Rabbit abundance relative to rainfall and plant production in northern Chihuahuan Desert grassland and shrubland habitats

David C. Lightfoot  
*University of New Mexico*, dlightfo@unm.edu

Ana D. Davidson  
*Universidad Nacional Autónoma de México, México*

Christopher M. McGlone  
*Ecological Restoration Institute, Northern Arizona University, Flagstaff, AZ*, wnan@byu.edu

Dara G. Parker  
*New Mexico State University, Las Cruces, NM*

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ABSTRACT.—Rabbit abundance patterns in the North American Southwest are poorly known, particularly relative to the potentially important influences of rainfall and plant production. We conducted a 10-year study on the comparative densities of black-tailed jackrabbits (*Lepus californicus*) and desert cottontail rabbits (*Sylvilagus audubonii*) in relation to annual rainfall and plant production in both black grama (*Bouteloua eriopoda*) grassland and creosotebush (*Larrea tridentata*) shrubland habitats in the northern Chihuahuan Desert. We found that both species of rabbits were more abundant in black grama grassland than in creosotebush shrubland and that plant production was positively correlated with rainfall at both habitats. However, rabbit densities were not positively correlated with rainfall or plant production in either habitat. In fact, both species of rabbits exhibited gradual, multiyear changes in abundance rather than the patterns of high annual variation observed in desert rodent communities. Our findings of no significant relationship between plant production and rabbit densities indicate that rabbit abundance in the northern Chihuahuan Desert may not be strongly regulated by short-term, bottom-up effects of annual variation in plant production. Long-term trends indicate that other factors such as disease or predation may be equally or more important. We suggest that future research evaluate both top-down and bottom-up influences to elucidate the temporal dynamics of Chihuahuan Desert rabbit abundance.

Key words: aboveground net primary production, desert grassland, desert shrubland, ENSO events, hares, habitats, Lagomorpha, Lepus, densities, precipitation, rabbits, Sylvilagus.
related rabbit abundance trends to temporal awareness of any studies that have quantitatively tailed jackrabbit reproduction; yet, we are not between environmental variables and black- et al. (2004) examined short-term relationships abundance are few (Davis et al. 1975). Portales temporal patterns of Chihuahuan Desert rabbit ecological studies that document the multiyear Martinez and Lopez-Portillo 2003). However, additional studies that document the multiyear Press and Price 1999, Kotler et al. 1998, Meserve et al. 2003) influences on rodent densities. Generally, positive relationships exist between desert rodent densities and variation in annual rainfall, and presumably subsequent bottom-up effects of plant production (Beatley 1969, 1974, Whitford and Steinberger 1989, Ernest et al. 2000, Meserve et al. 2003). The relative importance of bottom-up and top-down regulation of population dynamics and community organization has been of great interest in ecology. Abiotic factors are thought to be especially important in arid systems because limited rainfall controls the availability of resources. The pulse-reserve model for arid systems suggests strong bottom-up control of net primary production and in turn abundance of consumers (Reynolds et al. 2004).

The black-tailed jackrabbit (Lepus califor- nicus) and the desert cottontail (Sylvilagus audubonii) are the 2 common desert rabbit (Lagomorpha) species of the North American Southwest. Both species occur across a wide variety of habitats but prefer open semidesert mixed grass and shrublands, with cottontails additionally favoring riparian and scrub wood- lands (Anderson 1972, Schmidly 1974, Findley et al. 1975, Desmond 2004). Jackrabbits and cottontails are abundant, medium-sized mammals of Chihuahuan Desert landscapes and are known to serve as important food resources for predators such as coyotes and raptorial birds (Wagner and Stoddart 1972, Henke and Bryant 1999, Cartron et al. 2004, Bartel and Knowlton 2005). Rabbits also are known to have significant impacts through herbivory on vegetation composition and structure (Norris 1950, Steinberger and Whitford 1983, Clark and Wagner 1984, Gibbens et al. 1993, Lopez-Portillo 1996, Martinez and Lopez-Portillo 2003). However, ecological studies that document the multyear temporal patterns of Chihuahuan Desert rabbit abundance are few (Davis et al. 1975). Portales et al. (2004) examined short-term relationships between environmental variables and black-tailed jackrabbit reproduction; yet, we are not aware of any studies that have quantitatively related rabbit abundance trends to temporal variation in bottom-up environmental influences, such as rainfall and plant production, over multiple years.

The Chihuahuan Desert is characterized by a mosaic of desert grasslands and shrublands. Black grama (Bouteloua eriopoda) grasslands and creosotebush (Larrea tridentata) shrublands are common throughout the northern Chihuahuan Desert (Brown 1982, Havstad et al. 2006), and each provides different habitat structure and functional responses to rainfall. Black grama grasslands provide relatively low and homogenous vegetation structural profiles, and the shallow-rooted herbaceous vegetation grows quickly after rainfall (Ludwig 1986, Whitford 2002). Creosotebush shrublands provide taller vegetation profile habitats, but with fewer herbaceous plants. The deep-rooted perennial shrubs do not respond immediately to rainfall but rather to longer-term changes in deep soil moisture (Schlesinger et al. 1990, Reynolds et al. 2004). Fundamental differences in these habitat types may differentially influence rabbit populations; yet, studies that comparatively examine temporal trends of rabbit densities in relation to rainfall and plant production in grassland and shrubland habitats are lacking.

Temporal variation in northern Chihuahuan Desert rainfall is high, but most annual rainfall occurs during late summer (Havstad et al. 2006). El Niño Southern Oscillation (ENSO) events cause additional variable, multiannual shifts from opposing El Niño to La Niña rainfall pat- terns (Holmgren et al. 2001, Brown and Ernest 2002, Meserve et al. 2003), which drastically change interannual ecosystem productivity. This process is particularly pronounced in arid sys- tems, resulting in pulse-reserve ecosystem dy- namics (Reynolds et al. 2004). Plant production in the northern Chihuahuan Desert is also variable over space and time relative to quantities, duration, and seasonal timing of precipitation and to spatial variation in landscape geomorphology and soils (Ludwig 1986, Havstad et al. 2006).

The purpose of this study was to examine the abundance of black-tailed jackrabbits and desert cottontails in both grassland and shrubland habi- tats from a site in the northern Chihuahuan Desert in relation to measures of precipitation and aboveground plant production. Our primary research questions were the following: (1) Do the 2 habitats support different densities of each rabbit species? (2) How do the changes
in density of each rabbit species over time compare between grassland and shrubland habitats? And (3) do densities of each rabbit species vary in relation to rainfall and plant production over time; and if so, how does this relationship compare between grassland and shrubland habitats?

**Methods**

**Study Sites**

This research was conducted in the Jornada Basin in the Rio Grande Rift valley of the northern Chihuahuan Desert in southern New Mexico (Fig. 1). One study site was located in black grama grassland on a sandy plain (32°35’N, 106°51’W, 1340 m elevation; Fig. 2A) and the other in adjacent creosotebush shrubland on a gravelly, lower piedmont slope (32°30’N 106°12’W, 1340 m elevation; Fig. 2B). Both study sites were situated adjacent to above-ground net primary production (ANPP) study plots on the existing National Science Foundation, Jornada Basin Long-Term Ecological Research site (Huenneke et al. 2001, Havstad et al. 2006). The creosotebush site was dominated by creosotebush shrubs along with the less common woody shrubs honey mesquite (*Prosopis glandulosa*) and tarbush (*Flourensia cernua*), the perennial grass bush muhly (*Muhlenbergia porteri*), and many species of annual forbs. The soils consisted of gravelly loams. The black grama grassland site was dominated by perennial black grama grass along with other common perennial grasses such as spike dropseed (*Sporobolus cryptandrus*) and threeawn (*Aristida purpurea*), widely scattered honey mesquite and broom snakeweed (*Gutierrezia sarothrae*), soaptree yucca (*Yucca elata*), and many species of annual and perennial forbs. The soils consisted of sandy loams.

**Sampling Design**

Data on rabbits, vegetation, and rainfall were collected over a 10-year period from 1996 to 2005. Rabbit species composition and densities were estimated from road-transect surveys (Buckland et al. 1993) at each study site, once during the middle of spring (April/May) and once during fall (October/November) to correspond with spring and fall plant measurements. Over the 10-year study period, the seasonal surveys resulted in 20 samples from each of the 2 study sites. The survey roads were remote, secondary gravel roads on the Chihuahuan Desert Rangeland Research Center (New Mexico State University) and the Jornada Experimental Range (USDA–ARS), with no nighttime automobile traffic to confound observations. These were the only roads available to survey for each study area. One 10-km road transect was sampled through each creosotebush and grassland habitat. Although each of the two 10-km transects were technically a sample size of one each, both 10-km transects traversed varied landscapes and provided good representations of landscape variation across both the creosotebush shrubland and grassland environments.

A field crew consisting of a driver/data recorder and 2 observers drove slowly in an open-bed pickup truck (approximately 15 km·hour⁻¹) along each road segment, starting about one hour after sunset. The vehicle’s trip odometer was set to zero at the start of each road transect. The 2 observers in the back of the truck scanned the side of the road for rabbits, each using a 1,000,000-candlepower spotlight. The observers shined the light beam in a 90° arc from the road ahead to the area perpendicular to the truck and up to 100 m out from the road. Each time a rabbit was observed, the driver recorded the odometer mileage and time, and the rabbit was identified to species. The perpendicular distance from the center...
Fig. 2. Photographs of the Jornada Basin: (A) black grama grassland and (B) creosotebush shrubland study sites.
of the road to the rabbit was measured to the nearest meter with a metric infrared-laser range finder. Only observations within 100 m of the road were recorded to avoid potential errors associated with long-distance observations. Smith (1990) found black-tailed jackrabbits to be most active on nights with a moon, and we (McGlone and Lightfoot, unpublished data) found that rabbit activity in the Jornada Basin was significantly greater on nights with a full moon. In order to help standardize environmental conditions at sampling times, we sampled only on full-moon nights for this study. We also sampled only when skies were mostly clear and winds were light (<10 mph).

Precipitation was measured from LTER meteorological stations associated with ANPP plots located adjacent to each study site. Rainfall data were collected monthly at each of the sites by using tipping-bucket rain gauges. For this study, the monthly rainfall amounts from November to April were combined to represent the winter–spring rainfall period, and the monthly rainfall amounts from May to October were combined to represent the summer rainfall period. Over the 10-year study period, these seasonal time intervals resulted in a total of 20 precipitation samples from each of the 2 study sites.

ANPP was measured from the Jornada Basin LTER study plots, and biomass values were calculated from volumetric vegetation measurements as described by Huenneke et al. (2001). Three replicate vegetation measurement plots measuring 70 m on each side were established in each vegetation community type by the Jornada Basin LTER program. Each vegetation plot consisted of 21 permanently marked 1-m² quadrats. Vegetation was measured from each 1-m² quadrat according to methods of Huenneke et al. (2001), which consisted of volumetric measurements (vertical height and horizontal cover) of foliage canopy by species on each quadrat. Measurements were taken in January, April, and September of each year to capture seasonal plant growth. The same range of volumes of plant species foliage as measured on the study plots was harvested adjacent to, but not on, the study plots. Harvested plant biomass samples were oven dried, and regression equations of dry weight by volume were developed to relate the harvested plant biomass back to the volumetric values measured on the study plots. This procedure yielded biomass production, or ANPP (g · m⁻²) estimates for the study plots by plant species for each season or each year. The plant species were then categorized into functional groups by growth form (grasses, forbs, or shrubs) and by life history (annuals or perennials). ANPP was determined for the winter–spring period (November–April) and for the summer period (May–October). Over the 10-year study period, those 2 seasonal measurements per year resulted in a total of 20 samples for ANPP from each of the 2 study sites. ANPP data generated from the 2 years prior to the rabbit surveys were also used to evaluate lag responses to productivity of previous years.

Data Analyses

The rabbit count and perpendicular distance measures were used to estimate densities (number of individuals per hectare) for each of the 20 survey dates with the analytical program DISTANCE (Buckland et al. 1993). To avoid observer error between the grassland and shrubland habitats, due to shrubs affecting the abilities of observers to detect rabbits, we limited our data analysis to all observations ≤20 m from the road for both habitats.

We partitioned the rainfall and the ANPP data into 4 different time periods to examine lag-time relationships between previous rainfall and plant production and lag-time relationships between previous plant production and rabbit densities. We examined data at intervals from the previous 0–6 months, 6–12 months, 12–18 months, and 18–24 months. Over the 10-year study period with 2 seasons of data (spring and fall) per year, there were a total of 80 time intervals for lag and nonlag response times.

We used the Shapiro–Wilk test (SAS Institute, Inc. 2005; Proc Univariate) to test all data sets for normal distributions. The precipitation and ANPP data were normally distributed, so we used standard parametric linear regression (SAS Institute, Inc. 2005; Proc Reg) to test for significant relationships between rainfall (independent variable) and plant production (dependent variable). We examined relationships between the total ANPP summed over all plant species and the ANPP partitioned by plant life-forms (e.g., grasses, forbs, etc.) and plant life histories (annual or perennial). The rabbit density data were not normally distributed, so we used a nonparametric Spearman-rank correlation analysis (SAS Institute, Inc. 2005) to test for simple linear relationships between rainfall, ANPP, and rabbit density.
densities over the 10-year period. We used an alpha level of 0.05 for rejecting the null hypotheses of no significant regression or correlation relationships.

Results

Seasonal precipitation and ANPP varied considerably over the study period, but temporal patterns were similar for both study sites (Figs. 3A, 4A). Several ENSO events occurred over the duration of the study. The study began during a dry period in 1995, which was followed by wet periods in 1998, 1999, and 2005, with additional dry periods in 1997 and 2000–2003. Years 1998, 2004, and 2005 were El Niño periods, with above-average precipitation, and 1995, 1996, and 2000–2003 were La Niña years, with below-average winter precipitation. Summer rainfall relative to winter rainfall was particularly high during those years. Total ANPP at the grassland site was significantly positively predicted by the previous 6–12 months of rainfall (\(R^2 = 0.24, F_{19} = 5.68, P = 0.03\)), but not by the
previous 0–6 months of rainfall. Among the various plant life-forms and life histories, only perennial grass production was significantly predicted by rainfall at the grassland site ($R^2 = 0.19$, $F_{19} = 4.19$, $P = 0.05$), and only by the previous 6–12 months of rainfall. Total ANPP at the shrub site was significantly positively predicted by the previous 6–12 months of rainfall ($R^2 = 0.28$, $F_{19} = 7.19$, $P = 0.01$), and perennial shrubs ($R^2 = 0.34$, $F_{19} = 9.28$, $P = 0.007$) and perennial forbs ($R^2 = 0.19$, $F_{19} = 4.28$, $P = 0.05$) were significantly predicted by rainfall from the previous 6–12 months. Annual grass was the only vegetation at the shrubland site significantly predicted by rainfall from 0–6 months previous ($R^2 = 0.27$, $F_{19} = 6.62$, $P = 0.02$). We did not find any significant relationships between rainfall and ANPP beyond 12 months.

Both black-tailed jackrabbits and desert cottontails were generally more abundant at the grassland site than at the shrubland site, and black-tailed jackrabbits were consistently more abundant than desert cottontails (Figs. 3B, 4B).
Rabbit densities also changed over time at both study sites, but these changes occurred slowly over many years, with little annual fluctuation (Figs. 3B, 4B). Densities were consistently higher at the grassland site, where they peaked in 1999 near 1.0 rabbit \( \cdot \) ha\(^{-1}\), then gradually declined to about 0.3 rabbits \( \cdot \) ha\(^{-1}\) by 2005. Initial rabbit densities at the shrub site were near 0.3 \( \cdot \) ha\(^{-1}\) in 1995 and gradually declined to near 0.1 \( \cdot \) ha\(^{-1}\) in 2005. Total combined densities of both rabbit species and those for each rabbit species separately were not significantly \((P > 0.05)\) correlated with rainfall from the previous 6 months, 12 months, or >12 months from either the grassland or shrubland sites. Total rabbit densities and densities for each species were not significantly correlated with plant production from the previous 6 months or >6 months at either study site.

**DISCUSSION**

Over the 10-year study period, our data demonstrated that densities of both rabbit species changed gradually over time and that their densities were not positively correlated with short-term, bottom-up annual variation in either rainfall or plant production. This finding contrasts with the documented relationships between rainfall and Chihuahuan Desert rodent abundance (Whitford and Steinberger 1989, Ernest et al. 2000). Chihuahuan Desert rodent assemblages are dominated by granivorous heteromyids that forage largely on the seeds of annual herbs, which are greatly influenced by annual precipitation (Beatley 1969, 1974, Reichman 1975, Brown and Zeng 1990, Whitford and Steinberger 1989, Brown and Heske 1990). Black-tailed jackrabbits, and probably desert cottontails, forage on annual herbs as well as on woody perennial shrubs and are known to switch from foraging on herbs to woody shrubs during dry periods (Steinberger and Whitford 1983, Lopez-Portillo 1996, Martinez and Lopez-Portillo 2003). Woody shrubs provide temporally stable food and moisture resources regardless of short-term responses to variation in rainfall as predicted by the pulse-reserve model. Therefore, Chihuahuan Desert rabbits may not be as influenced by short-term, bottom-up annual plant production in the way rodents are. These results provide some of the first insights into relationships between rabbit abundance and bottom-up influences in North America.

Portales et al. (2004) demonstrated that the reproductive condition of the black-tailed jackrabbit was positively correlated with short-term rainfall and other environmental factors in the Mapimi Basin in the central Chihuahuan Desert. Rainfall and the immediate subsequent production of plant forage may be important to rabbit reproduction, as it is for desert rodents (Beatley 1969, 1974, Reichman 1975, Reichman and Van De Graff 1975), but other factors may be more important in determining the ultimate annual population levels in the Chihuahuan Desert. Predation and/or disease probably complicated the positive relationship between rainfall, plant production, and rabbit density increases. Predation has been documented as a factor in declining black-tailed jackrabbit densities (Clark 1972, Wagner and Stoddart 1972, Bartel and Knowlton 2005), and diseases, such as tularemia and myxomatosis, are known to negatively affect rabbit densities (Dwyer et al. 1990, Reddiex et al. 2002). However, we had no data on important predators or disease organisms from our study area. Still, the fact that densities of both rabbit species changed gradually over time indicates that some other factor(s), operating over time scales of more than one year were influencing the rabbit abundance. Both species of rabbits exhibited overall declines in density over the 10-year duration of the study.

Foraging behavior of black-tailed jackrabbits appears to be confounded by predation risk. McAdoo et al. (1987), working in Nevada, found black-tailed jackrabbit densities and foraging to be greatest near shrubs and less in large open grassland habitats with similar amounts of food. Ariadna et al. (2003) found higher densities of black-tailed jackrabbits in habitats with both greater food resources and lower predation risk. These studies demonstrate that top-down effects of predators can offset the positive effects of increased food availability. We observed rabbit predators such as coyotes, bobcats, hawks, Golden Eagles, and Great-horned Owls at our study site in the Jornada Basin. All of these predators likely affected the rabbit populations that we studied.

We also found that black-tailed jackrabbits and desert cottontails were more abundant in the relatively open grassland than in the adjacent shrubland. Black-tailed jackrabbits were more numerous than desert cottontail rabbits across both habitats. Black-tailed jackrabbits are known to prefer relatively open habitats with...
some shrub cover (Desmond 2004), presumably so they can visually detect and escape from predators (McAdoo et al. 1987). Desert cottontail rabbits prefer habitats with considerable shrub or low tree cover (Schmidly 1974) in which they presumably hide from predators; yet, we found higher densities of desert cottontail rabbits in the grassland than in the shrubland. The relatively open grama grasslands included scattered mesquite shrubs and soap-tree yucca plants which apparently provided adequate cover for both species of rabbits.

Our findings demonstrate that black-tailed jackrabbit and desert cottontail rabbit abundance did change over time, but the changes were long-term, multiyear trends rather than short-term, annual fluctuations. Our findings also demonstrate that rabbit densities were not positively correlated with either rainfall or plant production. Our 10-year sampling period may not have been long enough to provide sufficient data to detect subtle relationships between plant production and rabbit densities or to assess the importance of long-term patterns in rainfall and plant production. Other environmental factors such as predators and disease probably affected Chihuahuan Desert rabbit abundance. More long-term research is needed to understand the relationships between rabbits, climate, weather, plant production, predators, and disease.

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