Are ecosystems structured from the top-down or bottom-up: a new look at an old debate

Charles E. Kay

Abstract Whether ecosystems are structured from the top-down (i.e., predator driven) or bottom-up (i.e., food limited) has been debated by ecologists for nearly a century. Many marine and freshwater aquatic systems appear to be under top-down control, but less evidence exists that predators have had a similar effect in terrestrial systems, especially those systems involving large ungulates. Earlier research, however, omitted any serious discussion of Native Americans. Contrary to prevailing beliefs, Native Americans were not conservationists, and they had dramatic impacts on wildlife populations. Native Americans were the ultimate keystone predator and the ultimate keystone species through activities such as aboriginal burning. Moreover, the idea that North America was a "wilderness" untouched by the hand of man prior to 1492 A.D. is incorrect, as recent population estimates indicate that native people may have numbered as many as 100 million, or more, before they were decimated by introduced diseases and other colonial processes. Until the importance of aboriginal land management is recognized and modern management practices change accordingly, our ecosystems will continue to lose the biological diversity and ecological integrity they once had, even in national parks and other protected areas.

Key words aboriginal overkill, aspen, beaver, bison, elk, fire, keystone predation, keystone species, national park management, Native Americans, predation, willows

Estes (1995, 1996) recently discussed whether ecosystems are structured from the top-down, predator driven, or from the bottom-up, food- or resource-limited. This debate has a long history in ecology (Hairston et al. 1960, Hunter and Price 1992), and its solution is critical if biologists are to implement ecosystem management, especially in national parks and other protected areas (Estes 1996). Many marine and freshwater aquatic systems appear to be under top-down control, but there is less evidence that predators have a similar effect in terrestrial systems (Estes 1995, 1996). Earlier studies, however, omitted any serious discussion of Native Americans, who, I suggest, were the ultimate keystone predator (Mills et al. 1993, Power et al. 1996) that structured North American

ecosystems from ca. 12,000 before present (BP) to ca. 1870, especially in the western United States and Canada, where I have conducted most of my research. My views challenge the conventional belief that the impacts of native people on wildlife is unimportant or that they were conservationists (Arcese and Sinclair 1997, Vale 1998). Moreover, I suggest that through practices like aboriginal burning, Native Americans were the ultimate keystone species, creating the very ecosystems that we now consider "natural." I present various lines of evidence and reasoning that support my hypothesis. Because this is a synthesis paper, individual publications should be consulted for the methods that were used in those studies and for the details of their findings.

Author's address: Department of Political Science, Utah State University, Logan, UT 84322, USA,

Evidence historical and pre-Columbian ungulate populations were not food limited

There are several lines of evidence that suggest from ca. 1750 to 1870 ungulate populations were not food limited across the Intermountain West. These include, but are not limited to, first-person historical accounts, photographs from ca. 1870, berry-utilization data, and the abundance of beaver (*Castor canadensis*; Kay and Wagner 1994; Kay and White 1995; Kay 1995b,c, 1996b, 1997a,b,c,d,f,g;). In addition, archaeological data indicate that pre-Columbian ungulate populations were not food limited (Kay 1990, 1994a, 1997e; Kay et al. 1994; Allen 1996; Truett 1996).

First-person historical accounts

During the early 1990s, there were an estimated 100,000 clk (Cervus elaphus) in the Yellowstone ecosystem and an estimated 4,000 bison (Bison bison) in Yellowstone National Park (Harting and Glick 1994). According to the National Park Service, these large ungulate populations are assumed to be "natural" and to represent the "pristine" state of the ecosystem (Houston 1982, Despain et al. 1986). If that were true, then early explorers should have reported an abundance of game. Between 1835 and 1876, 20 different expeditions spent 765 days in the Yellowstone Ecosystem, yet they reported seeing elk only once every 18 party-days, and bison were seen on only 3 occasions, none of which were in Yellowstone Park itself (Kay 1990). In addition, no one reported seeing or killing even a single wolf (Cants lupus), another indication that game was scarce (Kay 1995b). Moreover, while the explorers were in Yellowstone, their journals contain 45 references to a lack of game or a shortage of food (Kay 1990). Recently, Schullery and Whittesey (1992) published an exhaustive compendium of early wildlife observations in Yellowstone, but their per-party sighting rates were actually only half those reported in firstperson journal accounts (Kay 1990).

Similarly, elk are now the most abundant ungulate in Banff National Park and other parts of the Canadian Rockies. Between 1792 and 1872, however, 26 different expeditions spent 369 days traveling through the mountains on foot or horseback yet reported seeing elk on only 12 occasions, or once every 31 partydays (Kay and White 1995). Thus, contrary to popular perception, first-person historical accounts provide no evidence that ungulates were once common in the Intermountain West (Koch 1941, Rawley 1985, Davis 1986, Allen 1996, Truett 1996).

Historical photographs

Historical photographs can also be used to judge the number of ungulates that occupied areas in the past and to determine whether those animals were food limited. If elk, for example, were as abundant historically as they are today in various western national parks (Hess 1993, Wagner et al. 1995, Allen 1996), then favored forage species, like aspen (*Populus tremuloides*) and willows (*Sallx* spp.), should show that those communities were as heavily browsed during the 1800s as they are at present (Kay 1990, Kay et al. 1994, Kay and White 1995). If aspen and willows depicted in historical images do not show evidence of repeated browsing, that would indicate that fewer ungulates used the range in the past and that factors other than food limited those herbivores.

Ungulates in Yellowstone National Park have reportedly been food limited for many years under what is termed "natural-regulation" management (Houston 1982, Despain et al. 1986). According to this paradigm, predation is an assisting but nonessential adjunct to the regulation of ungulate populations by food limitation. If wolves are present, they will only take the ungulates slated to die from other causes, such as starvation, and, hence, predation will not cause population declines of ungulates (Houston 1982, Despain et al. 1986). In the current debate over reintroduction of wolves to Yellowstone, the Park Service has denied that wolves are needed to control the park's elk herds or that wolves will have any significant impact on the number of ungulates (Boyce 1992, Kay 1996a). Instead, the Park Service believes that it is natural for thousands of elk and other ungulates to starve to death during winter. The agency contends that those animals have always heavily impacted the vegetation, including high-lining conifers, which is now widespread (Houston 1982, Despain et al. 1986, Kay 1990).

Historical photographs, however, show no evidence of ungulate browsing (Kay and Wagner 1994; Fig. 1). In addition, photographs taken over time of tall willows (n = 44) and aspen (n = 81) indicate that the areas occupied by those species have declined 95% since the late 1800s due to repeated ungulate browsing, not other factors (Chadde and Kay 1988, 1991; Kay and Chadde 1992; Kay 1994b, 1995c, 1996b, 1997d,g; Kay and Walker 1997). Thus, ungulate high-lining of conifers and repeated browsing of other woody vegetation represent a departure from conditions that existed prior to the establishment of Yellowstone National Park (Kay and Wagner 1994). Moreover, because conifers and other woody species



Fig. 1. Unbrowsed condition of aspen in Yellowstone Park during the late 1800s. The aspen in this and other early photographs show no evidence of ungulate browsing. Repeated browsing has now eliminated the aspen shown here. In the foreground, Company D of the Minnesota National Guard is on patrol—the military administered Yellowstone from 1886 until 1916, when the National Park Service was created. Photo (ca. 1893) by F. J. Haynes (H-3069), courtesy of the Montana Historical Society, Helena, Montana.

depicted in early images were 50–100 years old when they were photographed and because they show no evidence of ungulate use, this would suggest that few, if any, elk wintered in Yellowstone from the late 1700s through the 1870s (Kay and Wagner 1994).

Historical photographs taken in the Canadian Rockies show the same pattern (Kay et al. 1994; Kay and White 1995), as do photos taken ca. 1870 throughout the West (Kay 1997a; historical and repeat-photo studies in progress for the Agric. Res. Scrv., U.S. For. Serv., and Ut. Div. of Wildl. Resour.; Kay and Walker 1997). Early photographs show no evidence of ungulate browsing, the exact opposite of conditions today, especially in national parks and preserves where ungulate populations are food limited.

Use of berries

Berry production and utilization data also suggest that historical ungulate populations were low. Ethnographic accounts and archaeological studies reveal that Native Americans routinely consumed large quantities of berries, such as serviceberries (*Amelanchier alnifolia*) and chokecherries (*Prunus virginiana*; Lowie 1909, Chamerlin 1911). In September 1869, for instance, the Cook-Folsom-Peterson Expedition met Native Americans who were gathering and drying large quantities of chokecherries at the mouth of Tom Miner Creek just north of Yellowstone Park (Haines 1965). The Washburn Expedition of 1870 reported that near Yellowstone Park "we crossed a small stream

bordered with black cherry trees [chokecherries], many of the smaller ones broken down by bears, of which animal we found many signs" (Langford 1972:13). Because shrubs have to be ≥2 m in height before branches are commonly broken down by feeding bears (*Ursus* spp.), Yellowstone's chokecherry plants in 1870 not only produced abundant berries, but were apparently rather robust, tall shrubs.

Conditions today are quite different. Serviceberry and chokecherry plants in Yellowstone now average <50 cm tall and produce virtually no berries because they are repeatedly browsed by elk and other resource-limited ungulates (Kay 1990, 1995c). At the Lamar-West exclosure on Yellowstone's northern range, 100 protected serviceberry plants produced 111,047 berries, while 100 browsed plants produced no berries, and 100 protected chokecherry plants produced 212,178 berries, while 100 browsed plants produced none (Kay 1995a,c, 1997d).

Resource-limited ungulate populations and large quantities of berries are mutually exclusive on western ranges. Even moderate numbers of ungulates curtail berry production because those shrubs provide highly preferred forage, especially during winter (Kay 1995c). Ungulate-induced berry reduction is even reflected in grizzly bear (*Ursus arctos*) diets. While bears in other ecosystems commonly consume large quantities of berries (Le Franc et al. 1987), grizzlies in Yellowstone cat virtually none (Mattson et al. 1991; Kay 1995c, 1997d, f). In historic and prehistoric times, peoples in the West consumed large quantities of berries; this suggests that ungulate populations were small and that those animals were not limited by food (Kay 1994a, 1995a,c, 1996b).

The abundance of beaver

Beaver also provide evidence that historical ungulate populations were not food limited. Millions of beaver inhabited western North America prior to the fur trade (Johnson and Chance 1974, Kay 1994b). Beaver commonly inhabited mountain streams, but large numbers were also found along water courses on the Canadian and United States prairies, especially in Canada's aspen parklands. One Hudson Bay Company fur brigade, for instance, caught 511 beaver from a small northern Utah drainage in just 5 days (Kay 1994b). To support these large numbers of beaver, woody vegetation that beaver need for food and dam building materials, like aspen, willows, and cottonwoods (Populus spp.), must have been plentiful. Moreover, those plants could not have been subjected to repeated browsing by large numbers of resourcelimited ungulates, because those species are quickly eliminated by high levels of herbivory (Keigley 1997).

Yellowstone provides an excellent example of the impact food limited ungulates have on beaver populations. During the early 1800s, Russell (1965) spent weeks trapping beaver on what is now the park's horthern range. Even after Yellowstone was established as the world's first national park in 1872, there were hundreds, if not thousands, of beaver on the northern range (Kay 1990, 1997d). Today, however, beaver are ecologically extinct on Yellowstone's northern range because repeated browsing by the park's resource-limited ungulates has eliminated the tall willows and aspen beaver need for food (Chadde and Kay 1988, 1991; Kay and Chadde 1992; Kay 1997e; Kay and Walker 1997). Ungulate populations have also had a negative impact on beaver in Rocky Mountain National Park (Hess 1993) and in Banff National Park (Flook 1964). Thus, if large numbers of beaver were once common in the past; then ungulates must have been limited by factors other than food.

Archaeological data

Archaeological evidence indicates that pre-Columbian ungulate populations were also not resource limited. In food limited intermountain systems, large ungulates such as elk, bison, and moose (*Alces alces*) competitively exclude smaller, less efficient herbivores like deer (*Odocoileus bemionus* and *O. virginianus*) and bighorn sheep (*Ovis canadensis*; Cliff 1939, Cowan 1947, Flook 1964, Olmsted 1979, Parker et al. 1984, Telfer and Kelsal 1984). If pre-Columbian intermountain ungulate populations were food limited and native people randomly harvested those animals, then archaeologically recovered ungulate faunal remains should be dominated by elk, bison, and moose, whereas deer and bighorn sheep should be less abundant (Kay 1990, 1994a). The opposite pattern, however, has been observed.

Of nearly 60,000 ungulate bones unearthed at >400 archaeological sites in the United States and Canadian Rockies, <3% were elk, and only about 10% were bison (Kay 1990, 1994a, 1997e; Kay et al. 1994; Kay and White 1995). Only I moose bone has been recovered in the western United States, an area now inhabited by an estimated 25,000 moose (Kay 1997e). Instead, at intermountain archaeological sites in the southern Canadian and United States Rockies, deer and bighorn sheep are the most frequently recovered ungulates (Kay 1990, 1994a, 1995a, 1997e; Kay et al. 1994; Kay and White 1995). Even in the Greater Yellowstone Ecosystem, where elk presently constitute around 80% of the total ungulate fauna, elk are rare to nonexistent in archaeological sites (Wright 1984; Kay 1990, 1992, 1994a). Elk currently dominate ungulate communities in Idaho's River of No Return Wilderness and in Oregon's Blue Mountains, but few elk bones have ever been recovered from archaeological sites in those areas (Kay 1990). Similarly, elk now dominate the ungulate fauna in Banff, Jasper, Yoho, and Kootenay national parks, but elk is among the least frequent ungulate species recovered from archaeological remains (Kay et al. 1994, Kay and White 1995, Kay 1997c). Similar situations occur in other western states, including Arizona and New Mexico (Allen 1996, Truett 1996). Bandelier National Monument, for instance, now supports high densities of largely resource-limited elk, yet only 9 elk bones have been recovered from 45 archaeological sites (Allen 1996).

Many ecologists harbor a mistaken belief that aboriginal diets were primarily meat (McCabe and Mc-Cabe 1984, Reeves and McCabe 1997). Anthropologists, though, have long argued that native people should more appropriately be called gatherer-hunters, instead of hunter-gatherers, because ≥90% of most historic and prehistoric diets were non-ungulate foods (Lee and DeVore 1968, Sahlins 1972). This was especially true in the Intermountain West (Hunn and French 1981, Wright 1984). According to optimal-foraging models, plant foods, small mammals, and fish are lower ranked diet items (i.e., they provide lower caloric return rates per unit time) than ungulates (Smith 1983, Simms 1984, Smith and Winterhalder 1992), which implies that, historically and prehistorically, ungulate population levels were low; i.e., those animals were not food limited (Kay 1990, 1994a, 1997e). Optimal-foraging theory (Stephens and Krebs 1986) predicts that ungulates will be taken by aboriginal peoples whenever those animals are encountered; a diet of low-ranked items, such as that experienced by Native Americans in the West for >10,000 years, indicates that high-ranked ungulates were rare or absent (Smith 1983; Simms 1984; Broughton 1994a, 1994b, 1995, 1997; Janetski 1997).

As noted above, elk in Yellowstone National Park are food limited and now winter at densities of 20-40 elk/km² (Houston 1982, Singer and Norland 1994, Singer et al. 1997). If this was true in earlier times, optimal-foraging models predict that aboriginal diets should have been nearly 100% elk, but archaeological finds indicate that was not the case (Kay 1990). Thus, it is likely that few elk were actually available to prehistoric hunters and that today's ungulate populations are not representative of pre-Columbian conditions (Kay 1994a, 1997e). This was true in Yellowstone, and throughout the Intermountain West (Kay 1994a, 1997e).

Furthermore, the condition of archaeologically recovered faunal remains supports the interpretation that pre-Columbian ungulate populations were low (Broughton 1994*a*,*b*, 1995; Potter 1995). Most bone recovered

from intermountain archaeological sites is highly fragmented due to processing by the native peoples who left those cultural deposits (Kay 1990); i.e., Native Americans broke-up the bones and then extracted the grease via boiling (Lecchman 1951; Vehick 1977; Binford 1978, 1981). Bone-grease processing, however, is labor intensive and may have been done at a net loss of energy (Kay et al. 1994). Thus, native peoples may have been experiencing nutritional stress, or at least a shortage of critical animal fats (Binford 1978, 1981; Olson 1983; Schalk and Micrendorf 1983), which, in turn, indicates that ungulates were not abundant (Broughton 1994a, b, 1995; Potter 1995). That bone-grease processing was the norm throughout much of western North America suggests that ungulate populations were being kept at low levels by factors other than food limitation (Kay et al. 1994, Kay and White 1995).

Aboriginal overkill

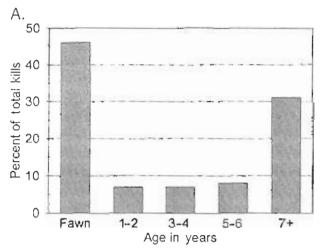
Carnivore predation (Estes 1995, 1996; Kay 1996a) and native hunting are factors that may once have limited ungulate numbers. The presence of aboriginal buffer zones, however, indicates that predation by wolves and other carnivores was not the primary factor limiting pre-Columbian ungulate populations. Hickerson (1965:45) noted that "Warfare between members of the two tribes had the effect of preventing hunters from occupying the best game region intensively enough to deplete the [deer] supply... In the one instance, in which a lengthy truce was maintained between certain Chippewas and Sioux, the buffer, in effect a protective zone for the deer, was destroyed and famine ensued." Lewis and Clark (1893:1179) noted that "With regard to game in general, we observe that the greatest quantities of wild animals are usually found in the country lying between two nations at war." Similarly, Palliser (1969:266-267) reported that game on the Canadian prairies was more abundant in aboriginal buffer zones: "... I must admit, we ran some risk of being surprised by an Indian war-party... As a general rule, the more dangerous the country the greater the probability of finding [an] abundance of game, showing in more ways than one the truth of the old sportsmen's adage, the more danger the more the sport. This part of the country is so evidently the line of direction [demarcation] between the three hostile tribes, that none of their dare venture into it for hunting, except when driven to desperation by hunger ... Much therefore as I enjoyed the [present] locality for a hunting camp, seeing buffalo on all sides, elk feeding in the distance, and fresh deer tracks in every direction ... Boucharville [my guide) did not relish it at all, and began already to calculate how soon we were to go away."

Thus, historical sources indicate that aboriginal hunting tended to extirpate or to drive-out game animals, and resource depletion around camps and villages has frequently been reported in studies of modem hunter-gatherers (Smith and Winterhalder 1992; Kay 1994a). This pattern would be expected if people pursued an optimal-foraging strategy with no effective conservation practices. Tribal territory boundary zones also explain how early explorers could encounter an abundance of game in a few locations and lack of game elsewhere (Steffian 1991, Hammett 1992). Many aboriginal buffer zones were ≥200 km wide. During the 1800s, for instance, all the land between the Yellowstone and Missouri rivers on the Montana prairies was an aboriginal buffer zone, where game was relatively more abundant (but not food limited), and native use was low due to constant warfare between the Blackfoot Confederation (5 tribes); the Shoshone; the Crow; the Salish, Flatheads, and their Kootenay allies; and the Sioux and their Cheyenne allies (C. E. Kay, unpubl. data). Similarly, West (1995) concluded that bison would not have survived on the central Great Plains without aboriginal buffer zones.

In addition, the age of their respective kills indicates that Native Americans were more efficient predators than wolves. The more difficult it is for a predator to capture a particular prey, the more that predator will take substandard individuals and young (Temple 1987, Kunkel 1997). So, if ≥2 predators are preying upon the same species, the least efficient predator will tend to kill fewer prime-age animals (Okarma 1984). Whereas wolves and other carnivores kill primarily young-ofthe-year and old animals, Native Americans killed mostly prime-age ungulates (Stiner 1990).

Ungulates recovered from intermountain archaeological sites invariably exhibit mortality profiles dominated by prime-age animals (Kay 1994a, 1995a, 1997e), which suggests that Native Americans were more efficient predators than wolves or other carnivores (Stiner 1990; Fig. 2). Killing mostly prime-age animals, however, runs contrary to any maximum sustained-yield strategy (Hastings 1983, 1984) and suggests that Native Americans had a major impact on pre-Columbian ungulate populations, especially when one considers that Native Americans also killed a disproportionate number of females, a preference that runs counter to any conservation strategy (Kay 1994a, 1995a, 1997e).

It is often claimed, however, that Native Americans' religious beliefs prevented those peoples from overutilizing their resources (Speck 1939; Nelson 1982, 1983). Native Americans tended to view wildlife as their spiritual kin where success in the hunt was obtained by following prescribed rituals and



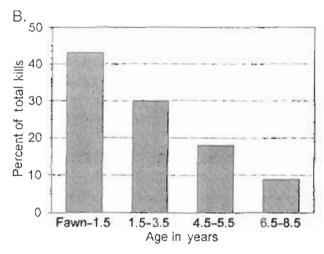


Fig. 2. Age structure of ungulates killed by wolves and Native Americans. (A) Age of white-tailed deer (n=48) killed by wolves in Minnesota (Frits and Mech 1981). (8) Age of mule deer (n=60) uneanhed from the 4,200-year-old Dead Indian Creek archaeological site in northwest Wyoming just east of Yellowstone Park (Simpson 1984). Unlike carnivores, which tend to select prey among young and old animals, Native Americans killed predominantly prime-age ungulates—an indication that Native Americans were more efficient predators. Moreover, the deer at this archaeological site were killed with spears or atlatts, which are less efficient hunting instruments than the bow and arrow that came into use around 1,500 BP (Blitz 1988).

atonement after the kill (Feit 1987). A scarcity of animals or failure in the hunt were not viewed as biological or ecological phenomena, but rather as a spiritual consequence of social events or circumstances (Reeves and McCabe 1997). If a Native American could not find any game, it was not because his people had overharvested the resource, but because he had done something to displease his gods. Because Native Americans saw no connection between their hunting and game numbers, their system of religious beliefs actually fostered the overexploitation of ungulate populations (Kay 1994a, 1995a, 1996b, 1997e).

Native hunters were opportunistic and tended to take high-ranking ungulates, regardless of the size of the prey populations or the likelihood of those animals becoming extinct (Alvard 1993, 1994, 1995; Broughton 1994a, 1994b, 1995, 1997). Native Americans had no concept of maximum sustained yield and did not manage ungulate populations to produce the greatest offtake. In addition, human hunting and predation by carnivores are generally additive and work in concert to reduce ungulate numbers (Walters et al. 1981, Kunkel 1997). Moreover, competition from carnivores would have tended to discourage any possible ungulate conservation practices (Kay 1994a, 1995a, 1996b, 1997e). Because Native Americans could prev-switch between trophic levels to small animals, plant foods, and fish, they could hunt their preferred ungulate prey to low levels or extinction without having an adverse effect on human populations. In fact, once Native Americans killed off most of the ungulates, human populations rose (Hawkes 1991, 1992, 1993).

Although the demonstrated lack of elk and other ungulates in archaeological sites may at first appear to refute the aboriginal overkill hypothesis, the opposite is true. Optimal-foraging theory (Smith 1983, Simms 1984, Smith and Winterhalder 1992, Broughton 1997, Janetski 1997) predicts that highranked items, like elk and other ungulates, are more susceptible to overexploitation than low-ranked items, such as plant foods, small mammals, or fish. According to optimal-foraging models, high-ranked items will seldom appear in the diet if they are being overexploited (Broughton 1994a, 1994b, 1995, 1997). So, ungulate species unearthed with the lowest frequency in archaeological sites, such as moose and elk, were probably subjected to extreme overexploitation (Kay 1994a, 1997e).

Birkedal (1993) reported that Native Americans, armed with no more than spears and hunting dogs, once kept grizzly bear populations at very low levels throughout much of Alaska, and Taulman and Robbins (1996) suggested that native hunters limited the distribution and abundance of the 9-banded armadillo (Dasypus novemcinctus) in the southern United States. Moreover, aboriginal effects were not limited to terrestrial communities. Broughton (1997), for example, found that Native Americans had a significant impact on the population of white sturgeon (Acpenser transmontanus) in San Francisco Bay, and Hewes (1973) and Schalk (1986) suggested that aboriginal fishing had a detrimental effect on the number of salmon in the Columbia Basin. Archaeological information also indicates that native hunters significantly reduced pinniped populations along the California and Oregon coasts (Hildebrandt and Jones 1992, Jones and Hildebrandt 1995) and that native peoples had a significant effect on freshwater and marine shellfish populations (Botkin 1981). Adler (1970) found that the eastern box turtle (Terrapene carolina) was eliminated from many areas by aboriginal peoples.

There are, however, exceptions to aboriginal overkill. According to predator-prey theory, prey populations will increase if they have a refugium where they are safe from predation (Taylor 1984). Therefore, ungulates that could escape aboriginal hunters in time or in space were more abundant. Moreover, refugia do not have to be complete to be effective. Partial refugia also enable prev populations to survive (Taylor 1984). This may explain why larger numbers of ungulates existed on the Great Plains and in the Arctic. By undertaking long-distance migrations, bison and caribou (Rangifer tarandus) were able to outdistance many of their human and carnivorous predators (Bergerud 1990, 1992; Kay 1994a, 1995a, 1996b, 1997e); however, even migratory ungulates historically were not food limited.

Aboriginal populations

Ahoriginal populations were also much larger than commonly believed. Until recently, it was thought that only about 2,000,000 natives inhabited North America prior to the arrival of Columbus (Stannard 1989, 1992). Dobyns (1983), however, postulated that native people, who were attempting to escape Spanish exploitation in Cuba, fled to Florida in oceangoing canoes and brought European-introduced smallpox with them to the mainland during the early 1500s. This and other diseases, to which aboriginal inhabitants had no immunological resistance, then ravaged native people, reducing aboriginal populations by ≥90% before the Pilgrims landed at Plymouth Rock. Subsequently, Ramenofsky (1987), Smith (1987), and Campbell (1990) tested Dobyns' hypothesis using the archaeological record and concluded that ca. 1550–1600 a major collapse of native populations had occurred in North America-100 to 200 years prior to direct contact of Europeans with native people in many areas, especially the Intermountain West; i.e., European diseases were transmitted from native group to native group across all of North America—termed pandemics. Based on this and other evidence, it is now believed that in 1492 there may have been as many as 100 million native people in North America with perhaps an even larger numher in South America (Stannard 1992). Although Dobvos' hypothesis is still debated (Snow 1995), in

general, estimates of pre-European native populations have steadily been revised upward.

There was no "wilderness." In fact, the idea that North America was a "wilderness" untouched by the hand of man before 1492 is a myth, a myth that may have been created, in part, to justify appropriation of aboriginal lands and the genocide that befell native peoples (Denevan 1992, Gomez-Pompa and Kaus 1992, Simms 1992, Cronon 1995). Moreover, there is no evidence that native people ever purposefully limited their populations to avoid environmental impacts or that, if they tried, they were successful (Cohen 1977, 1989; Cohen and Armelagos 1984; Blurton-Jones 1986, 1987; Diamond 1992b; Smith and Winterhalder 1992). As resource use intensified over the last several thousand years, a parallel increase in the violence within and between prehistoric socicties occurred (i.e., people were fighting over scarce resources; Lambert 1997).

The Serengeti myth

Ecologists and wildlife biologists often cite Africa's Screngeti as an example of how North America must have looked before it was despoiled by Europeans (Frank et al. 1998). It has been claimed, for instance, that Yellowstone National Park is the last remnant of North America's Screngeti (Anonymous 1996). Today's Serengeti, however, is not a natural ecosystem, nor is it a vignette of "wilderness" Africa. Instead the Screngeti is a romantic, European, ethnocentric view of how "primitive" Africa should have looked (Adams and McShane 1992); one of the first things that colonial governments did when they created Screngeti and other African national parks was to forcefully remove all of the indigenous peoples. Hominoid predators, however, have existed in Africa for ≥3.5 million years, and it is thought that Homo sapiens evolved in Africa about 100,000 years ago (Shreeve 1995, Tattersall 1995). Thus, I suggest that here is nothing more unnatural than an African ecosystem without hominoid predators; the Serengeti, therefore, is not a "natural" ecosystem nor is it an example of how North America teemed with wildlife before the arrival of Columbus.

In all the ecological studies that have been done on the Serengeti, native people have rarely been mentioned, or if they have, it has usually been in the pejorative sense, as "poachers" (Sinclair and Norton-Griffiths 1979, Sinclair and Arcese 1995). Simulation models have indicated that Serengeti's wildlife populations will collapse if present levels of "poaching" increase by 10% (Sinclair and Arcese 1995:617-637). I would suggest, however, that this may simply be a case of native people exercising their aboriginal rights.

The 60 million bison myth

Similarly, it is often claimed that nearly 60 million bison roamed the North American plains until decimated by advancing European civilization (Roe 1951). That 60-million figure, though, was based on maximum carrying capacity (i.e., food limited) and made no allowance for either the impacts of carnivore predation or native hunting (Roe 1951). Shaw (1995) recently questioned this interpretation, and Geist (1996), concluded that the factor driving bison evolution, ecology, and behavior for the last 12,000 years was hunting by native peoples. Both Shaw (1995) and Geist (1996) revised bison population estimates downward to 10–15 million animals, but I believe that number is still too high (Kay 1996b).

first, buffer zones, berry use, and beaver populations on the prairies all suggest that early wildlife populations were not food limited. Second, widespread burning of the prairies in historical and pre-Columbian times, provides another line of evidence that large numbers of resource-limited bison did not inhabit the plains. Early historical observations provide ample evidence that during the late 1700s and early 1800s, prairie fires often burned for days, and single fires covered huge areas, often running for 100-200 km (Nelson and England 1971, Thomas 1977, Higgins 1986). Large numbers of ungulates and large prairie fires, however, are mutually exclusive because heavy grazing reduces standing plant biomass, prevents the accumulation of plant litter, and creates discontinuous fuel patterns, all of which prevent the growth and spread of large fires (Norton-Griffiths 1979, McNaughton 1992, Hobbs 1996). If there were large fires on the prairies (Nelson and England 1971, Higgins 1986) bison and other ungulates could not have been food limited. Fidler (1990), who traveled with Piegan natives during 1792-1793, reported virtually no unburnt ground on the Canadian prairies from the Oldman River to Buckingham House, a distance of several hundred kilometers. Fidler (1990) observed that most of those fires had either purposefully or accidentally been set by native people. Much of that burning, in fact, occurred during winter when there was no lightning to start fires.

Circumstances were the same throughout the Intermountain West, historically and prehistorically, for many of those plant communities were also once swept by frequent but low-intensity fires (Kay et al. 1994, Kay 1995a, Kay and White 1995). This could not have been true, though, if large numbers of food limited ungulates had been present (Savage and Swetnam 1990; Touchan et al. 1995, 1996; Swetnam and

Baisan 1996). In Yellowstone, for instance, grazing by resource-limited elk and bison has created discontinuous fuel patterns that have changed historical burning patterns and limited the spread of fires (Kay 1997g; Fig. 3). The area along Blacktail Creek was overrun by wildfires in 1988, yet much of the range did not actually burn because ungulate grazing had prevented the accumulation of plant litter necessary to carry fire through these grassland communities. Instead, 100- to 160-km/hour winds drove the fire from patch of fuel to patch of fuel in swales and on north aspects where snow limited forage removal by elk and other wintering ungulates. Historically, Yellowstone's northern range had a fire frequency of once every 25 years (Houston 1973), yet despite the park's "let-burn" policy, virtually none of the northern range has burnt in the last 30 years, except for 1988, but that is thought to be a 100-300 year event (Romme and Despain 1989), and, thus, similar fires could not have created the original fire-return interval. Despain et al. (1986:109) suggested that the park's grasslands have failed to burn during the past 30 years because "lightning has chosen not to strike very often on the northern range," but that hypothesis is not supported by data from the Bureau of Land Management's Automatic Lightning Strike Detection System which shows that, on average, lightning strikes four times per km2 per year (Kay 1990:136-137). Instead, the range does not burn because the park's food limited elk and bison have overgrazed Yellowstone's grasslands, unlike conditions in the past when native hunting limited ungulate numbers.

The ultimate keystone species

Native people were not only the ultimate keystone predator, they were also the ultimate keystone species



Fig. 3. Ungulate-induced burn pattern in Yellowstone National Park. Photo by C. E. Kay.

(Bonnicksen et al. In press). The Americas, as first seen by Europeans, had largely been created by native peoples, not crafted by nature. Native people modified their environments in many ways (Bonnicksen et al. In press), but I will discuss only competition between humans and wildlife for food and aboriginal burning.

Passenger pigeon myths

Neumann (1984, 1985, 1989, 1995) has written extensively about native people, who, by consuming certain foods, limited various wildlife populations. His most interesting example involves the passenger pigeon (Ectopistes migratorius), often cited as an example of how pre-Columbian America teemed with wildlife before Europeans drove that and other species to extinction. But as Neumann (1985) has chronicled, native populations in pre-Columbian times were so large that those people consumed most of the nuts, fruits, and berries that passenger pigeons needed for food. It was only after European diseases decimated Native American populations, and thereby freed the mast crop for wildlife, that the passenger pigeon grew to unprecedented numbers. Thus, the large flocks of passenger pigeons that reportedly darkened the skies during the 1700s and 1800s were an artifact of the "American Holocaust" (Stannard 1992), not an example of how America teemed with wildlife before Europeans arrived.

Aboriginal burning

Native Americans also had a major impact on ecosystems by repeatedly burning the vegetation. They did this to modify plant and animal communities for human benefit and to increase productivity (Pyne 1995). In California, for instance, native peoples had 70 reasons for burning the vegetation (Lewis 1973), and even in northern Canada, where the vegetation is less diverse, Native Americans still set fires for at least 17 different reasons (Lewis and Ferguson 1988). Although aboriginal burning has been widely reported in the anthropological literature (e.g., Lewis 1985; Boyd 1986; Turner 1991; Pyne 1993, 1995; Gottesfeld 1994), those findings have been largely ignored by ecologists (Kay 1995a).

Determining how fires started, though, is critical because "fires set by hunter-gatherers differ from [lightning] fires in terms of seasonality, frequency, intensity, and ignition patterns" (Lewis 1985:75). The majority of aboriginal fires were set in the spring, between snowmelt and vegetation greenup, or late in the fall when burning conditions were not as severe (Pyne 1995). Unlike lightning fires, which tend to be infrequent and of high intensity, native burning produced a high frequency of low-intensity fires. Abo-

riginal burning and lightning fires created different vegetation mosaics, and in many instances, entirely different plant communities (Blackburn and Anderson 1993). Moreover, aboriginal burning reduced or eliminated the number of high intensity, lightning-generated fires (Reid 1987; Pyne 1993, 1995). Once aboriginal fires opened up the vegetation, then subsequent lightning fires behaved like those set by Native Americans (Pyne 1993, 1995).

Pleistocene considerations

Others, however, contend impacts of Native people were insignificant because these were grazing (i.e., food limited) systems during the Pleistocene (Frank et al. 1998), and, thus, intermountain plant communities are preadapted to withstand high levels of large mammal herbivory (Burkhardt 1996). According to this view, there has not been a long enough period of time since the extinction of the Pleistocene megefauna for plants in the West, and the rest of North and South America as well, to adapt to the low levels of ungulate herbivory caused by native hunters (Burkhardt 1996). However, this contention is incorrect for at least 2 reasons. First, 10,000 years is more than enough time for evolution to work, and second, systems during the Pleistocene were predator limited, not food limited, as is usually assumed.

The rate of evolution depends not on time alone, but on the intensity of selection pressure (Weiner 1994). Evolution can occur quite quickly if natural selection is intense. On the Islc of Jersey off the coast of France, for instance, red deer (Cervus elephus) "became reduced to one-sixth of their body weight in less than six thousand years" (Lister 1989:539). On other islands, various poboscideans evolved into forms only 1–2 m tall in <10,000 years (Stuart 1991, 1993). Thus, sufficient time may have passed since the megafauna extinctions for plants to evolve from heavily grazed to lightly grazed forms, especially since heavily grazed plant communities quickly revert to other forms once herbivory is removed (Kay 1990, 1995c, 1997g; Kay and Chadde 1992).

Moreover, Pleistocene animals were termed megafauna for a reason; they were huge. For herbivores to obtain large body size, however, they must have more than adequate forage intake and nutrition (Geist 1971, 1986, 1987a, 1987b). Historically, when herbivores reached islands and predators did not, the herbivores invariably became smaller. The reason red deer lost five-sixth of their "normal" body size on the Isle of Jersey was because they were being limited by food, not predation (Lister 1989). As Geist (1987b:1067) noted, "Island dwarfs appear to

be shaped by efficiency [food] selection in the absence of predation." Thus, if herbivores had been food limited during the Pleistocene, they would not have achieved megafauna size.

Some authors believe that climate change-induced food limitation drove the megafauna to extinction. Others contend that predation, and human predation in particular, caused the megafauna extinctions, not only in the Americas, but around the world (Martin 1967, 1973; Martin and Wright 1967; Martin and Klein 1984; Flannery 1990, 1994; Stuart 1991). Fisher (1996) recently tested the foodextinction and predator-extinction hypotheses by measuring dentinal growth lines in ca. 12,000 BP proboscidean tusks, and found that prehistoric mammoths and mastodons were well fed and had been reproducing at near their maximum theoretical rate, before they went extinct (Fisher 1996, Ward 1997). Not only did native people structure entire ecosystems for the last 12,000 years, but they also may have caused the megafauna extinctions, as well, and, therefore, might be considered the ultimate keystone predator.

Management implications

Whether ecosystems are structured from the topdown or the bottom-up is more than a theoretical debate because it will influence how we manage the Earth's ecosystems, especially in preserves and other protected areas (Diamond 1992a, Estes 1996). Most national parks, wilderness areas, and nature reserves, for instance, are supposedly manged to represent the conditions that existed in pre-Columbian times (i.e., so-called natural or pristine conditions; Arcese and Sinclair 1997). But what is natural? If Native Americans determined the structure of entire plant and animal communities by firing the vegetation and limiting ungulate numbers, among other activities, then that is a completely different situation than we have today (Martinez 1993; Wagner and Kay 1993; Kay 1997a, 1997b). Thus, a handsoff or "natural-regulation" approach by modern land managers will not duplicate the ecological conditions under which those communities developed (Wagner et al. 1995).

Native Americans were not only the ultimate keystone predator but also the ultimate keystone species, whose removal has altered North American ecosystems, even in protected areas. Unless the importance of aboriginal land management is recognized, and modern management practices changed accordingly, our ecosystems will continue to lose the biological diversity and ecological integrity they once had. For, as Aldo Leopold noted >40 years ago, "if we are serious about restoring (or maintaining) ecosystem health and ecological integrity, then we must first know what the land was like to begin with" (Covington and Moore 1994). Native Americans owned, used, and modified nearly all of the New World for ≥12,000 years; to dismiss those people as having had little effect on their environment (Arcese and Sinclair 1997, Vale 1998) is in White's (1995:175) words an act of "immense condescension."

How night these concepts apply to park management? Under the Treaties of 1851 (Kappler 1904:594-596) and 1868 (Kappler 1904:1008-1011), various native people already claim hunting rights in Yellowstone National Park. Thus, I way to reduce overgrazing in that park (Wagner et al. 1995) would be to honor the United States' previous commitment to Yellowstone's original owners and to allow them once again to hunt in the park and surrounding areas (Czech 1995, Kay 1997b). Native people successfully managed Yellowstone and other North American ecosystems for at least the last 10,000 years, and, although they were not conservationists as that term is commonly used, by keeping ungulate populations low, Native Americans promoted biodiversity, which is the hallmark of a keystone predator (Mills et al. 1993, Power et al. 1996). Similarly, aboriginal burning will have to be reinstated if communities are to retain their ecological integrity (Kay and White 1995). And, finally, it should be remembered that allowing nature to take its course under present conditions (Arcese and Sinclair 1997), so called "natural-regulation" or "hands-off" management, is really a value judgment and a decision that has wide-ranging consequences (Wagner et al. 1995), because areas that today are structured from the bottom-up are entirely different from the ecosystems that were historically and prehistorically structured from the top-down (Wagner and Kay 1993).

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Literature cited

- Adams, J. S., and T. O. McShane. 1992. The myth of Wild Africa: Conservation without illusion. W. W. Norton and Company, New York, New York.
- ADIER, K. 1970. The influence of prehistoric man on the distribution of the box turtle. Annals of Carnegie Museum 41:263–280.
- ALLIN, C. D. 1996. Elk response to the La Mesa fire and current status in the Jemez Mountains. Pages 179–195 in C. D. Allen, editor. Fire effects in southwestern forests: Proceedings of the second La Mesa fire symposium. U.S. Forest Service, General Technical Report RM-286.
- ALVARD, M. S. 1993. Testing the "ecologically noble savage" hypothesis: Interspecific prey choice by Piro hunters of Amazonian Peru. Human Ecology 21:355-387.
- ALVARD, M. S. 1994. Conservation by native peoples: Prey choice in a depleted habitat. Human Nature 5:127-154.
- ALMARD, M. S. 1995. Intraspecific prey choice by Amazonian hunters. Current Anthropology 36:789–818.
- Anonymous, 1996. Grazing and Yellowstone. Yellowstone Science 4(1):12-17.
- ARCESE, P., AND A. R. E. SINCLAR. 1997. The role of protected areas as ecological baselines. Journal of Wildlife Management 61:587-602.
- Вевоевир, А. Т. 1990. Rareness as an anti-predator strategy to reduce predation risk. Population Dynamics 1:15–25.
- Bergerup, A.T. 1992. Rareness as an antipredator strategy to reduce predation risk for moose and caribou. Pages 1008-1021 in D. M. McCullough and R. Barrett, editors. Wildlife 2001: Populations. Elsevier Applied Science, New York, New York.
- BINFORD, L. R. 1978. Nunamiut ethnoarchaeology. Academic Press, New York, New York.
- BINFORD, L. R. 1981. Bones: Ancient men and modern myths. Academic Press, New York, New York.
- Birkedal, T. 1993. Ancient hunters in the Alaskan wilderness: Human predators and their role and effect on wildlife populations and the implications for resource management. Pages 228–234 in W. E. Brown and S. D. Veirs, Jr., editors. Partners in stewardship: Proceedings of the seventh Conference on Research and Resource Management in Parks and on Public Lands. The George Wright Society, Hancock, Michigan.
- BLACKBURN, T. C., AND K. ANDERSON, editors. 1993. Before the wilderness: Environmental management by native Californians. Ballena Press, Menlo Park, California.
- Burtz, J. H. 1988. Adoption of the bow in prehistoric North America. North American Archaeologist 9:123-145.
- BLURTON-JONES, N. G. 1986. Bushman birth spacing: A test for optimal interbirth intervals. Ethology and Sociobiology 7:91-105.
- BLUXTON-JONES, N. G. 1987. Bushman birth spacing: Direct tests of some simple predictions. Ethology and Sociobiology 8:183-203.
- BONNEKSEN, T. M., M. K. ANDERSON, H. T. LEWIS, C. E. KAY, AND R. KNUDSON. In press. American Indian influences on the development of North America's native forest ecosystems. Proceedings of the U.S. Forest Service Ecological Stewardship Workshop, 4–16 December 1995, Tueson, Arizona.
- BOTKIN, S. 1981. Effects of human exploitation on shellfish populations at Malibu Creek, California. Pages 121-139 in T. K. Earle and A. L. Christenson, editors. Modeling change in pre-historic subsistence economies. Academic Press, New York, New York.
- Boyce, M. S. 1992. Wolf recovery for Yellowstone National Park:
 A simulation model. Pages 123–138 in D. M. McCullough and
 R. Barrett, editors. Wildlife 2001: Populations. Elsevier Applied Science, New York, New York.

- BOYD, T. 1986. Strategies of Indian burning in the Willamette Valley. Canadian Journal of Anthropology 5:65-86.
- BROCGITTON, J. M. 1994a. Declines in mammalian foraging efficiency during the late Holocene, San Francisco Bay, California. Journal of Anthropological Archaeology 13:371-401.
- BROUGITTON, J. M. 1994b. Late Holocene resource intensification in the Sacramento Valley, California: The vertebrate evidence. Journal of Archaeological Science 21:501-514.
- BROUGITON, J. M. 1995. Resource depression and intensification during the Late Holocene, San Francisco Bay: Evidence from the Emeryville shellmound vertebrate. Dissertation, University of Washington, Seattle.
- BROUGHTON, J. M. 1997. Widening diet breadth, declining foraging efficiency, and prehistoric harvest pressure: lehthyofaunal evidence from the Emeryville Shellmound, California. Antiquity 71:845-862.
- BURKHARDT, J. W. 1996. Herbivory in the Intermountain West: An overview of evolutionary history, historic cultural impacts and lessons from the past. Idaho Forest, Wildlife, and Range Experiment Station, Bulletin 58.
- CAMPBELL, S. K. 1990. Post Columbian cultural history in northern Columbia Plateau A.D. 1500–1900. Garland Publishing, New York, New York.
- CHARDE, S. W., AND C. E. KAY. 1988. Willows and moose: A study of grazing pressure, Slough Creek exclosure, Montana, 1961–1986. University of Montana, Montana Forest and Conservation Experiment Station, Research Note 24.
- Chadde, S. W., and C. E. Kay. 1991. Tall willow communities on Yellowstone's northern range: A test of the "natural regulation" paradigm. Pages 231–264 in R. R. Keiter, and M. S. Boyce, editors. 'The Greater Yellowstone Ecosystem: Redefining American's wilderness heritage. Yale University Press, New Haven, Connecticut.
- CHAMBERLIN, R. V. 1911. The ethno-botany of the Gosiute Indians of Utah. Memoirs of the American Anthropological Association 2:331-405.
- Cliff, E. P. 1939. Relationship between elk and mule deer in the Blue Mountains of Oregon. Transactions of the North American Wildlife Conference 4:560-569.
- COHEN, M. N. 1977. The food crisis in prehistory. Yale University Press, New Haven, Connecticut.
- COHEN, M. N. 1989. Heath and the rise of civilization. Yale University Press, New Haven, Connecticut.
- COHEN, M. N., AND G. J. ARMILAGOS. 1984. Palepathology at the origins of agriculture. Academic Press, New York, New York.
- COVINGTON, W. W., AND M. M. MOORF. 1994. Southwestern ponderosa forest structure: Changes since Euro-American settlement. Journal of Forestry 92:39–47.
- Cowan, I, McT. 1947. Range competition between mule deer and bighorn sheep and elks in Jasper Park, Alberta. Transactions of the North American Wildlife Conference 12:223–227.
- CRONON, W. 1995. The trouble with wilderness: Or, getting back to the wrong nature. Pages 69-90 in W. Cronon, editor. Uncommon ground: Toward reinventing nature. W. W. Norton and Company, New York, New York.
- CZECH, B. 1995. American Indians and wildlife conservation. Wildlife Society Bulletin 23:568–573.
- DAYIS, G. P. 1986. Man and wildlife in Arizona: The American exploration period 1824–1865. Arizona Game and Fish Department, Phoenix.
- DENEVAN, W. 1992. The pristine myth: The landscape of the Americas in 1492. Association of American Geographers Annals 82:369-385.
- DESPAIN, D. G., D. HOUSTON, M. MEAGHER, AND P. SCHULLERY. 1986.
 Wildlife in transition: Man and nature on Yellowstone's northern range. Roberts Rinehart, Inc., Boulder, Colorado.

- Demonp, J. 1992a. Must we shoot deer to save nature? Natural History 104(8):2-8.
- DIAMOND, J. 1992b. The third chimpanzee: The evolution and future of the human animal. Harper Collins Publisher, New York, New York.
- DONYNN, H. F. 1983. Their numbers become thinned: Native American population dynamics in eastern North America. University of Tennessee Press, Knoxville.
- Estris, J. A. 1995. Top-level carnivores and ecosystem effects: Questions and approaches. Pages 151–158 in C. G. Jones and J. H. Lawton, editors. Linking species and ecosystems. Chapman and Hall, New York, New York.
- ESTES, J. A. 1996. Predators and ecosystem management. Wildlife Society Bulletin 24:390–396.
- FEIT, H. A. 1987. North American native hunting and management of moose populations. Swedish Wildlife Research Supplement 1:25-42.
- FIDLER, P. 1990. A look at Peter Fidler's journal: Journal of a journey over land from Buckingham House to the Rocky Mountains in 1792 and 1793. Historical Research Centre, Lethbridge, Alberta.
- FISHER, D. C. 1996. Extinction of proboscideans in North America. Pages 296-315 in J. Shoshani and P. Tussy, editors. The proboscidea: Evolution and palaeoecology of elephants and their relatives. Oxford University Press, New York, New York.
- FLANNERY, T. F. 1990. Pleistocene faunal loss: Implications of the aftershock for Australia's past and future. Archaeology in Oceania 25:45-67.
- FLANNERY, T. F. 1994. The future caters: An ecological history of the Austrulasian islands and people. Reed Books, Chaswood, New South Wales, Australia.
- FLOOK, D. R. 1964. Range relationships of some ungulates native to Banff and Jasper National Parks, Alberta. Pages 119-128 in D. J. Crisp, editor. Grazing in terrestrial and marine environments. Blackwell Press, Oxford, United Kingdom.
- FRITTS, S. H., AND L. D. MECH. 1981. Dynamics, movements, and feeding ecology of a newly protected world population in northwestern Minnesota. Wildlife Monographs 80.
- FRANK, D. A., S. J. McNaugiffon, and B. F. Tracy. 1998. The ecology of the Earth's grazing ecosystems. BioScience 48:513–521.
- Gust, V. 1971. Mountain sheep. University of Chicago Press, Chicago, Illinois.
- GEST, V. 1986. The paradox of the great Irish stags. Natural History 95(3):54-64.
- Gust, V. 1987a. On the evolution and adaptations of Alces. Swedish Wildlife Research Supplement 1:11-23.
- Gust, V. 1987b. On speciation in Ice Age mammals, with special reference to cervida and caprids. Canadian Journal of Zoology 65:1067–1084.
- Geist, V. 1996. Buffalo nation: History and legend of the North American bison. Voyageur Press, Stillwater, Michigan.
- GOMEZ-POMPA, A., AND A. KAUS. 1992. Taming the wilderness myth. Bioscience 42:271–279.
- GOTTESFEID, L. M. J. 1994. Aboriginal burning for vegetative management in northwest British Columbia. Human Ecology 22:171–188.
- Haines, A. L. 1965. Valley of the upper Yellowstone. University of Oklahoma Press, Norman.
- HAIRSTON, N. G., F. E. SMITH, AND L. B. SLOBODKIN. 1960. Community structure, population control and competition. American Naturalist. 94:421–425.
- HAMMETT, J. E. 1992. The shapes of adaptation: Historical ecology of anthropogenic landscapes in the southeastern United States. Landscape Ecology 7:121-135.
- HARTING, A., AND D. GUCK. 1994. Sustaining greater Yellowstone,

- a blueprint for the future. Greater Yellowstone Coalition, Bozeman, Montana.
- Hastings, A. 1983. Age-dependent predation is not a simple process. I. Continuous time models. Theoretical Population Biology 23:347-362.
- HASTINGS, A. 1984. Age-dependent predation is not a simple process. II. Wolves, ungulates, and a discrete time model for predation on juveniles with a stabilizing tail. Theoretical Population Biology 26:271–282.
- HAWKEN, K. 1991. Showing off: Tests of a hypothesis about men's foraging goals. Ethology and Sociobiology 12:29-54.
- HAWKES, K. 1992. On sharing and work. Current Anthropology 33:404-407.
- HAWKES, K. 1993. Why hunter-gatherers work. Current Anthropology 34:341–361.
- HESS, K., Jr. 1993. Rocky times in Rocky Mountain National Park: An unnatural history. University Press of Colorado, Niwot.
- HEWTS, G. 1973. Indian Fisheries productivity in pre-contact times in the Pacific salmon area. Northwest Anthropological Research Notes 7:133-155.
- HICKERSON, H. 1965. The Virginia deer and intertribal buffer zones in the upper Mississippi Valley. Pages 43–65 in A. Leeds and A. P. Vayda, editors. Man, culture and animals: The role of animals in human ecological adjustments. American Association for the Advancement of Science 78.
- Higgins, K. F. 1986. Interpretation and compendium of historical fire accounts in the northern Great Plains. U.S. Fish and Wildlife Service, Resource Publication 161.
- HILDEBRANDT, W. R., AND T. L. JONES. 1992. Evolution of marine mammal hunting: A view from the California and Oregon Coasts. Journal of Anthropological Archaeology 11:360–401.
- Honns, N. T. 1996. Modification of ecosystems by ungulates. Journal of Wildlife Management 60:695-713.
- Housron, D. B. 1973. Wild fires in northern Yellowstone National Park. Ecology 54:1111-1117.
- HOUSTON, D.B. 1982. The northern Yellowstone elk: Ecology and management. MacMillan Publications, New York, New York.
- HUNN, E. S., AND D. H. FRENCH. 1981. Lomatium: A key resource for Columbia Plateau nature subsistence. Northwest Science 55:87-94.
- HUNTER, M. D., AND P. W. PRICE. 1992. Playing chutes and ladders: Bottom-up or top-down forces in natural communities. Ecology 72:724-732.
- JANETSKI, J. C. 1997. Fremont hunting and resource intensification in the eastern Great Basin. Journal of Archaeological Science 24:1075–1088.
- JOHNSON, D. R., and P. H. CHANGE. 1974. Presettlement over harvest of upper Columbia River beaver populations. Canadian Journal of Zoology 52:1519–1521.
- JONES, T. L., AND W. R. HILDERMANDE. 1995. Reasserting a prehistoric tragedy of the commons: Reply to Lyman. Journal of Anthropological Archaeology 14:78–98.
- KAPPLER, C. J., editor. 1904. Indian affairs: Laws and treaties. Volume, II—Treaties. Government Printing Office, Washington D.C.
- KAY, C. E. 1990. Yellowstone's northern elk herd: A critical evaluation of the "natural regulation" paradigm. Dissertation, Utah State University, Logan.
- KAY, C. E. 1992. Book review—The Jackson Hole elk herd: Intensive wildlife management in North America. Journal of Range Management 45:315–316.
- KAY, C. E. 1994a. Aboriginal overkill: The role of Native Americans in structuring western ecosystems. Human Nature 5:359-398.
- KAY, C. E. 1994b. The impact of native ungulates and heaver on ri-

- parian communities in the Intermountain West. Natural Resources and Environmental Issues 1:23-44.
- KAY, C. E. 1995a. Aboriginal overkill and native burning: Implications for modern ecosystem management. Western Journal of Applied Forestry 10:121–126.
- Kay, C. E. 1995b. An alternative interpretation of the historical evidence relating to the abundance of wolves in the Yellowstone Ecosystem. Pages 77–84 in L. D. Carbyn, S. H. Fritts, and D. R. Seip, editors. Ecology and conservation of wolves in a changing world. Canadian Circumpolar Institute, Edmonton, Alberta.
- KAY, C. E. 1995c. Browsing by native ungulates: Effects on shrub and seed production in the Greater Yellowstone Ecosystem. Pages 310–320 in B. A. Roundy, E. D. McArthur, J. S. Haley, and D. K. Mann, editors. Proceedings: Wildland shrub and arid land restoration symposium. U.S. Forest Service, General Technical Report INT-GRR-315.
- Kay, C. E. 1996a. Wolf recovery, political ecology, and endangered species. The independent Institute, Oakland, California.
- KAY, C. E. 1996b. Ecosystems then and now: A historical-ecological approach to ecosystem management. Pages 79-87 in W. D. Willims, and J. F. Dormaar, editors. Proceedings for the fourth prairie conservation and endangered species workshop. Provincial Museum of Alberta Natural History, Occasional Paper 23.
- KAY, C. E. 1997a. Is Aspen doomed? Journal of Forestry 95(5):4-11.
- KAN, C. E. 1997b. Aspen: A new perspective—implications for park management and ecological integrity. Proceedings of the ninth conference on research and resource management in parks and on public lands, Hancock, Michigan, The George Wright Society.
- KAY, C.E. 1997c. The condition and trend of aspen, Populus tremuloides, in Kootenay and Yoho National Parks: Implications for ecological integrity. Canadian Field-Naturalist 111:607-616.
- KAY, C. E. 1997d. Yellowstone; Ecological malpractice—photo excerpts from a forthcoming book. Political Economy Research Center Reports 15(2):1–40.
- Kay, C. E. 1997e. Aboriginal overkill and the biogeography of moose in western North America. Alces 35:141–164.
- Kay, C. E. 1997f. Testimony presented at the Oversight Hearing on Science and Resource Management in the National Park Service field by the U.S. House of Representatives Subcommittee on National Parks and Public Lands, 25 February 1997, Washington, D.C., U.S. Government Printing Office, Serial Number 195-3.
- Kay, C. E. 1997g. Viewpoint: Ungulate herbivory willows, and political ecology in Yellowstone. Journal of Range Management 50:139-145.
- Kay, C. E. 1997b. Conflicting blueprints for protecting National Parks. Forum for Applied Research and Public Policy 12(4):138-139.
- KAY, C. E., AND S. W. CHARDE. 1992. Reduction of willow seed production by ungulate browsing in Yellowstone National Park. Pages 92–99 in W. P. Clary, E. D. McArthur, D. Bedunah, and C. L. Wambolt, editors. Proceedings of the Symposium on ecology and management of riparian shrub communities. U.S. Forest Service, General Technical Report INT-289.
- KAY, C. E., B. PATTON, AND C. A. WHITE. 1994. Assessment of long-term terrestrial ecosystem states and processes in Banff National Park and the central Canadian Rockies. Resource Conservation, Parks Canada, Banff National Park, Banff, Alberta.
- KAY, C. E., AND C. A. WHITE. 1995. Long-term ecosystem states and processes in the Central Canadian Rockies: A new perspective on ecological integrity and ecosystem management. Pages

- 119-132 in R. M. Linn, editor. Sustainable society and protected areas. The George Wright Society, Hancock, Michigan.
- KAY, C. E., AND F. H. WAGNER. 1994. Historic condition of woody vegetation on Yellowstone's northern range: A critical test of the "natural regulation" paradigm. Pages 151-169 in D. G. Despain, editor. Plants and their environments-Proceedings of the first biennial scientific conference on the Greater Yellowstone Ecosystem. U.S. National Park Service, Denver, Colorado, Technical Report NPS/NRYELL/NRTR-93/XX.
- KAY, C. E., AND J. W. WALKER. 1997. A comparison of sheep and wildlife grazed willow communities in the Greater Yellowstone Ecosystem. Sheep Research Journal 13:6–14.
- Keigley, R. B. 1997. An increase in herbivory of cottonwood in Yellowstone National Park. Northwest Science 71:127–136.
- Kocu, E. 1941. Big game in Montana from early historical records. Journal of Wildlife Management 5:357–370.
- KUNKEL, K. E. 1997. Predation by wolves and other large carnivores in northwestern Montana and southeastern British Columbia. Dissertation, University of Montana, Missoula.
- LAMBERT, P. M. 1997. Patterns of violence in prehistoric huntergatherer societies of coastal southern California. Pages 77–109 in D. L. Martin and D. W. Frayer, editors. Troubled times: Violence and warfare in the past. Gordon and Breach Publishers, New York, New York.
- LANGLORD, N. P. 1972. The discovery of Yellowstone Park. University of Nebraska Press, Lincoln.
- ELE, R. B., AND I. DEVORE, editors. 1968. Man the hunter. Aldine Publishing Company, Chicago, Illinois.
- LEECHMAN, D. 1951. Bone grease. American Antiquity 16:355–356.
 LEFRANC, M. N., JR., M. B. Moss, K. A. PATNODE, AND W. C. SNGG. III., editors. 1987. Grizzly bear compendium. National Wildlife Federation, Washington, D.C.
- LEWIS, H. T. 1973. Patterns of Indian burning in California: Ecology and ethno-history. Anthropological Papers No. 1. Ballena Press, Ramona, California.
- Lewis, H. T. 1985. Why Indians burned: Specific versus general reasons. Pages 75–80 In J. E. Lotan, B. M. Kilgore, W. C. Fischer, and R. W. Mutch, editors. Proceedings-symposium and workshop on wilderness fire. U.S. Forest Service, General Technical Report INT-182.
- Lewis, H. T., and T. A. Ferguson. 1988. Yards, corridors and mosaics: How to burn a boreal forest. Human Ecology 16:57–77.
- LEWIS, M., AND W. CIARK. 1893. The history of the Lewis and Clark expedition. 1964, reprint. Dover Publications, New York, New York.
- LISTUE, A. M. 1989. Rapid dwarfing of red deer on Jersey in the last interglacial. Nature 342:539-542.
- Lowie, R. H. 1909. The northern Shoshoni. American Museum of Natural History Anthropological Papers 2(3):165–306.
- McCane, R.E., and T. R. McCane. 1984. Of slings and arrows: A historical retrospection. Pages 19–72 in L. K. Halls, editor. White-tailed deer: Ecology and management. Wildlife Management Institute and Stackpole Books, Harrisburg, Pennsylvania.
- McNaugiron, S. J. 1992. The propagation of disturbance in savannas through food webs. Journal of Vegetation Science 3:301–314.
- MARTIN, P. S. 1967. Pleistocene overkill. Natural History 76(9):32-38.
- MARTIN, P. S. 1973. The discover of America. Science 179:969-974.
- MARTIN, P. S., AND H. E. WRIGHT. 1967. Pleistocene extinctions: The search for a cause. Yale University Press, New Haven, Connecticut.
- MARTIN, P. S., AND R. G. KILIN, editors. 1984. Quaternary extinctions: A prehistoric revolution. University of Arizona Press, Tucson.

- MARTINEZ, D. 1993. Managing a precarious balance: Wilderness versus sustainable forestry. Winds of Change 8(3):23-28.
- Marison, D. J., B. M. Bianchard, and R. R. Knight. 1991. Food habits of Yellowstone grizzly bears, 1977–1987. Canadian Journal of Zoology 69:1619–1629.
- Mais, L. S., M.E. Sotur, and D. F. Doak. 1993. The keystone-species concept in ecology and conservation. Bioscience 43:219-224.
- Nelson, J. G., and R. E. ENGLAND. 1971. Some comments on the causes and effects of fire in the northern grasslands area of Canada and the nearby United States. 1750–1900. Canadian Geographer 15:295–306.
- Netson, R. K. 1982. A conservation ethic and environment: The Koyukon of Alaska. Pages 211-228 in N. M. Williams and E. S. Hunn, editors. Resource managers: North America and Australian hunter gatherers. American Association for the Advancement of Science, Selected Symposium Number 67.
- NUSON, R. K. 1983. Make prayers to the raven. University of Chicago Press, Chicago, Illinois.
- NELMANN, T. W. 1984. The opossum problem: Implications for human-wildlife competition over plant food. North American Archaeologist 5:287–313.
- NEUMANN, T. W. 1985. Human-wildlife competition and the passenger pigeon: Population growth from system destabilization. Human Ecology 4:389–410.
- NEUMANN, T. W. 1989. Human-wildlife competition and prehistoric subsistence: The case of the eastern United States. Journal of Middle Atlantic Archaeology 5:29-57.
- Nrumann, T. W. 1995. The structure and dynamics of the prehistoric ecological systems in the eastern woodlands: Ecological reality versus cultural myths. Journal of Middle Atlantic Archaeology 11:125–138.
- Norton-Griffits, M. 1979. The influence of grazing, browsing, and fire on the vegetation dynamics of the Screngeti. Pages 310-352 in A. R. E. Sinclair and M. Norton-Griffiths, editors. Screngeti: Dynamics of an ecosystem. University of Chicago Press, Chicago, Illinois.
- OKANNA, H. 1984. The physical condition of red deer falling prey to the wolf and lynx and harvested in the Carpathian Mountains. ACTA Therologica 29:283–290.
- OLISSTED, C. E. 1979. The ecology of aspen with reference to utilization by large herbivores in Rocky Mountain National Park. Pages 89-97 in M. S. Boyce and L. D. Hayden-Wing, editors. North American elk: Ecology, behavior, and management. University of Wyoming, Laramie.
- Oison, D. L. 1983. A descriptive analysis of the faunal remains from the Miller site, Franklin County, Washington. Thesis, Washington State University, Pullman.
- Parista, J. 1969. Solitary rambles and adventures of a hunter in the prairies. Charles E. Tuttle Company, Rutland, Vermont.
- Pannin, M. L., C. T. Romins, and T. A. Hander. 1984. Energy expenditures for locomotion by mule deer and elk. Journal of Wildlife Management 48:474-488.
- POTTER, J. M. 1995. The effects of sedentism on the processing of hunted careasses in the southwest: A comparison of two Pueblo IV sites in central New Mexico. Kiva 60:441-428.
- Power, M. E., D. Timas, J. A. Espis, B. A. Minge, W. J. Bond, L. S. Mills, G. Daley, J. C. Cestile, J. Lubcherco, and R. T. Pane. 1996. Challenges in the quest for keystones. Bioscience 46:669-620.
- Proc. S. J. 1993. Keeper of the flame: A survey of anthropogenic fire. Pages 245–266 in P. J. Cruzan and J. G. Goldanmer, editors. Fire in the environment: Its ecological, climatic, and atmospheric chemical importance. John Wiley and Sons, New York, New York.
- Prest, S. J. 1995. Vestal fires and virgin lands: A reburn. Pages

- 15–21 in J. K. Brown, R. W. Mutch, and C. W. Spoon, and R. H. Wakimoto, editors. Proceedings of the Symposium on fire in wilderness and park management. U.S. Forest Service, General Technical Report INT-320.
- RAMENOINKY, A. F. 1987. Vectors of death: The archaeology of European contact. University New Mexico Press, Albuquerque.
- RAWLLY, E. V. 1985. Early records of wildlife in Utah. Utah Division of Wildlife, Resources Publication 86-2.
- REIPLS, H. M., AND R. E. McCabe. 1997. Of moose and man. Pages 1–75 in A. W. Franzmann and C. C. Schwartz, editors. Ecology and management of North American moose. Smithsonian Institution Press, Washington, D.C.
- Reio, D. K. 1987. Fire and habitat modification: An anthropological inquiry into the use of fire by indigenous people. Thesis, University of Alberta, Edmonton.
- Roe, F. G. 1951. The North American buffalo: A critical study of the species in its wild state. University of Toronto Press, Toronto, Ontario.
- ROMME, W. H., AND D. G. DESPAIN. 1989. Historical perspective on the Yellowstone fires of 1988. Bioscience 39:695–699.
- RUSSELL, O. 1965. Journal of a trapper, 1834–43. A. L. Haines, editor. University of Nebraska Press, Lincoln.
- Samuss, M. D. 1972. Stone age economics. Aldine Press, Chicago, Illinois.
- SAVAGE, M., AND T. W. SWETNAM. 1990. Early mineteenth-century fire decline following sheep pasturing in Navajo ponderosa pine forest. Ecology 71:2374–2378.
- SCHALK, R. F. 1986. Estimating salmon and steelhead usage in the Columbia Basin before 1850: The anthropological perspective. Northwest Environment Journal 2:1-29.
- SCHALK, R. F., AND R. R. MIERENDOWF, editors. 1983. Cultural resources of the Rocky Reich of the Columbia River. Center for Northwest Anthropology, Washington State University, Project Report Number 1.
- SCHULLERY, P., AND L. WHITTLESEY. 1992. The documentary record of wolves and related wildlife species in the Yellowstone National Park area prior to 1882. Pages 1-3 to 1-173 in J. D. Varley and W. G. Brewster, editors. Wolves for Yellowstone? A report to the United States Congress. Volume IV—Research and analysis. National Park Service, Yellowstone National Park, Mammoth, Wyoming.
- Shaw, J. H. 1995. How many bison originally populated western rangelands? Rangelands 17:148–150.
- SHREEVE, J. 1995. The Neanderthal enigma. Avon Books, New York, New York.
- Simms, S. R. 1984. Aboriginal Great Basin foraging strategies: An evolutionary analysis. Dissertation, University of Utah, Salt Lake City.
- Simms, S. R. 1992. Wilderness as a human landscape. Pages 183-201 in S. I. Zeveloff, L. M. Vause, and W. H. McVaugh, editors. Wilderness tapestry. University of Nevada Press, Reno.
- Simbon, T. 1984. Population dynamics of mule deer. Wyoming Archaeologist 27(1-2):83-96.
- Sinclair, A. R. E., and M. Norton-Grieffin, editors. 1979.
 Serengeti: Dynamics of an ecosystem. University of Chicago Press, Chicago, Illinois.
- Sinciair, A. R. E., and P. Arcese, editors. 1995. Screngeti II: Dynamics, management and conservation of an ecosystem. University of Chicago Press, Chicago, Illinois.
- SINGER, F. J., AND J. E. NORLAND. 1994. Niche relationships within a guild of ungulate species in Yellowstone National Park, Wyoming, following release from artificial controls. Canadian Journal of Zoology 72:1383-1394.
- SINGER, F. J., A. HARTING, K. K. SYMONDS, AND M. B. COUGHENOUR. 1997. Density dependence, compensation, and environmental

- effects on calf mortality in Yellowstone National Park. Journal of Wildlife Management 61:12–25.
- SMITH, E. A. 1983. Anthropological applications of optimal foraging theory: A critical review. Current Anthropology 24:625-651.
- SMITH, E. A., AND B. WINTERHALDER, editors. 1992. Evolutionary ecology and human behavior. Aldine de Gruyter, New York, New York.
- SMITH, M. T. 1987. Archaeology of aboriginal culture change in the interior southeast. Florida State Museum, Gainesville.
- Snow, D. R. 1995. Microchronology and demographic evidence relating to the size of pre-Columbian North American Indian populations. Science 268:1601–1604.
- SPECK, F. G. 1939. Aboriginal conservators. Bird Lore 40:258–261.
 STANNARD, D. E. 1989. Before the horror: The population of Hawaii on the eve of western contact. Social Science Research Institute, University of Hawaii Press, Honolulu.
- STANNARD, D. E. 1992. American holocaust. Oxford University Press, New York, New York.
- STUTIAN, A. F. 1991. Territorial stability as a factor in the occurrence and perpetuation of inter-group buffer zones. Michigan Discussions in Anthropology Hunter-Gatherer Studies 10:89-105.
- STEPHENS, D. W., and J. R. KREBS, 1968. Foraging theory. Princeton University Press, Princeton, New Jersey.
- STINER, M. C. 1990. The use of mortality patterns in archaeological studies or hominid predatory adaptations. Journal of Anthropological Archaeology 9:205–251.
- STUART, A. J. 1991. Mammalian extinctions in the late Pleistocene of northern Eurasia and North America. Biological Review 66:453–562.
- STUART, A. J. 1993. The failure of evolution: Late Quaternary mammalian extinctions in the holarctic. Quaternary International 19:101–107.
- SWEENAM, T. W., AND C. H. BAISAN. 1996. Historical fire regime patterns in the southwestern United States since A.D. 1700. Pages 11–32 in C. D. Allen, editor. Fire effects in southwestern forests: Proceedings of the second La Mesa fire symposium. U.S. Forest Service, General Technical Report RM-286.
- TATTERAL L. 1995. The fossil trail: How we know what we think we know about human evolution. Oxford University Press, New York, New York.
- Tauman, J. E., and L. W., Rouman. 1996. Recent range expansion and distributional limits of the nine-banded armadillo (Dasypus novemcinctus) in the United States. Journal of Biogeography 23:635-648.
- TAYLOR, R. J. 1984. Predation. Chapman and Hall, New York, New York.
- TELFER, E. S., AND J. P. KEINLL. 1984. Adaptation of some large North American mammals for survival in snow. Ecology 65:1828-1834.
- Terres, S. A. 1987. Do predators always capture substandard individuals disproportionately from prey populations? Ecology 68:669-674.
- Thosas, G. 1977. Fire and for trade: The Saskatchewan district: 1790–1840. The Beaver (Antumn):32-39.
- TOUCHAN, R., C. D. AULEN, AND T. W. SWETNAM. 1996. Fire history and climatic patterns in ponderosa pine and mixed-conifer forests of the Jemez Mountains, northern New Mexico. Pages 33-46 in C. D. Allen, editor. Fire effects in southwestern forests: Proceedings of the second La Mesa fire symposium. U.S. Forest Service, General Technical Report RM-286.

- TOUCHAN, R., T. W. SWITNAM, AND H. D. GRISSINO-MAYER. 1995. Effects of livestock grazing on pre-settlement fire regimes in New Mexico. Pages 268-272 in J. K. Brown, R. W. Mutch, C. W. Spoon, and R. H. Wakimoto, editors. Proceedings: Symposium on fire in wilderness and park management. U.S. Forest Service, General Technical Report INT-320.
- TRUETT, J. 1996. Bison and elk in the American southwest: In search of the pristine. Environmental Management 20:195–206.
- TURNER, N. J. 1991. Burning mountain sides for better crops: Aboriginal landscape burning in British Columbia. Archaeology in Montana 32:57-73.
- VALE, T. R. 1998. The myth of the humanized landscape: An example from Yosemite National Park. Natural Areas Journal 18:231–236.
- Vanes, S. C. 1977. Bone fragments and bone grease manufacturing: A review of their archaeological use and potential. Plains Anthropologist 22(77):169–182.
- Wagner, F. H., and C. E. Kay. 1993. "Natural" or "healthy" ecosystems: Are U.S. national parks providing them? Pages 257–270. In M. J. McDonnell and S. T. Pickett, editors. Humans as components of ecosystems. Springer-Verlag, New York, New York.
- WAGNER, F. H., R. FORESTA, R. B. GILC, D. R. McCLILLOUGH, M. P. PEL-TON, W. F. PORTER, AND H. SALWASSER. 1995. Wildlife policies in the U.S. national parks. Island Press, Washington, D.C.
- WALTERS, C. J., M. STOCKER, AND G. C. HABER. 1981. Simulation and optimization models for a wolf-ungulate system. Pages 317–337 in C. W. Fowler and T. D. Smith, editors. Dynamics of large mammal populations. John Wiley and Sons, New York, New York.
- WARD, P. D. 1997. The call of distant mammoths: Why the Ice Age mammals disappeared. Copernicus Springer-Verlag, New York, New York.
- WILNER, J. 1994. The beak of the finch. Vintage Book, New York, New York.
- Wist, E. 1995. The way to the West: Essays on the central plains, University of New Mexico Press, Albuquerque.
- Winte, R. 1995. Are you an environmentalist or do you work for a living?—Work and nature. Pages 171–185 in W. Cronon, editor. Uncommon ground: Toward reinventing nature. W. W. Norton and Company, New York, New York.
- WRIGHT, G. A. 1984. People of the high country: Jackson Hole before the settlers. Peter Lang, New York, New York.

Charles E. Kay has a B.S. in wildlife biology and an M.S. in environmental studies, both from the University of Montana, and a Ph.D. in wildlife ecology from Utah State University. He is an adjunct assistant professor in the Department of Political Science and a senior environmental scholar at Utah State University's Institute of Political Economy. Charles takes a "consilience" approach to science, integrating wildlife and vegetation ecology with archaeological and historical research. In 1990 he completed a detailed, independently funded evaluation of Yellowstone National Park's "natural-regulation" management program. He continues to study Rocky Mountain ecosystems through contracts with Parks Canada, the U.S. Agricultural Research Service, the U.S. Forest Service, and large ranch owners. Charles is currently writing a book for Oxford University Press on Native Americans' roles in structuring western North American ecosystems, and he is compilling ecological observations from the original Lewis and Clark journals. As a long-time researcher, hunter, wildlife photographer, and natural history writer, Charles is familiar with the Rocky Mountains from Arizona to Alaska.

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