

## COEXISTENCE OF TERRITORIALITY AND HAREM DEFENSE IN A RUTTING FALLOW DEER POPULATION

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Male fallow deer (*Cervus dama*) on the Swedish island of Öland (Espmark and Brunner, 1974) and in South Weald, England (Chapman and Chapman, 1975) apparently defend territories with a central area (rutting stand) heavily trampled and marked with the products of scent glands and urine. Courtship and copulation with visiting females occur there. Lek-breeding has been reported for fallow deer in Schleswing-Holstein, north Germany (Heidemann, 1973), Jaegersborg, Denmark (Schaal, 1986; Schaal and Bradbury, 1987), Gyulaj, Hungary (Pemberton and Balmford, 1987), San Rossore, central Italy (Apollonio et al., 1989, 1990), and Petworth, England (Clutton-Brock et al., 1988; Pemberton and Balmford, 1987); some males at the last two localities also defended individual-resource territories.

At Doñana National Park, in southern Spain, males follow and defend groups of females in a typical deer strategy of harem following (Alvarez et al., 1975). In some other areas the establishment of leks apparently is associated with high population densities (Clutton-Brock et al., 1988). The absence of any previous observation of leks at Doñana in spite of relatively high concentration of animals led us to investigate the mating system of individually recognizable males. Because females concentrate in highly productive patches of grass at the study area during the rut, a time of low food availability, the space perhaps becomes economically defensible for males at this time.

Doñana National Park is located at the estuary of the Guadalquivir River (37°00'N, 6°26'W). The climate is Mediterranean, with hot dry summers and mild wet winters. Annual rainfall is about 530 mm, falling mostly in winter. Mean temperatures range from a January low of 5°C to July and August highs of 35°C.

The dominant features of the park are a seasonal marshland and an area of sands covered by low shrub; the study was concentrated in the ecotone between the two, where fallow deer commonly occur (Braza and Alvarez, 1987). A 260-ha area located in the less disturbed zone, the inner reserve in its southern part, was chosen for our observations. The vegetation consists of low shrubs, brackens, meadows, meadows with rushes, and dry marsh. The area was fenced in a short (700 m) curvilinear segment on its SE side (Fig. 1).

The rut of fallow deer at the study area starts at the beginning of October and lasts about 3 weeks; antler casting occurs between the end of March and the beginning of May. Young are born at the end of May and beginning of June (Braza et al., 1986). At rutting time after the dry summer, the marsh (of lowest elevation among the habitat types) becomes a vast hard-baked flat covered with dead vegetation. The only relatively moist zones are the meadows and the meadows with rushes, because of moisture reserves from a higher water table beneath the more elevated shrub area. As a result, in October, the meadows are well defined with high productivity (mean aboveground dry biomass in October 1982 = 299.3 g/m<sup>2</sup>) as compared to the shrub area (biomass too small to measure) and the dry marsh (60 g/m<sup>2</sup>). Fallow deer do not eat bracken (Palacios et al., 1984; Venero, 1984).

At 10-day intervals from 3 August 1986, all individuals of each class (adult and subadult males, females >1 year of age, fawns, and yearlings) were counted and their positions plotted on a map of the study area (1:2,000) divided into 1-ha squares. Rutting-season observations started on 1 October 1986 and lasted 20 days. By video-recording males observed in the study area and observing the antlers, most of the 49 males could be recognized individually.

Unlike those in England and central Italy (Apollonio et al., 1989, 1990; Clutton-Brock et al., 1988), fallow deer on our study area are not active at midday during the rut. Their activity is largely nocturnal (Braza et al., 1984), a situation apparently common for mammals of the National Park during warm and dry months (Alvarez et al., 1981). Accordingly, we observed the fallow deer from 0600 to 0900 h and from 1600 to 1930 h daily during the 20-day rut.

The study area was divided into four zones, each assigned to an observer, who, positioned in a tower (4-30 m high), registered the aggressive behavior between males (fighting, chasing, displacing, and parallel walking). The positions of all animals present at 0630 h and 1700 h were plotted on a map. Each observation bout lasted 10-15 min and was divided into 1-min intervals. Each male, in turn, was observed.

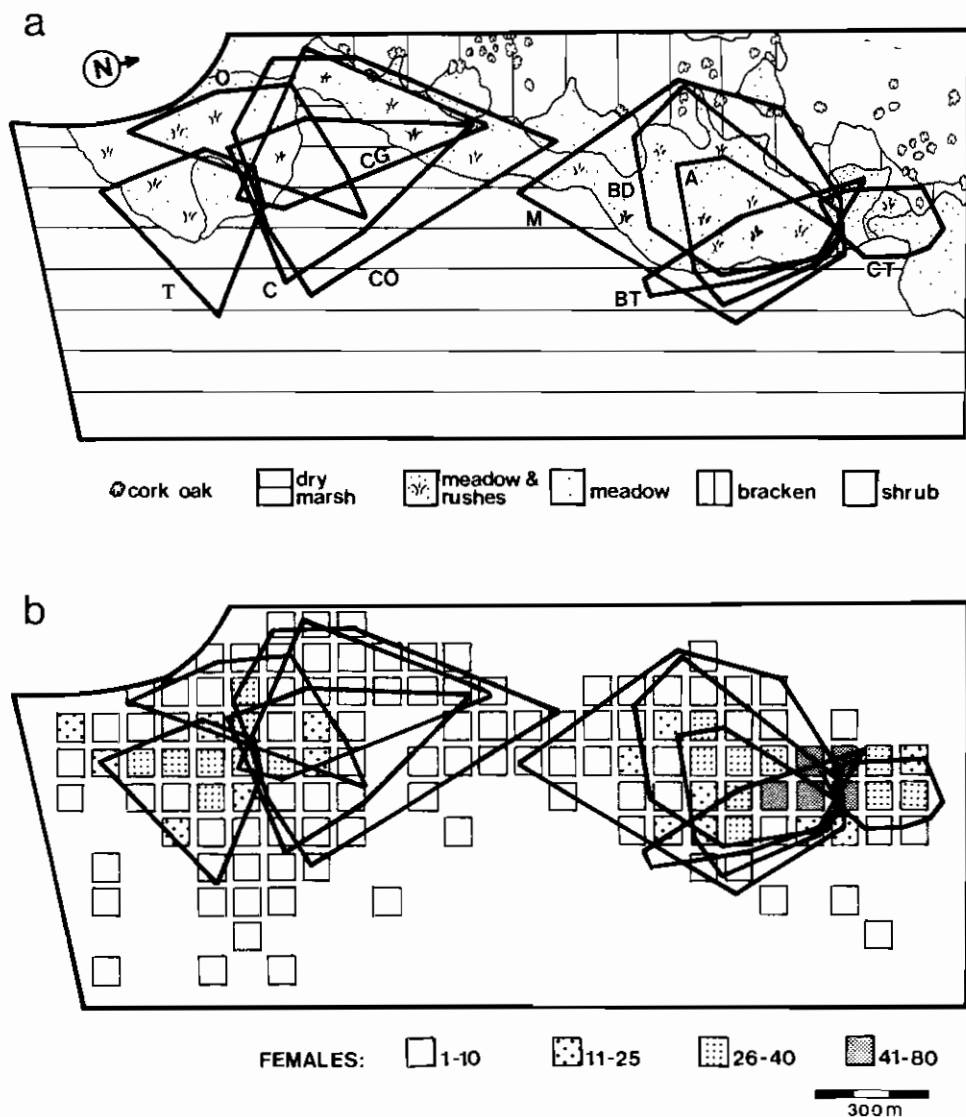


FIG. 1.—Convex polygons of: a, the 10 sedentary fallow deer males during the 1986 rut at Coto Doñana, SW Spain, in the environmental units of the study area based on sighting frequencies; and b, of the 10 sedentary males superimposed upon cumulative number of sightings of females during the rut in each 1-ha square. No female was observed in the open spaces. Letters refer to individual males; CT and T were the two territorial males.

An exhaustive search for deer in the study area every 10 days from 2 months before the rutting started provided the number of males and females for each of the 11 sampling days (males,  $\bar{X} = 64.4$ ,  $SD = 16.9$ ; females,  $\bar{X} = 124.8$ ,  $SD = 13.0$ ). Neither the number of animals of each sex nor the sex ratio during the same period ( $\bar{X} \pm SD = 1:2.1 \pm 0.7$ ,  $n = 11$ ) showed great variation. The density on the study area was about 73 adults/km<sup>2</sup>, which, together with fawns and yearlings, totaled 160 animals/km<sup>2</sup>.

Of the 710 sightings of males during October, 266 were of 10 males located in the study area at least 20 times ( $\bar{X} = 26.6$ ,  $SD = 6.1$ ; range, 20–39). Analysis of behavior was undertaken on these sedentary males, all of them  $\geq 5$  years of age. The limits of the rutting areas of these males were defined by connecting the

outer locations (convex polygons); surface areas of the polygons were measured by dividing each into a network of triangles and summing their areas.

In October, both sexes were found most often in the meadows with rushes, meadows, and dry marsh, in decreasing order. The same applied for the 10 sedentary males, for which geometric centers of locations were mainly in meadows with rushes (six), dry marsh (three), and meadows (one;  $\bar{X} = 15.4$  ha,  $SD = 9.5$ ; range, 4.7–31.5). These 10 rutting areas were arranged in two clusters (Fig. 1a).

Although there is an overlap in rutting areas of individual males, there is much variation in the areas of exclusive use ( $\bar{X} = 29.6\%$ ,  $SD = 35.2$ ,  $n = 10$ ; range, 0–90.8); only one individual in each cluster showed almost exclusive use of its rutting area (Fig. 1a).

Most aggressive interactions among males of each cluster involved the territorial male and the next most aggressive individual (39.2% in the northern cluster and 43.9% in the southern one) although the two territorial males fought all their neighbors. At no time during the rutting season were either rutting stands (Espmark and Brunner, 1974) or leks observed in the study area. The sedentary males spent most of the most of the time in their rutting areas, following and vigorously herding females feeding therein. Each male retained only temporary control over female groups because females moved freely in and out of a male's zone. The 10 sedentary males occupying the study area, therefore, can be classified as either territorial or harem follower. No other mating strategy was observed.

The location of females in the study area corresponded to a great extent to the rutting areas of the 10 sedentary males (Fig. 1b). Of 1,562 sightings of females during the rutting period, 46% were within the convex polygons of the two territorial males.

Contrary to the high number of copulations observed by Clutton-Brock et al. (1988) in England and by Apollonio et al. (1989, 1990) in central Italy, where fallow deer appear to be active for the whole daylight period, we observed only seven copulations. This led us to think that they must copulate mostly at night. Of the seven copulations observed, five involved territorial males.

The two territorial males defended their rutting areas even when no females were present; they did not attempt to court females outside their spatial boundaries. This characterizes their behavior as defense of the space itself for reproductive purposes, hence territoriality (Noble, 1939). Because females use the area for obtaining food resources, no leks occur (Bradbury and Gibson, 1983).

Inasmuch as the zones of highest density of females are mostly within the two observed territories, defense of those high resource areas, with the resulting increase of interceptions of female movements, is probably the first option for any male. As a second choice, a male may resort to following female groups, defending them against other males. Thus, concentration of females and their predictable movements probably are factors promoting rutting territoriality in male fallow deer, as proposed for various species by Emlen and Oring (1977), Gosling (1986), and Owen-Smith (1977). Red deer (*C. elaphus*) living in the study area overwhelmingly resort to resource territoriality as a mating strategy (Carranza et al., 1990), establishing their territories also in the productive ecotone between the dry marsh and the shrub.

According to Bradbury et al. (1986), at relatively low densities of females leks would not occur, because the derived reproductive benefits would not exceed the cost of aggregation to males. However, density of females is not the only factor bringing about leks in fallow deer. Leks are common at San Rossore, where a density as low as 19 animals/km<sup>2</sup> and sex ratio of 1:1.2 in favor of females exists (Apollonio et al., 1989, 1990).

If concentration of food resources leads to territoriality, this mating strategy should not be restricted, among the Cervidae, to roe deer (*Capreolus capreolus*—Bramley, 1970; Kurt, 1968), sika deer (*C. nippon*—Miura, 1984) and red deer (Carranza et al., in press), but should be a common strategy.

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