during the dry season following the RHD outbreak (Table I).

The percentage of rabbit biomass in the diet was negatively correlated with food diversity (r_s = —1, n = 6, P < 0.001; Fig. 2), suggesting that dietary diversity was related to the availability of young rabbits and relatively independent of the availability of food types other than rabbits. There was a significant positive correlation between diet diversity and percentage of bird biomass (P < 0.05), suggesting that birds were a typical complementary food.

**DIFICUSSION**

Given the remarkable geographical, seasonal and individual variations detected in Eurasian badger diets (e.g. Lüps & Wandeler, 1993), some authors support the traditional view that the species is an opportunistic food generalist (e.g. Roper, 1994; Roper & Mickevicius, 1995). Other authors describe the foraging behaviour of some Eurasian badger populations as being more characteristic of specialized feeders, relying locally on single resources, such as earthworms (Lruuk & Parish, 1981), some fruit (Lruuk & de Lock, 1981), insects and fruit (Pigozzi, 1991) or young rabbits (Martín et al., 1995).
The more simplistic interpretation of our results suggests that badgers in the Doñana area are opportunistic feeders, because they changed their diet in DBR after rabbits became scarce. However, such a change in diet diversity of facultative specialists when the encounter rate with preferred prey declines is predicted by most theoretical models of food specialization (e.g. Stephens & Lurue, 1986). In fact, one of the axioms of classical foraging theory is that dietary diversity of specialists is governed by encounter rate with the more profitable prey, regardless of abundance of other food resources (Futuyma & Moreno, 1988). In our case, availability of rabbits seems to be the key to badger’s diet diversity (Fig. 2). Despite the change in rabbit numbers, there was no possibility of a change in the availability of rabbits per individual badger, if the numbers of badgers had also decreased. However, we were unable to detect a short-term numerical response of predators, including badgers, to the rabbit decline after RHD (Delibes et al., 1992).

Our results suggest that, only when rabbit became scarce in DBR after RHD, was the diet diversity increased. We assume that the seasonal availability of insects and fruit in DBR was not noticeably modified by the RHD outbreak; nevertheless, these seasonal changes in secondary prey abundance, very conspicuous in the DBR badger diet after RHD (Table 1), were not reflected in the diets where and when rabbits were abundant.

According to Brown & Morgan (1995), the forager’s diet results from a two-step decision process: select where to seek food, and once there, select what to eat. Probably Doñana badgers could be considered selective predators because they start searching for young rabbits by foraging on good rabbit breeding grounds, independently of the abundance of alternative foods which could be eaten in an opportunistic way, as they were found while searching for rabbits (Martin et al., 1995; Rodríguez, Martín & Delibes, 1996).

A question remains as to why Eurasian badgers apparently behave as a selective predator in different places. Obviously, they cannot be considered morphological, physiological or even behavioural specialists in the evolutionary sense of Futuyma & Moreno (1988). Paradoxically, it seems they are opportunistic to a point which enables them to adapt their feeding behaviour to act locally as true specialists, on such different food resources as earthworms or rabbits.

The described dietary response of the Doñana badgers to a decline in preferred prey was similar to that of some Scottish badger populations when earthworms became scarce (Lurue, 1989). In the long term, we could expect a change in the Doñana badger population size, although we hope the species will not disappear, as happened in the Ardnish study area of Lurue (1989) following a change in land use which gave rise to a decrease in the availability of earthworms.

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Diet of two species of mink in Estonia: displacement of Murtela lutveola by M. riron

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Abstract

During a period in the 1980s when both European mink Murtela lutveola and American mink M. riron were present in Estonia, their food was significantly different. European mink ate a greater proportion of fish and crustaceans, whereas American mink took relatively more mammals and frogs. This was probably related to a difference in habitat selection. After the disappearance of the European mink, the diet of the American mink in our main study area was similar to that of the European in the same area previously. Two alternative hypotheses are presented for the mechanisms which led to the replacement of European mink by the American species: (i) the two species have a different niche, and the American mink could replace the European mink after the latter had disappeared for unrelated reasons, or (ii) the American mink aggressively ousted the European mink, a process starting in the American mink’s preferred habitat (slow flowing rivers). At present there are insufficient data to reject either of these scenarios.

Ley wordz: diet, mink, habitat, competition, extinction

INTRODUCTION

The decline of the European mink Murtela lutveola in the second half of the 20th century has been catastrophic, and now this species is at risk of being the next mammalian extinction in Europe. Amongst the hypotheses that seek to explain the decline, several concern the relationship between the endemic European mink and the introduced American mink M. riron. In particular, questions arise of evidence of competition for food, niche overlap and displacement, between the species. One way to test such a competition hypothesis is to make use of the natural experiment provided by the disappearance of the European mink, and compare the American mink’s diet before and after.

In this paper, we provide data on the food of the European mink and the American mink over a large area of Estonia. After the European mink disappeared from the country, data were collected on diet of the American mink along one of the rivers, and they were compared with the diet of European mink during its occupancy of the same river in the preceding period.

The course of the decline since the early 1970s, and final disappearance of the European mink from Estonia after 1992, have been well documented (Maran, 1991; Maran & Henttonen, 1995). It parallels the rapid decline of the species elsewhere in Eastern Europe, in contrast to its performance in Spain where numbers are steady or increasing (Bárta, 1956; Szunyoghy, 1974; Schreiber et al., 1989; Romanowski, 1990; Palomares, 1991). Despite our rather detailed knowledge of the Eastern European decline and enumeration of possible causes (Maran & Henttonen, 1995; Maran et al., in press), no satisfactory single explanation has been put forward, and no simple solution for the conservation of European mink is yet available.

The demise of European mink in Estonia coincided with the establishment there, in the early 1970s, of the American mink (Maran, 1991), although the European mink had been in long-term decline even before the arrival of its exotic congener. One of the possible, final causes of the European mink’s disappearance, therefore, could be simple competition for resources with a