What Remains

Species Identification and Bone Histology

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Microscopic problems of historical research can and should be made macroscopic—capable of reflecting worlds larger than themselves. It is in this reflected flicker of truth, the revelations of the general in the particular, that the contribution of the historical method to social science will be found.

> M. M. Postan 1939

Thousands of bone fragments were recovered during excavation of the Donner family camp site at Alder Creek (figure 6.1). Determining what animals were present in the assemblage was a priority, but it was not an easy ta mply distinguishing human from nonhuman bone was challenging vecause of the extensive fragmentation, butchering, and burning that took place during entrapment, in addition to physical and chemical changes that occurred after burial. Fortunately, other archaeologists and forensic scientists have developed molecular, genetic, and histological techniques that can be used to discriminate small, burned fragments of human bone from those of other large mammals. In circumstances like those at Alder Creek—where the organic component is almost completely destroyed—microscopic study of the bone tissue (histology) is the simplest and most effective technique for identification.¹ We used this technique to identify the largest and best preserved bone fragments

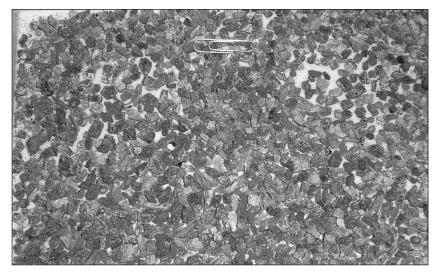


Figure 6.1. Fragments of bone recovered from the Alder Creek site.

from the Alder Creek assemblage with the goal of understanding some of the strategies employed for survival by members of the party. We begin by explaining quantitative (based on measurements) and qualitative (based on observation of qualities) methods for differentiating human bone from that of other animals, emphasizing large mammals likely to be present in the Donner assemblage—cattle (*Bos*), horses (*Equus*), and deer (*Odocoileus*). After presenting the results of our study, we compare the findings to previous bone chemistry analyses from the Donner Lake site and to historical accounts of dietary adaptations at Alder Creek. The identification of bone fragments from the campsite hearth at Alder Creek can be used to understand not only the species utilized for food but also the strategies employed for survival by members of the party.

Macroscopic Osseous Analysis (contributed by Guy Tasa)

The first step in the analysis of a large collection of fragmentary burned bones is to roughly characterize the bone assemblage by counting and weighing the material.² In total, 16,204 bone fragments (2,281.45 grams, or 5.03 pounds) were recovered from the 2003 (1,085 fragments, 249.69 grams) and 2004 (15,119 fragments, 2,031.76 grams) excavations at

Size class	Weight	Examples		
Class I	<100 g	Meadow mice, shrews		
Class II	100–700 g	Squirrels, chipmunks, gophers		
Class III	700 g–5 kg	Rabbits, hares, skunks		
Class IV	5–25 kg	Coyote, bobcat, dog		
Class V	25–225 kg	Deer, bear, sheep, antelope, humans		
Class VI	>225 kg	Elk, horse, cow, bison		
Class X	N/A	Unidentifiable		

Table 6.1. Mammalian size classes	Table	6.I .	Mammalia	an size	classes
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Source: After Thomas, "Great Basin Hunting Patterns"; Schmitt, "Zooarchaeology of Times Square Rockshelter."

Alder Creek. The vertebrate faunal material was sorted and quantified by visual inspection first. Based on the overall robusticity, shaft curvature, and bone thickness, each fragment was assigned to a very general bone/ animal size class (table 6.1). This approach may be problematic for our assemblage, however, because bone is known to shrink with burning;³ as a result, larger animals may be classified into smaller size categories.⁴ Thus, the category assignments for the Alder Creek assemblage should be considered an initial approximation. The number of identifiable specimens per taxon (NISP) was established using reference materials and collections.⁵

Although extensive fragmentation precluded the identification of size class or skeletal element for most of the fragments in the assemblage (87 percent), 2,046 long bone shaft fragments, 18 tooth fragments, and 33 articular/other fragments could be identified (table 6.2). These identified pieces were among the largest and best-preserved fragments, largely derived from the half-inch and quarter-inch screens (table 6.3). They comprised 13 percent of the total number of pieces and 43 percent of the volume by weight (table 6.4). There was a single bone fragment in Class II (rodent-sized mammals), 3 fragments in Class III (rabbit-sized mammals), and 15 in Class IV (dog-sized mammals). The majority of identifiable bone in the assemblage was in Class V (deer-, human-, and bear-sized mammals), and there were 3 fragments in Class VI (elk-, cow-,

	Count		Weight (g)		
Element	n	%	g	%	
Unidentifiable	14,108	87.1	1,257.35	55.I	
Long bone fragment	2,046	12.6	976.05	42.8	
Tooth fragment	18	<1.0	2.43	<1.0	
Articular/other	33	<1.0	45.60	2.0	
Total	16,204		2,281.45		

Table 6.2. Identified skeletal elements from Alder Creek Camp

and horse-sized mammals). A single bone was identified as *Bos taurus* (domestic cow) based on a nearly complete lateral malleolus (*os maleo-lare*). In addition, 15 fragments of mostly tooth enamel were consistent with the order Artiodactyla (even-toed ungulates including cow, deer, and sheep). Of these, 7 specimens were suspected to belong to the genus *Odocoileus* (deer). Macroscopic observations left some 13,562 bone fragments classified simply as "unidentified mammal," and another 537 fragments remained entirely unidentifiable.⁶

Taxonomic Identification and Characteristics of Mammalian Bone

Because macroscopic analysis left much of the bone unidentified or broadly categorized, we turned to light microscopy in an attempt to identify species beyond the level of the size classes.⁷ We first sought to determine whether the hearth assemblage contained any human bones. Human bones can be discriminated from those of other large mammals using quantitative histological techniques to measure microarchitectural features of bone tissue, that is, the size of basic structural units of the bone.⁸

Once the human versus nonhuman question was addressed, we sought to understand more about the starvation diet during the winter of 1846– 47. Our basic question was, what other animals can be identified in the osseous remains? Focusing on the nonhuman mammalian limb bone fragments, we identified taxa using a qualitative approach that is based on previously described ancestral or phylogenetic differences in the bone microarchitecture.⁹

This qualitative approach is not based on the presence or absence of

		Mesh size			
Description	Common name	½" (G2)	¼" (G3)	⅓" (G4)	
Order Artioda	ictyla				
Unspecified	Artiodactyla		6	9	
Bos taurus	Domestic cow	I.			
Class II	Rodent-sized mammals			I	
Class III	Rabbit-sized mammals		I	2	
Class IV	Dog-sized mammals		5	10	
Class V	Deer and human-sized mammals	56	1,419	592	
Class VI	Elk and cow-sized mammals		3		
Class X	Unidentifiable mammals	I.	696	12,865	
Unidentifiable				537	
#Total		58	2,130	14,016	

Table 6.3. Identified vertebrate species at Alder Creek Camp by screen size

Table 6.4. Summary of identified vertebrate species at Alder Creek Camp

		NISP		Weight	
Description	Common name	n	%	g	%
Order Artio	dactyla				
Unspecified Artiodactyla		15 ^a	<1.0	4.10	<1.0
Bos taurus	Domestic cow	I	<1.0	4.83	<1.0
Class II	Rodent-sized mammals	I	<1.0	0.05	<1.0
Class III	Rabbit-sized mammals	3	<1.0	0.48	<1.0
Class IV	Dog-sized mammals	15	<1.0	4.01	<1.0
Class V	Deer and human-sized mammals	2,067	12.8	1,020.87	44.7
Class VI	Elk, horse, and cow-sized				
	mammals	3	<1.0	5.96	<1.0
Class X	Unidentifiable mammals	13,562	83.7	1,228.87	53.9
Unidentifiabl	e	537	3.3	12.41	<1.0
Total		16,204		2,281.45	

Note: All the identified vertebrate species are in the class Mammalia. ^aSeven specimens strongly suspected to be deer (*Odocoileus* sp.).

unique types of bone tissue; instead, variation occurs in the arrangement, distribution, and life history of bony tissue.¹⁰ These differences were first described in the early part of the nineteenth century,¹¹ and a basic review is provided here to explain the rubric we developed to identify the nonhuman bone specimens.

Bone is divided into primary and secondary tissue. Primary bone includes three types: lamellar, laminar (plexiform), and primary osteons. *Lamellar* bone is a rapidly formed, relatively less organized type of bone tissue. Small regions of this type of bone are common in mammals; it is found in pockets at the internal (endosteal) surfaces of the bone shaft, in circumferential layers at the external (periosteal) surface of the shaft, and in the interstitial spaces between more organized types of tissue.¹² *Laminar* bone has a more organized, predictable structure with primary bone tissue arranged around a network (plexus) of intercommunicating blood vessels that resemble a brick fireplace in cross section.¹³ This type of bone, also known as plexiform bone, occurs as the primary tissue type in cows, deer, elk, antelope, sheep, and some large canines.¹⁴ Laminar (plexiform) bone is only rarely seen in humans.¹⁵

Bone tissue is also organized into structures called osteons. Unlike the above tissue types, where bone is organized circumferentially around the shaft, osteons are composed of bone tissue organized around a central canal. There are two types of osteons (primary and secondary) relevant to distinguishing human from nonhuman long bones, and both are easily observed in the light microscope.

Primary osteons consist of lamellar bone tissue arranged around a central canal.¹⁶ Secondary osteons differ from the primary type in several ways. Unlike primary osteons, which have very little internal organization,¹⁷ secondary osteons are highly organized internally and can be differentiated by a series of concentric rings encircling a central canal.¹⁸ Each secondary osteon is also separated from the surrounding lamellar bone by a distinct translucent "cement" layer. Secondary osteons form as an animal grows and matures through a process of remodeling the primary bone tissue. Specialized bone cells remove regions of primary bone and fill in those spaces with the new, highly organized bone tissue.

Because of the nature of this remodeling process, secondary osteons appear to overlap in a seemingly haphazard arrangement when viewed in cross section. Secondary osteons overlie regions of primary tissue as well. Primary osteons often occur in large regions in horse and pig bone, but they also have been documented occasionally in human tissue.¹⁹ Secondary osteons are common in rat, rabbit, dog, horse, deer, cow, and human bone.²⁰

Alder Creek Bone Histology

To carry out our inquiry, we began by calculating the desired sample size of Class IV and V long bone fragments based on the following rationale: (1) the assemblage includes 2,067 fragments (see table 6.3) of Class V bones—the category that could include human; and (2) the possible proportion of human in the diet was estimated to be only 1 percent, calculated by the number of months of entrapment and the probable onset of cannibalism between February 23 and March 3, 1847.²¹

Using these parameters, statistician Terry Allen (of the Department of Sociology, University of Utah) ran one thousand simulations to determine the number of fragments we had to sample in order to be 95 percent confident that human bone would be found in that sample if such tissue were being processed in and around the hearth. He determined that by randomly selecting 104 fragments (the upper limit of the confidence interval) from the Class V sample, we could be 95 percent confident of getting a human bone if human tissue constituted 1 percent of the bone refuse. By randomly selecting 106 bones, we could be 99 percent confident; 108 bones would provide 99.9 percent confidence. Given the possibility that the human bone makes up less than 1 percent of the collection, we selected 120 as the number of Class IV and V bone fragments that would have to be profiled in order to be statistically confident that there is no human bone in the intensively processed assemblage from Alder Creek.

Unfortunately, given poor preservation and the fragmentary nature of the assemblage, randomly selecting 120 pieces of bone from the appropriate size class was not possible. After the bones in size classes IV, V, and VI were examined for preservation that could withstand histological processing, only 85 long bone fragments could be identified. The sample was derived from the one-half-inch and one-quarter-inch fractions and consisted of 10 bones in Class IV and 74 bones in Classes V and VI.²² This sample represents 23.5 percent of the 362 bones with evidence of processing scars, 0.04 percent (85/2,046) of the total number of long bone fragments, 0.04 percent (85/2,070) of the total sample of Class V

and VI bone fragments, and 0.005 percent of the total assemblage from the Donner family camp. The remaining bones in the assemblage were either too small to process or had had extensive damage from breaking, burning, cryoturbation (that is, freeze-thaw action), and cracking.

Thus, if our sample size included 90 bone fragments, we could claim 68 percent confidence that one human bone would be included in our sample if 1 percent of the refuse at Alder Creek actually contained human bone. With a sample size of 85, therefore, the power of our study is diminished. Yet this small sample size is supplemented by the previous macroscopic analysis, which excluded another 20 fragments as being nonhuman based on gross morphology (15 Artiodactyla, 1 *Bos*, 1 Class II, and 3 Class III; see table 6.3). This means that the combined microscopic and macroscopic findings provided a total of 105 fragments for analysis²³—just above the 104 fragments calculated as the minimum number for 95 percent certainty that human remains were not part of this assemblage.

Identification Rubric

Laboratory methods are outlined in the notes section of this chapter.²⁴ Bone samples were first evaluated quantitatively for the presence of human specimens. Human bones can be distinguished by osteon and Haversian canal diameters as well as the number of osteons per square millimeter. Quantitative, morphometric considerations useful for discriminating humans from other mammals have been described by forensic scientists and archaeologists.²⁵ As a supplement to the published literature on histomorphometrics,²⁶ we compared values from histomorphometric analysis of ten adult human femora (n=200 osteons measured).

Secondary osteons and their central canals are significantly larger in adult humans than in the other large mammals likely to be in this assemblage: horses (*Equus ferus caballus*), cows (*Bos taurus*), and deer (*Odocoileus* sp.) (table 6.5). Adult human osteons commonly range in diameter from 174 micrometers to 506 micrometers²⁷ and Haversian canal diameters range in size from 33 micrometers to 100 micrometers.²⁸ Secondary osteon density in long bones is usually one to two Haversian canals per square millimeter. Osteon diameters, particularly the minimum diameters, are on average smaller in horses (158–223 µm), deer (67–165

Table 6.5. Osteon and Haversian canal diameters

from published sources and this study

	Osteon diameter		Haversian canal diameter	
	Min.	Max.	Min.	Max.
Description	(µm)	(µm)	(µm)	(µm)
Cattaneo et al., "Determining the Human	n Origin"			
Human	277	353	58	77
Nonhuman ^a	223	297	34	45
Dittmann, "Histomorphometric Investiga	ations"			
Human	174	281	33	50
Horse	158	205	26	33
Cow	121	157	18	23
This study ^b				
Human	244	506	56	74
Cow	52	163	9	38
Deer	67	143	17	29
Elk	108	157	14	30

^aThe nonhuman sample in Cattaneo et al., "Determining the Human Origin" consisted of 5 cows, 6 sheep, 1 horse, 1 dog, and 1 cat.

^bThe reference sample in the current study included 1 cow, 1 deer, 1 elk, and 10 adult human femurs.

 μ m), and cows (79–250 μ m), which can have as many as twenty to thirty Haversian canals in a square millimeter. Minimum Haversian canal diameters are similarly smaller in horses (24–54 μ m), deer (17–29 μ m), and cows (9–23 μ m). On the basis of measurements of the minimum and maximum diameters of osteons and their central canals, human long bone fragments can be discriminated from nonhuman fragments with a high level of precision.²⁹

Once the specimens were evaluated for the presence of human remains, the nonhuman component of the sample was qualitatively sorted into family-level taxonomic groups. The major qualitative difference distinguishing human from nonhuman mammalian long bone fragments is a predominance of large, densely packed, overlapping, Haversian systems.³⁰ Primary lamellar bone is only commonly found as new bone

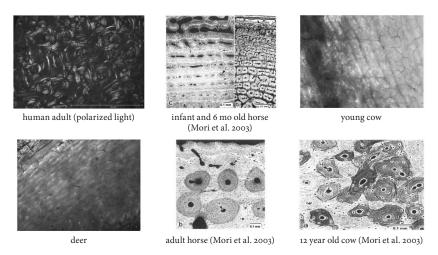


Figure 6.2. Faunal comparison collection: cow, deer, human, horse. Images from this study and Mori et al., "Comparative Histology," 43.

apposition at the periosteal surface (or outer circumference),³¹ although occasionally there are pockets of lamellar bone at the endosteal surface or in regions of the cross section that have yet to be remodeled. Primary band between the presence of plexiform bone makes it very unlikely that a particular bone fragment belongs to a human.³² Plexiform bone is only rarely seen in humans, but such bone may form when a child is going through a very rapid growth spurt or experiences some form of nutritional stress, particularly iron-deficiency anemia.³³ If human children were present in the assemblage, we would expect to see primary bone, and perhaps plexiform bone given the likelihood of nutritional stress. In a long bone fragment the size of an adult human's, the presence of plexiform bone is not human.

Qualitative criteria for identifying bones in the nonhuman component of the sample were taken from the published literature and from bone samples of fauna processed for reference use in this analysis: elk, bear, deer, cow, antelope, pig, and coyote (figure 6.2).³⁴ The nonhuman component of the Alder Creek sample was evaluated to discriminate perissodactyls (odd-toed hoofed animals, such as horses) from artiodactyls (even or two-toed, hoofed animals, such as cows and deer). Young equids (horses) were identified on the basis of large areas of primary osteons, banded primary osteons, and "budding" at the periosteal surface.³⁵ Nonhuman primates and occasionally humans can also have large areas of primary osteons in some skeletal elements, but banding is rarely seen in adult humans (an estimated 3.3 percent of individuals), and it generally occurs in an isolated row within the periosteal lamellar region.³⁶

Cervids (deer) were identified from specimens with secondary osteon remodeling at the interior of the cortex, situated between layers of circumferential lamellae and plexiform bone at the outer surface.³⁷ Bovids (cows and oxen) also have well-organized plexiform tissue, with large lamellar regions up to 200 micrometers in diameter, which persists through most of their adult life.³⁸ In older bovine individuals, secondary osteons can develop near the endosteal surface, eventually remodeling the entire surface.³⁹ Bovine secondary osteons have few lamellae and relatively few osteocytes,⁴⁰ and they also tend to have a thick osteoid perimeter around the Haversian canal in contrast to the thinner perimeter of cervids and equids.

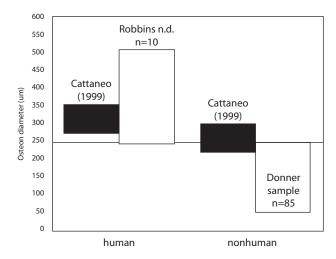
Using the observations about bone biology described above, we developed the following rubric.

Our criteria for assigning a bone sample to the "human" category were (1) lack of plexiform bone, (2) a substantial amount of secondary osteon remodeling, (3) Haversian canal diameters of 50–100 micrometers for secondary osteons, (4) secondary osteon diameters of 240–500 micrometers, and (5) an average of one or two secondary osteon canals per square millimeter of bone surface.

Bones assigned to the equid (horse) category had (1) large fields of primary or secondary osteons arranged in bands circumferentially within annulations (or rings) of lamellar bone, (2) evidence of budding, and (3) banded secondary osteons with dimensions in the range of equids.

Bones were assigned to the cervid (deer) category when (1) the plexiform bone had lamellar sheets of smaller diameter (75–125 μ m wide) than those of cows (175-250 μ m), (2) remodeling occurred such that the highest concentrations of (and the oldest) secondary osteons were situated between plexiform regions at the periosteal and endosteal envelopes, and (3) osteons were within the dimensions accepted for deer.

Bones were assigned to the bovid (cow) category when they had (1) large areas of plexiform bone with a widely spaced plexus of canals (lamellar regions 175–250 μ m wide), (2) secondary remodeling with sporadic,



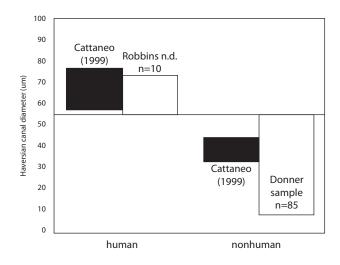
(above and opposite) **Figure 6.3.** Histomorphometrics in contemporary reference samples. Data from Cattaneo et al., "Determining the Human Origin."

large secondary osteons, (3) relatively large osteocytes arranged in a few circumferential lamellar sheets, and (4) the presence of a thick layer of osteoid around the Haversian canal.

Results

Histomorphometrics demonstrated that none of the bone fragments had measurements consistent with human tissue (table 6.5). Osteon diameters were limited to a range of $48-250 \mu$ m; Haversian canal diameters were 5–55 µm. These results fall at the low end or outside the range expected for humans (osteons, 175+ µm; Haversian canals, 30–75 µm) (figure 6.3). In addition, none of the samples studied demonstrated the tightly packed, overlapping osteon pattern expected in human bone tissue (except for one Class IV bone [5 mm in diameter], which is discussed below). Thus, all of the specimens had features inconsistent with human bone: large regions of plexiform tissue, banded osteons, and/or averages of ten to forty osteons per square millimeter.

Therefore, on the basis of the microscopic quantitative and qualitative expectations outlined above, investigators identified no adult human



remains in this assemblage of 85 large mammalian long bone fragments. In addition, the macroscopic analysis confirmed another 20 fragments as being nonhuman (15 Artiodactyla, 1 *Bos*, 1 Class II, and 3 Class III). So the combined microscopic and macroscopic findings identified a total of 105 fragments as nonhuman.⁴¹

To more closely examine this nonhuman bone, fragments were sorted into categories at the level of the family based on qualitative assessment of the primary bone tissue architecture. Of the total sample (n=85), 12 (14.11 percent) individuals had primary and/or secondary banded osteons; 61 (71.76 percent) individuals had plexiform bone and/or secondary remodeling. Of the latter category, 32 (37.65 percent) had plexiform bone and secondary osteons with large lamellar sheets (>200 µm in width), large osteocytes, and thick osteoid perimeters (figure 6.4). Based on our qualitative rubric, we estimate that there are 12 horse, 29 deer, and 32 cow bone fragments in this sample. This is not to suggest that is the number of individual animals—just the number of specimens belonging to those kinds of animals. There were two additional samples in Class V that could not be assigned a taxonomic category because diagenesis obliterated the microstructure.

Ten of the fragments (11.76 percent) considered here belonged to size class IV. For this portion of the analysis, we were looking to identify

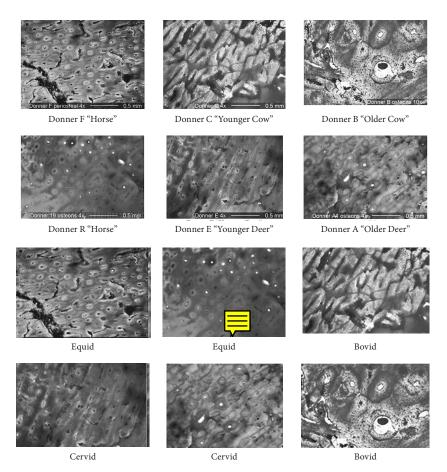


Figure 6.4. Images from Donner Party bone samples. From Dixon et al., "Men, Women, and Children," 640.

nonhuman mammals and for evidence of infant humans; thus, the rubric for identifying the bone as human was different. Human infant long bones are composed primarily of lamellar woven bone, and secondary remodeling begins late in infancy and continues during the second or third year of life. A Class IV long bone fragment primarily constituted of lamellar woven bone would be consistent with an identification of human, in that infant humans could not be eliminated as a possibility. Upon examining these Class IV fragments in the microscope, we found that three were significantly affected by diagenesis and could not be further identified. Six bone fragments in this category had primary lamellar bone tissue that was not the disorganized woven bone diagnostic of human infants. The nine fragments we examined may have come from any of the following Class IV animals that would have been active in the Sierra Nevada at that time: gray, kit, and red fox; wolf, coyote, and dog; and porcupine, muskrat, and beaver.

One Class IV fragment (with a total bone cross-section diameter of 5 mm) had butchering marks (see chapter 7). Histological analysis demonstrated that the bone had undergone diagenesis. The surface could be imaged using reflected light; however, the poor quality of the resulting image made histomorphometrics impossible. Qualitatively, this bone demonstrated dense Haversian remodeling, with secondary osteons present across the entire cross section. This bone tissue type is inconsistent with a human long bone fragment of this size. The bone's diameter could be consistent with a human infant less than one year old, but at this age, we would expect little to no remodeling. On the basis of the diameter and qualitative features of the bone cross section, we can tentatively attribute this bone to the family Canidae. We can therefore rule out the family Leporidae because rabbits and hares should be classified in Class III, rather than Class IV.⁴²

Discussion

Our results indicate that the bone we sampled from the Alder Creek assemblage did not include human tissue. The bone fragments did, however, include remains from horse, deer, cattle, and dog (equids, cervids, bovids, and a canid). While our findings allow us to provide a descriptive account of some of the larger mammals present in the bone refuse at Alder Creek, there are also limitations to our interpretation. We cannot determine how many individuals are represented or how old the animals were when they died. Moreover, we cannot determine the proportion of the diet that these different species constituted. Our analysis demonstrates part of the starvation diet at the Donner campsite, but there is likely a more diverse array of species represented in the thousands of bone fragments too small or too fragile to be processed for histology. Moreover, other animals were probably consumed but did not leave recognizable traces in the archaeological record. How do our findings compare to earlier excavations of Donner Party encampments? Small, weathered bone fragments were recovered during Donald Hardesty's excavations at Donner Lake.⁴³ Forensic anthropologist P. Willey examined bone fragments from the Murphy cabin assemblage but found them to be too small and deteriorated for identification or to recognize processing scars from cannibalism (see chapter 3). Radio-immunoassay techniques (used for measuring antigen concentrations in the blood) were subsequently carried out on bones from Donner Lake and revealed that some of the bones were human.⁴⁴ Radioimmunoassay analysis was not carried out on the severely weathered bone excavated by Hardesty at Alder Creek.

If cannibalism occurred at the Alder Creek site, there are numerous possible explanations for why no human remains were identified in this analysis. Perhaps human remains were processed away from the habitation area, at a locality that remains unexcavated (see chapters 5 and 7). It is possible that human bodies were treated differently, out of respect for those who remained alive or in an attempt to conceal such activities from others in the camp. Cannibalism may have occurred for only a short time at the end of the occupation of the site (see chapter 1), and thus the bodies were unlikely to have been processed down to the bone.⁴⁵ If cannibalism occurred, the remains may not have been fully exploited in the manner of the animal carcasses. If humans were not reduced to bones that were then crushed and boiled for marrow and grease, they are less likely to be represented in this assemblage. Thus, there are many explanations for the absence of human remains in the hearth deposits; therefore, the result of this study does not confirm or deny the practice of cannibalism at the site.

The other remains we were able to identify most likely consist of cow, deer, horse, and dog, which were presumably consumed by the individuals trapped at Alder Creek. We know that the Donner Party left for California with a substantial number of cattle, some horses, and dogs. We also know that horses were left by the relief parties in March 1847. Eliza Donner Houghton's and Patrick Breen's diaries speak mainly of cattle being used for food, but they both describe the relief parties organizing horses to be taken to Donner Lake to be left with the survivors and eaten once the other provisions ran out.⁴⁶ However, Eliza Donner described Mr. Eddy having meat from a bear that was killed early on during the winter. She also described carcasses of bullocks and horses among the

remains at the Donner Lake camp when the last relief party arrived to rescue Mr. Keseberg,⁴⁷ and we know that at least one deer was killed during the "Forlorn Hope" escape attempt. In addition, acquaintances of the survivors have recounted that horses were eaten. For example, G. W. Thissell, who was the neighbor of George Donner, Jr., reported, "They saw their provisions fading away. First on half rations, then just enough to sustain life. The last ox and horse had been killed, then their faithful dogs, and, lastly, they boiled the old ox and horse hides, and were living on them when they were rescued."⁴⁸

This archaeological project was designed to expand our understanding of the Donner Party's experience. The osseous remains represent the starvation diet of the camp. They demonstrate a struggle to survive that transgressed social and cultural boundaries, such as those that separate horses and dogs from other animals commonly used for food (see also chapter 7). The bones also suggest differential treatment afforded to the human remains, which also made them less likely to be preserved or discovered.

Even within the extremes of cold and starvation, meaning persists. Culture is the only force in nature that can define food as a moral substance even in the face of incredibly harsh biological circumstances.⁴⁹ The absence of human remains in the food refuse is not evidence that the Donner family never resorted to cannibalism. It represents instead a period of time when these families starved themselves and their children to avoid cannibalism. The fragments of bone in the hearth represent the desire to treat the dead as different from other kinds of food. Although osteologists rarely feel this way, in the case of the Alder Creek Camp, the *absence* of human tissue in the food refuse is the most interesting result.

Acknowledgments

Chapter contributor Guy Tasa conducted macroscopic analysis of the bone while working at the University of Oregon's Museum of Natural and Cultural History. Tasa's original macroscopic analysis was included in a paper presented to the Society for Historical Archaeology in 2006, entitled "The Donner Family Camp Site Bone." Contributors Ryne Danielson and Matt Irish assisted Dr. Robbins with the bone histology related to this project while working on their undergraduate degrees in anthropology at Appalachian State University. The authors thank Dr. John Lukacs for providing lab space and equipment for histological analysis at the Department of Anthropology, University of Oregon. We also thank Eric Altman, Tammy Edwards, Melissa Hanks, Melody Heath, and Mary Allison Jobe for photography, sectioning, and other assistance with the preparation and analysis phase of the project.

Notes

The chapter epigraph is from Postan, Historical Method.

1. Owsley, Mires, and Keith, "Case Involving Differentiation"; Ubelaker, *Human Skeletal Remains*; Dix, Stout, and Mosley, "Bones, Blood, Pellets"; Cattaneo et al., "Determining the Human Origin," 181; Mulhern and Ubelaker, "Differences in Osteon Banding"; Dittmann, "Histomorphometric Investigations"; Hillier and Bell, "Differentiating Human Bone," 249.

2. "Macroscopic" refers to analyses in which objects are observable with the unaided eye. For this project, macroscopic analyses of bone were conducted by Guy Tasa at the University of Oregon; see Tasa, "Donner Family Camp Site." An elaboration of Tasa's findings is presented in Dixon et al., "Men, Women, and Children."

3. See, for example, Shipman, Foster, and Schoeninger, "Burnt Bones and Teeth."

4. In the end, the histological analyses supported the size classifications based on visual observation.

5. See Grayson, Quantitative Zooarchaeology; Lawrence, Post-Cranial Skeletal Characters; Sisson and Grossman, Anatomy; Olsen, Mammal Remains; Schmid, Atlas of Animal Bones; Gilbert, Mammalian Osteo-Archaeology; Brown and Gustafson, Key.

6. Although the initial findings at Alder Creek suggest some similarity to Hardesty's Murphy cabin site (see chapter 3 and Hardesty, *Archaeology*) with cow bones being present, the recent analysis exhibits more contrast: Alder Creek Camp had probable deer along with rodent-, rabbit-, and dog-sized mammals, while the Murphy cabin had horse/ mule, bear, and human. The results suggest that members of the Donner family camp at Alder Creek procured a greater variety of wild animals for food in addition to the domestic species that traveled with them. Human remains may be present in the assemblage, but they are unidentifiable by visual means. It is also possible that the stranded emigrants treated human remains differently; they may have been butchered differently or only tissue and not bone was removed or consumed. A detailed comparison between the Alder Creek camp and Donner Lake camp bone assemblag

7. Histological analyses were conducted by Gwen Robbins, wratt Irish, Kelsey Gray, and Ryne Danielson.

8. Owsley, Mires, and Keith, "Case Involving Differentiation"; Cattaneo et al., "Determining the Human Origin," 181; Dittmann, "Histomorphometric Investigations."

9. Foote, "Contribution"; Enlow and Brown, "Comparative Histological Study," parts 1–3; Enlow, "Study"; Martin and Burr, *Structure*; Skedros, Su, and Bloebaum, "Biomechanical Implications"; Hidaka et al., "Histomorphometrical Study"; Whyte, "Distinguishing Remains"; Frank et al., "Microdamage in Canine Bone"; Mori et al., "Comparative Histology"; Locke, "Structure of Long Bones." See also Owsley, Mires, and Keith, "Case Involving Differentiation"; and Mulhern and Ubelaker, "Differences in Osteon Banding." Bone tissue types vary among skeletal elements; we limited this analysis to bone tissue types found in the walls of long bone shafts (diaphyseal compact bone).

10. Martin and Burr, Structure.

11. Foote, "Contribution"; see Enlow and Brown, "Comparative Histological Study," part 1. For a historical perspective, see Martin and Burr, *Structure*.

12. Currey, Bones.

13. Martin and Burr, Structure.

14. Enlow and Brown, "Comparative Histological Study," part 3.

15. Zoetis et al., "Species Comparison"; Enlow, "Study." Small regions may form at the periosteal surface when a child is going through a very rapid growth spurt or experiences some form of nutritional stress, particularly iron-deficiency anemia.

16. These osteons are not discrete, bounded units of bone, and in fact, they are not easily distinguished from one another at all; the borders of one primary osteon will be indistinct from its neighbor. Bone tissue formed from primary osteons is not highly organized. These osteons are arranged in a relatively open, loosely organized configuration or sometimes in densely packed bands. See Currey, *Bones*.

17. Enlow and Brown, "Comparative Histological Study," part 3; Hall, *Developmental and Skeletal Biology*; Martin and Burr, *Structure*; Currey, *Bones*. In general, there will be a lot of diversity in the size and shape of secondary osteons within a single bone or region of bone.

18. Currey, Bones.

19. Mulhern and Ubelaker, "Differences in Osteon Banding."

20. Enlow and Brown, "Comparative Histological Study," part 3.

21. Stewart, Ordeal by Hunger, 193, 215-16; Mullen, Donner Party Chronicles, 203.

22. The longest dimension of the specimens included measured between 0.77 centimeters and 2.73 centimeters.

23. See also Dixon et al., "Men, Women, and Children," 647-48n21.

24. Ground sections were prepared using a protocol developed by G. J. R. Maat ("Practicing Methods," 293) for calcined and fragile archaeological bone specimens. Lowviscosity cyanoacrylate, a fast-setting adhesive (most often called by the brand name Super Glue), was applied to the surface of the bone fragments and allowed to harden for twenty-four hours. A diamond-impregnated blade on a slow rotating saw (Buehler Minimet) was used to make the initial sections to approximately 4-millimeter thickness perpendicular to the long axis of the bone. One cut face was then ground and polished with successively finer grit sandpaper (220–600 grit) and then mounted on a glass slide using the cyanoacrylate. After at least two hours of hardening time, the opposite face was ground down to a final thickness of approximately 50–100 micrometers (μ m) and polished with 800-grit sandpaper. The sections were examined using a transmitted light microscope under 40–100X final magnification. Images made from the microscope, known as micrographs, were obtained using a mounted Polaroid digital camera, and

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the images were transmitted to a computer. The osteons were examined both directly through the microscope, and using the micrographs. A 0.5-millimeter scale was calibrated and digitally placed on the images. These images were imported into Photoshop xs and converted to gray scale. The midrange contrast, light, and dark ranges were also adjusted to provide maximum visibility of the microstructure. For archival purposes, the sections were removed from the slide using the solvent xylene, in which they were immersed for twenty-four hours. The finished sections were then permanently mounted onto glass slides using Entellen, a synthetic resin used to mount a bone section to the slide. Sections were examined for the presence of lamellar and laminar bone and for primary and secondary osteons. The entire cross section of the bone was evaluated, including the periosteal and endosteal surfaces and the interior architecture. Qualitative descriptions of the bone architecture were recorded. When secondary osteons were present, digital images obtained from a microscope, called digital micrographs, were imported into ImageJ (National Institute of Health software) for histomorphometrics. After the scale was calibrated, measurements were recorded for all intact osteons and Haversian canals visible in the field $(2.15 \times 1.61 \text{ millimeters})$. The minimum and maximum diameters were measured at the cement line. Records were kept of the number of features measured per specimen, minimum and maximum diameters, the mean measure (and standard deviation) for each feature in each individual, and the number of canals per square millimeter.

25. Owsley et al., "Case Involving Differentiation"; Cattaneo et al., "Determining the Human Origin," 181; Dittman, "Histomorphometric Investigations"; Mori et al., "Comparative Histology."

26. Cattaneo et al., "Determining the Human Origin," 181; Dittmann, "Histomorphometric Investigations"; Walter, Paine, and Horni, "Histological Examination."

27. Secondary remodeling is a process that increases with age, and the density of secondary osteons is variable. In addition, the size of osteons in humans varies with age; however, see Pfeiffer, "Variability in Osteon Size." Osteons in older individuals tend to be smaller than those of younger individuals, with more overlapping and fragmentary osteons that have been remodeled; see Kerley, "Microscopic Determination of Age." Skeletal element is also important, as femoral osteons have a greater diameter than rib osteons, for example; see Pfeiffer, "Variability in Osteon Size."

28. Owsley et al., "Case Involving Differentiation"; Cattaneo et al., "Determining the Human Origin," 181; Dittmann, "Histomorphometric Investigations."

29. Cattaneo et al., "Determining the Human Origin," 181.

30. Foote, "Contribution"; Enlow, "Study"; Enlow and Brown, "Comparative Histological Study," parts 1–3; Owsley et al., "Case Involving Differentiation"; Martin and Burr, *Structure*; Ubelaker, *Human Skeletal Remains*; Skedros, Su, and Bloebaum, "Biomechanical Implications"; Hidaka et al., "Histomorphometrical Study"; Mulhern and Ubelaker, "Differences in Osteon Banding"; Whyte, "Distinguishing Remains"; Frank et al., "Microdamage in Canine Bone"; Mori et al., "Comparative Histology"; Locke, "Structure of Long Bones." 31. Enlow and Brown, "Comparative Histological Study," part 3.

32. Mulhern and Ubelaker, "Differences in Osteon Banding."

33. Zoetis et al., "Species Comparison"; Enlow, "Study."

34. Walter and colleagues (in "Histological Examination") observe that coyote long bones do undergo secondary remodeling later in life, with secondary osteons of 0.1734 average diameter.

35. Enlow and Brown, "Comparative Histological Study," part 3; Mulhern and Ubelaker, "Differences in Osteon Banding"; Mori et al., "Comparative Histology." "Budding" refers to the way in which the primary osteons appear as they form adjacent to plexiform regions; see Currey, *Bones*.

36. Enlow and Brown, "Comparative Histological Study," part 3; Mulhern and Ubelaker, "Differences in Osteon Banding."

37. Enlow and Brown, "Comparative Histological Study," part 3; Owsley et al., "Case Involving Differentiation"; Ubelaker, *Human Skeletal Remains*; Mulhern and Ubelaker, "Differences in Osteon Banding."

38. Currey, Bones; Mori et al., "Comparative Histology."

39. Enlow and Brown, "Comparative Histological Study," part 3; Mori et al., "Comparative Histology."

40. An osteocyte is a star-shaped cell that occupies a hole (lacuna) within mature bone tissue.

41. See also Dixon et al., "Men, Women, and Children," 647–48n21.

42. Enlow and Brown, "Comparative Histological Study," part 3; Hidaka et al., "Histomorphometrical Study"; Frank et al., "Microdamage in Canine Bone"; see also Hillier and Bell, "Differentiating Human Bone."

43. Hardesty, Archaeology, 47–48.

44. Hardesty, Archaeology, 47-48.

45. See, for example, Dixon et al., "Men, Women, and Children."

46. Houghton, Expedition, 78.

47. Houghton, Expedition, 78, 356.

48. Houghton, Expedition, 154.

49. Zentall and Galeff, Social Learning, 166.

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