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Breeding season avifauna of prairie dog colonies and non-colonized areas in shortgrass prairie

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Avian assemblages of black-tailed prairie dog (*Cynomys ludovicianus*) colonies were compared to those present at two types of non-colonized grassland, during the breeding season, in southwestern Kansas and southeastern Colorado. Relative abundances were quantified in 1996 during a period of drought and in 1997 during a period of above average precipitation. We detected fewer bird species at all sites in 1996 than in 1997, and the total number of bird species detected on prairie dog colonies was lower than that detected on both types of non-colonized areas during both years. Horned larks (*Eremophila alpestris*) had higher relative abundances on non-colonized sites in 1996, but were more abundant on prairie dog colonies in 1997. Lark buntings (*Calamospiza melanocorys*) and grasshopper sparrows (*Ammodramus savannarum*) were not recorded at the study sites in 1996, but appeared in high numbers in 1997. These two species, and western meadowlarks (*Sturnella neglecta*), had higher relative abundances on non-colonized sites in 1997 than on prairie dog colonies. A comparison of our results with those of other studies suggest that the effects of prairie dogs on associated biota might not be identical in all regions of the Great Plains, or under all environmental conditions.

Keywords: prairie dogs, grassland birds, shortgrass prairie.

INTRODUCTION

Prairie dogs (*Cynomys* spp.), through their burrowing activities and herbivory, have the ability to create localized communities of plants and animals that differ considerably from surrounding grasslands (Whicker and Detling 1988). The importance of prairie dog ecosystems to numerous vertebrates in the Great Plains has been demonstrated (Kotliar, Baker *et al.* 1999), and increased awareness of this role has prompted appeals for heightened protection of prairie dogs and changes to the management of public and private rangelands (Miller, Wemmer *et al.* 1990; Miller, Ceballos and Reading 1994;

Davitt, Grandi *et al.* 1996). In spite of their importance, black-tailed prairie dogs (*Cynomys ludovicianus*) have declined by 98% in the past century, and due to continued recent declines, the U.S. Fish and Wildlife Service designated the species as a candidate for listing in 2000 (U.S.D.I. 2000).

Accurate information on how prairie dogs affect flora and fauna in different areas of their range will contribute significantly to land management efforts. To date, only two studies have been conducted which compare the avian assemblages of prairie dog colonies to non-colonized sites (Agnew, Uresk and Hansen 1986; Barko, Shaw and Leslie 1999).

Our purpose was to quantify differences in breeding season avian assemblages among prairie dog colonies and non-colonized sites in the shortgrass prairie of southwestern Kansas and southeastern Colorado. These findings will enhance prairie dog conservation as well as cooperation of land managers, livestock producers, conservationists, and policy makers in ensuring the maintenance of biotic integrity and economic vitality of western rangelands.

STUDY SITES AND METHODS

Study sites were located on Cimarron National Grassland and adjacent private land in Morton County, southwestern Kansas and Baca County, southeastern Colorado. Cimarron National Grassland comprises greater than 43,700 ha of predominantly grazed land administered by the U.S. Forest Service. Most of the surrounding private land is cropped but some areas remain in perennial grass cover and are grazed by cattle. Study sites were on silty loam soils with 0 to 6% slopes. Study sites were located in a relatively flat landscape north of the Cimarron River, characterized by short and mid-height grasses. Although croplands were adjacent to or near many of the study sites, all study sites were contiguous with larger blocks of grassland vegetation. Woody vegetation in the study landscape was scarce and restricted to plantings around farmsteads, and likely had no measurable influence on our results. Mean yearly precipitation recorded at the Elkhart weather station in Morton County between 1901 and 1996 was 44.75 cm (National Weather Service Cooperative Observer Network, pers. com.). Precipitation during the 12 months preceding the 1996 sampling period was 34.3 cm, 76% of the long-term mean. Precipitation during the 12 months preceding the 1997 sampling period was 68.0 cm, 152% of the long-term mean.

Whereas many areas at Cimarron National Grassland are characterized by mid-height

vegetation, which might reflect the composition of seed mixes used to reclaim abandoned farmland in the 1930s to 1950s (Schumacher and Atkins 1965), our initial sampling efforts were directed at non-colonized areas that we assumed were characteristic of a shortgrass region. In 1996, we selected for sampling every site we were aware of on Cimarron National Grassland that was dominated by the perennial shortgrasses *Bouteloua gracilis* and *Buchloe dactyloides* ($n = 5$, mean = 42.10 ha, range = 32 - 64 ha, total approximately 210 ha). For comparison we selected eight of the largest prairie dog colonies on Cimarron National Grassland (mean = 41.49 ha, range = 20 - 64 ha, total approximately 332 ha).

All sites sampled in 1996 were re-sampled in 1997 except two prairie dog colonies; one colony had been destroyed by plague (Cully, Carter and Gage 2000) and the boundaries of the other had changed sufficiently to make it unsuitable for sampling. Completion of the 1996 field season had indicated that it would be logistically possible to sample more sites. In 1997, seven additional prairie dog colonies on Cimarron National Grassland and adjacent private lands ($n = 13$, mean = 39.25 ha, range = 8 - 125 ha, total approximately 555 ha) and one additional non-colonized shortgrass site on private land ($n = 6$, mean = 43.18 ha, range = 32 - 64 ha, total approximately 260 ha) were sampled.

In 1997, we thought it would be informative to quantify the avian assemblages of a third habitat type present in the study area, non-colonized areas characterized by the presence of mid-height grasses, and compare the data to that of each of the other two treatments in the study. Using a list of potential sites with soil types and slopes identical to those found on the prairie dog colonies, we randomly selected eight non-colonized sites on Cimarron National Grassland for sampling in 1997 (mean = 56.68 ha, range = 32 - 64 ha, total approximately 454 ha). Data collected

in 1997 from non-colonized shortgrass sites and non-colonized random sites were not pooled for analyses, each of the non-colonized treatments was analyzed separately.

Vegetation sampling that occurred subsequent to our study (mid-June to mid-July) in 1996 and 1997, demonstrated that each of the three treatments were characterized by distinct vegetation structure and composition (Winter, Cully and Pontius 2002). Dominant vegetation of prairie dog colonies consisted of the perennial shortgrasses *Aristida purpurea*, *Bouteloua gracilis* and *Buchloe dactyloides*. Dominant vegetation of the non-colonized shortgrass sites was *Bouteloua gracilis* and *Buchloe dactyloides*, while the vegetation of non-colonized random sites typically included the mid-height perennial grass *Bouteloua curtipendula* as a co-dominant with *Bouteloua gracilis* and *Buchloe dactyloides*. Vegetation composition during the period when vegetation sampling was conducted likely did not differ greatly from that present one and one-half months earlier when avian sampling occurred during our study. The vegetation that would most likely contribute to a marked seasonal difference, cool season perennial grasses and winter annual grasses, are minor components of the vegetation of this region (Lauenroth and Milchunas 1992), and of our study sites specifically (Winter 1999; Winter, Cully and Pontius 2002).

Strip transects (200 m wide) were established at each study site and varied in length from 0.4 to 1.6 km. One to five parallel transects were sampled at each study site; the length and number of transects varied in proportion to the size of the site, and transects were separated by 250 m. In 1996 we surveyed 12.5 km of transects on prairie dog colonies and 8.5 km of transect at non-colonized shortgrass sites. In 1997 we surveyed 11.3 km of transects on prairie dog colonies, 9.5 km of transects on non-colonized shortgrass sites and 13.8 km of transects on non-

colonized random sites. We standardized the data to the number of individuals detected/km of transect (relative abundance). Relative abundances at each site were used to calculate mean relative abundance for each treatment.

Birds were surveyed along the transects at each study site only once each year, between 27 May and 18 June, and all data were collected by SLW. Data were collected between sunrise and 10:30 am CDST, when it was not raining and winds were less than 30 km/hr. All sitting, perching and flushing birds, perpendicular or forward of the observer, and within the boundaries of the transect, were recorded. Birds flying over the study site and birds perched on or within 20 m of livestock watering facilities were not recorded.

Our impression in 1996 was that vegetation structure of the prairie dog colonies was identical to that found on the non-colonized shortgrass sites, and we chose not to measure vegetation structure that year. In 1997, vegetation structure was measured using a visual obstruction pole modified from Robel, Briggs *et al.* (1970) at randomly located points along each transect. The number of points ranged from 35 to 120, in proportion to the area of the site. Observations were made one meter from the pole at a height of one meter. Vegetation height was determined by recording the highest point at which vegetation crossed between the observer and the pole. Visual obstruction was determined by recording the lowest point at which the pole was visible.

The response variable for statistical tests was relative abundance, expressed as number of individuals detected per kilometer of transect at each site. Two-tailed tests were used to test null hypotheses that there were no significant differences among the treatments (two treatments in 1996 - prairie dog colonies and non-colonized shortgrass sites; three treatments in 1997 - prairie dog colonies, non-colonized shortgrass sites and

non-colonized random sites). We chose two-tailed tests because we did not want to conduct the study with the assumption that avian abundances would be higher on one treatment than on another, in which case one-tailed tests would have been appropriate. For statistical analysis a one-factor analysis of variance (ANOVA), using SAS v. 6.11 proc glm (SAS Institute 1996), was used when the assumptions of the F-test were satisfied. When the assumptions of the F-test were not satisfied, the Kruskal-Wallis nonparametric test, using SAS v. 6.11 proc npar1way (SAS Institute, 1996), and a Tukey-type non-parametric multiple comparison method (Zar 1996) were used. We set $\alpha = 0.05$ for hypothesis tests. A test statistic value of $Q > 2.39$ generated by the Tukey-type test indicated significant differences. Tables 1 and 2 provide P-values from analysis of variance and Kruskal-Wallis tests. When data from three treatments were used in statistical tests, significant Q and P-values from pairwise comparisons are provided in the text of the results section. In instances where individuals of a particular species were not detected at any study replicates of a particular treatment, that treatment was not included in statistical analyses. This was done because the treatment containing all zeros contributes no within treatment variability to the test statistic but it does contribute to the among treatment variability.

RESULTS

In 1996, 136 individuals of five species were recorded at prairie dog colonies and 136 individuals of seven species were recorded at the non-colonized shortgrass sites (Table 1). The higher relative abundance of horned larks (*Eremophila alpestris*) at non-colonized shortgrass sites was significantly different from that found on prairie dog colonies that year. There was no significant difference in the relative abundances of all birds combined, killdeer (*Charadrius vociferus*), mourning

Table 1. Relative abundance (individuals/km of transect) and total number detected (#) on prairie dog colonies (n = 8) and non-colonized shortgrass sites (n = 5) in 1996. Model df = 1, error df = 11, P-values are from F-tests or Kruskal-Wallis tests. Treatments in which no individuals were detected were not included in statistical tests. Response variable is relative abundance.

	Prairie dog colonies		Shortgrass sites		P-value
	mean \pm SE	#	mean \pm SE	#	
All birds combined	11.11 \pm 1.82	136	16.07 \pm 1.13	136	0.072
Killdeer <i>Charadrius vociferus</i>	0.16 \pm 0.16	3	0.08 \pm 0.08	1	0.523
Mountain plover <i>Charadrius montanus</i>	0	0	0.08 \pm 0.08	1	
Mourning dove <i>Zenaida macroura</i>	0.23 \pm 0.16	4	0.08 \pm 0.08	1	0.692
Burrowing owl <i>Athene cunicularia</i>	0.98 \pm 0.76	7	0	0	
Common nighthawk <i>Chordeiles minor</i>	0	0	0.17 \pm 0.17	2	
Western kingbird <i>Tyrannus verticalis</i>	0	0	0.08 \pm 0.08	1	
Horned lark <i>Eremophila alpestris</i>	7.46 \pm 1.45	91	14.91 \pm 1.26	123	0.005
Western meadowlark <i>Sturnella neglecta</i>	2.28 \pm 0.7	31	0.66 \pm 0.38	7	0.158

doves (*Zenaida macroura*), and western meadowlarks (*Sturnella neglecta*) in 1996.

In 1997, 241 individuals of seven species were recorded on prairie dog colonies, 297 individuals of eight species were recorded on non-colonized shortgrass sites, and 552 individuals of nine species were recorded on non-colonized random sites (Table 2). The higher relative abundance of horned larks on prairie dog colonies was significantly different from that found on non-colonized random sites ($P = 0.003$) that year. In 1997, the higher relative abundance of lark buntings (*Calamospiza melanocorys*) on non-colonized shortgrass sites was significantly different from that found on prairie dog colonies ($P = 0.014$). The higher relative abundances of all birds combined ($P = 0.001$), western meadowlarks ($Q = 2.62$), lark buntings ($P < 0.001$), and grasshopper sparrows (*Ammodramus savannarum*, $Q =$

Table 2. Relative abundance (individuals/km of transect), total number detected (#), and vegetation height and density on prairie dog colonies (n = 13), non-colonized shortgrass sites (n = 6), and non-colonized random sites (n = 8) in 1997. Model df = 2, error df = 24, P-values are from F-tests or Kruskal Wallis tests. Different superscripts indicate significant differences from pair-wise comparisons. P and Q-values for significantly different pair-wise comparisons are provided in the text. Treatments in which no individuals were detected were not included in statistical tests. Response variable is relative abundance.

	Prairie dog colonies		Shortgrass sites		Random sites		P-value
	mean ± SE	#	mean ± SE	#	mean ± SE	#	
All birds combined	22.59 ± 1.88 ^a	241	30.88 ± 4.20 ^{a,b}	297	40.19 ± 4.65 ^b	552	0.002
Mountain plover <i>Charadrius montanus</i>	0.02 ± 0.02	1	0	0	0	0	
Long-billed curlew <i>Numenius americanus</i>	0	0	0.53 ± 0.31	5	0.38 ± 0.25	5	0.452
Mourning dove <i>Zenaidura macroura</i>	1.11 ± 0.51	9	0.23 ± 0.16	3	0.83 ± 0.45	10	0.717
Burrowing owl <i>Athene cunicularia</i>	0.54 ± 0.24	7	0	0	0	0	
Common nighthawk <i>Chordeiles minor</i>	0	0	0.07 ± 0.07	1	0.58 ± 0.19	7	0.054
Horned lark <i>Eremophila alpestris</i>	16.48 ± 1.64 ^a	172	10.51 ± 4.07 ^{a,b}	101	6.78 ± 1.13 ^b	89	0.008
Lark bunting <i>Calamospiza melanocorys</i>	2.18 ± 0.97 ^a	32	11.92 ± 3.43 ^b	115	18.31 ± 3.81 ^b	254	0.0002
Grasshopper sparrow <i>Ammodramus saviannarum</i>	0.27 ± 0.18 ^a	4	4.34 ± 1.49 ^{a,b}	44	8.22 ± 1.28 ^b	126	0.0001
Dickcissel <i>Spiza americana</i>	0	0	0	0	0.39 ± 0.39	5	
Red-winged blackbird <i>Agelaius phoeniceus</i>	0	0	0	0	0.23 ± 0.23	3	
Western meadowlark <i>Sturnella neglecta</i>	1.84 ± 0.89 ^a	16	2.91 ± 0.85 ^{a,b}	26	4.19 ± 0.72 ^b	51	0.028
Brown-headed cowbird <i>Molothrus ater</i>	0	0	0.28 ± 0.28	2	0	0	
Unidentified species	0	0	0	0	0.26 ± 0.17	2	
Vegetation height (cm)	9.48 ± 0.80 ^a		25.22 ± 5.50 ^{a,b}		33.59 ± 3.29 ^b		0.0008
Visual obstruction (cm)	2.49 ± 0.17 ^a		5.40 ± 1.22 ^{a,b}		6.41 ± 1.32 ^b		0.002

4.19) on the non-colonized random sites were significantly different from those found on the prairie dog colonies in 1997. Higher values of vegetation height (Q = 3.66) and visual obstruction (Q = 3.39) on the non-colonized random sites were significantly different from those found on the prairie dog colonies (Table 2) in 1997.

DISCUSSION

While the sample sizes in our study are relatively small, and sampling occurred only once during each of two years, our results provide useful insight into the ecology of grassland avifauna, and the influence that

prairie dogs might have on this community. During the two years of our study, which were characterized by markedly contrasting climatic conditions, we recorded a dramatic difference in the abundances of numerous species, especially lark buntings and grasshopper sparrows. There are numerous reports from throughout the Great Plains of grassland bird populations fluctuating dramatically in association with climatic variation (Wiens 1974; George, Fowler *et al.* 1992; Igl and Johnson 1995). The drastic population fluctuations that we recorded for lark bunting and grasshopper sparrow might represent a coping strategy whereby their populations are opportunistically “tracking”

optimum resources over a large scale as they shift in time and space (Wiens 1974; Cody 1985).

Similar to the results of other studies (Agnew, Uresk and Hansen 1986; Barko, Shaw and Leslie 1999; Manzano-Fischer, List and Ceballos 1999), horned larks were the most abundant species on prairie dog colonies that we sampled, comprising 67% and 71% of the individuals detected on colonies in 1996 and 1997, respectively. Like Agnew, Uresk and Hansen (1986) and Barko, Shaw and Leslie (1999), we also found burrowing owls (*Athene cunicularia*) associated only with prairie dog colonies. The relative abundance of burrowing owls on prairie dog colonies, however, was not suitable for statistical analysis because of their total absence from the other treatments. Additional surveys for burrowing owls in Morton and Baca counties, conducted concurrently with our study, found burrowing owls almost exclusively on prairie dog colonies (Winter 1999). Although prairie dog colonies occupied less than 2% of the land area at Cimarron National Grassland in Morton County in 1997 (J. Chynoweth, USDA Forest Service, pers. com.), burrowing owls were rarely sighted at non-colonized locations, regardless of the sampling or survey methodology (Winter 1999). Prairie dog colonies in our study area appear to be of great importance to burrowing owls during the breeding season.

Our results do not suggest that the number of bird species on prairie dog colonies was higher than that found on non-colonized sites, and this contrasts with the results of research conducted by Agnew, Uresk and Hansen (1986) in South Dakota mixed-grass prairie. In our study, the total number of bird species detected on prairie dog colonies, as well as the number of species/km of transect, was lower than that found on non-colonized shortgrass sites in 1996, and lower than that found on non-colonized shortgrass sites and non-colonized random sites in 1997. Further

contrasting with the results of Agnew, Uresk and Hansen (1986), in our study relative abundance of all birds combined was statistically similar between colonies and non-colonized shortgrass sites during both years, and higher on non-colonized random sites than on colonies in 1997. Agnew, Uresk and Hansen (1986) consistently found higher abundances of all birds combined on prairie dog colonies than on non-colonized mixed-grass prairie during all sample periods.

Agnew, Uresk and Hansen (1986) reported that abundances of five species [horned lark, mourning dove, killdeer, barn swallow (*Hirundo rustica*), and burrowing owl] were higher on prairie dog colonies, whereas only three species [red-winged blackbird (*Agelaius phoeniceus*), upland sandpiper (*Bartramia longicauda*), and lark bunting] had significantly higher abundances on non-colonized sites. In our study, three species (lark bunting, grasshopper sparrow, and western meadowlark), had a statistically significant higher relative abundance on at least one of the non-colonized treatments than on prairie dog colonies in 1997. In spite of the inability to statistically compare the relative abundances of common nighthawk (*Chordeiles minor*) on prairie dog colonies and non-colonized random sites in 1997, their relative abundance on non-colonized random sites seems biologically significant and deserves notice. Our data suggest common nighthawk avoid prairie dog colonies, preferring instead the habitat present on non-colonized random sites, which were characterized by mid-height grasses.

Horned lark was the only species in our study to have a statistically significant higher relative abundance on prairie dog colonies than on a non-colonized treatment (prairie dog colonies vs. random sites in 1997). While horned larks were the most abundant species on prairie dog colonies during both years of our study, their abundances were higher on non-colonized shortgrass sites than

on prairie dog colonies in 1996. In 1997 abundances of horned larks were statistically similar when colonies were compared to non-colonized shortgrass sites. This is surprising, considering this species affinity for habitats characterized by short, sparse vegetation and bare ground (Beason 1995), and previous research in a mixed-grass region which found much higher abundances of horned larks on prairie dog colonies relative to non-colonized areas (Agnew, Uresk and Hansen 1986). However, the prairie dog colonies and non-colonized shortgrass sites we sampled might have represented habitats for horned larks that are not as dramatically different as colonized and non-colonized habitats in mixed-grass regions.

Vegetation sampling at our study sites in 1996 and 1997, but subsequent to the avian sampling, indicated that prairie dog colonies were characterized by a greater percent cover of forbs, but similar percent cover of grasses and bare ground, than non-colonized shortgrass sites in 1996 (Winter, Cully and Pontius 2002). If these vegetation structural characteristics were present at the time avian sampling occurred as well, forb cover on prairie dog colonies might have provided a visual obstruction that caused prairie dog colonies to be less favored by horned larks during that year. In 1997, when our study found abundances of horned larks were statistically similar between prairie dog colonies and non-colonized shortgrass sites, vegetation height and visual obstruction were similar as well. Additionally, subsequent vegetation sampling that year demonstrated the two treatments also had statistically similar cover of forbs, grasses and bare ground (Winter, Cully and Pontius 2002).

The results of our study, and those of Barko, Shaw and Leslie (1999), suggested that the preference of horned larks for prairie dog colonies or non-colonized grasslands in a shortgrass region might be greatly influenced by the effects of climate on vegetation

productivity. During years of high vegetation productivity, there might be a marked contrast between the vegetation structure on prairie dog colonies and that found in adjacent non-colonized shortgrass prairie. During years when vegetation productivity is suppressed by drought, there might be little or no contrast in the vegetation structure of the two habitats.

Because of differences in the characteristic vegetation of shortgrass prairie and mixed-grass prairie, the influence of prairie dogs on bird populations in these two regions might be different. Mixed-grass prairie is characterized by mid-height grasses (Coupland 1992) and the characteristic effect of prairie dog herbivory is the displacement of mid-height grasses by short grasses (Coppock *et al.* 1983; Archer, Garret and Detling 1987). Both Agnew, Uresk and Hansen (1986) and Archer, Garret and Detling (1987) reported shorter vegetation on prairie dog colonies than non-colonized mixed-grass prairie. The low stature of vegetation on prairie dog colonies provides suitable habitat for species such as horned lark and killdeer, which otherwise might be unable to use mixed-grass prairie because of tall or dense vegetation (Agnew, Uresk and Hansen 1986). Conversely, the ability of prairie dogs to create patches of short-statured vegetation within shortgrass prairie is tempered by the fact that grasslands in this region of the Great Plains are characterized by grasses that are already short (Lauenroth and Milchunas 1992). Bird species that require short-statured vegetation might find non-colonized shortgrass prairie equally suitable habitat relative to prairie dog colonies.

The extent to which prairie dogs influence the bird community of a region might in large part be a function of the difference in the vegetation structure of prairie dog colonies and non-colonized sites. The vegetation structure of grasslands can be altered by climate, fire, and herbivory (Dix 1960; Sims, Singh and Lauenroth 1978; Brand and Goetz

1986; Coupland 1992; Lauenroth and Milchunas 1992). These factors, acting alone or together, might negate or enhance the contrast in vegetation structure of prairie dog colonies and associated non-colonized grasslands. The potential for a difference in the vegetation structure of a prairie dog colony and adjacent non-colonized grassland seems much greater in mixed-grass prairie than in shortgrass prairie, where lower precipitation amounts constrain primary productivity. In mixed-grass prairie, prairie dogs can create patches of short-statured vegetation that are attractive to bird species that do not readily use the taller, denser vegetation of associated non-colonized grassland. In shortgrass prairie, there is less of a structural contrast between the vegetation of colonies and non-colonized areas, and both habitats appear suitable for bird species, such as horned larks, that require short-statured vegetation.

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