FOLSOM MAMMOTH HUNTERS? THE TERMINAL PLEISTOCENE ASSEMBLAGE FROM OWL CAVE (10BV30), WASDEN SITE, IDAHO

L. Suzann Henrikson, David A. Byers, Robert M. Yohe II, Matthew M. DeCarlo, and Gene L. Titmus

The 1960s and 1970s excavations at Owl Cave (10BV30) recovered mammoth bone and Folsom-like points from the same strata, suggesting evidence for a post-Clovis mammoth kill. However, a synthesis of the excavation data was never published, and the locality has since been purged from the roster of sites with human/extinct megafauna associations. Here, we present dates on bone from the oldest stratum, review provenience data, conduct a bone-surface modification study, and present the results of a protein-residue analysis. Our study fails to make the case for mammoth hunting by Folsom peoples. Although two of the point fragments tested positive for horse or elephant protein, recent AMS dates indicate that all of the mammoth remains predate Folsom, and horse remains are absent from the Owl Cave collection. Further, no unambiguously cultural surface modifications were identified on any of the mammoth remains. Given the available data, the Owl Cave deposits are most parsimoniously read as containing a Folsom-age occupation in a buried context, the first of its kind in the desert West, but one nonetheless part of a palimpsest of terminal Pleistocene materials.

Durante excavaciones de Owl Cave (10BV30) en Idaho en las décadas de 1960 y 1970 fueron recuperados de los mismos estratos huesos de mamut y puntas de proyectil del estilo Folsom, sugiriendo que se tratara de un yacimiento matanza de mamuts de la era post-Clovis. Sin embargo, nunca se publicó una síntesis de los datos de la excavación y la localidad ha sido removida de la lista de sitios con evidencia de actividad humana asociada con megafauna extinta. Aquí presentamos el fechamiento de muestras de hueso del estrato más antiguo de la cueva, revisamos sus datos de procedencia, realizamos un estudio de la superficie de los huesos, y presentamos los resultados de un análisis de residuos proteicos. Nuestro estudio no logra comprobar la evidencia de cacería de mamut por la cultura Folsom. Aunque en dos de los fragmentos de proyectil se detectaron restos de proteína de caballo o elefante, fechados recientes por AMS indican que todos los restos de mamut preceden el yacimiento Folsom y no hay restos de caballo en la colección de Owl Cave. Además, no se identificó ninguna modificación de superficie de claro origen cultural en los restos de mamut. La interpretación más parsimoniosa de los datos disponibles es que los depósitos de Owl Cave contienen una ocupación de la época Folsom en un contexto enterrado, el primero de este tipo en el desierto del Oeste, pero que sin embargo es parte de un palimpsesto de materiales del Pleistoceno terminal.

Archaeologists know relatively little about the terminal Pleistocene archaeology of the Snake River Plain (SRP). This gap in knowledge stems from both a lack of field research and the often superficial and inadequate reporting of the collections from SRP sites known to contain terminal Pleistocene materials. These issues have prompted the archaeological community to disregard sites such as Jaguar Cave, Wilson Butte Cave, and Kelvin’s Cave as unequivocal associations between people and extinct mammals (Cannon and Meltzer 2004; Henrikson and Long 2007). None of this is to say that people were not present during the terminal Pleistocene in southern Idaho. In fact, fluted and stemmed...
Figure 1. Map of the Snake River Plain showing the location of Owl Cave and other sites mentioned in text: (1) Wasden site, (2) Jaguar Cave, (3) Wilson Butte Cave, (4) Kelvin’s Cave, (5) Lake Terreton, (6) Snake River.

projectile points are relatively common in some areas, especially around the margins of pluvial Lake Terreton (Armstrong et al. 2016). As typical of the desert West, though, such artifacts occur most frequently as isolated surface finds (Long 2007; Titmus and Woods 1991; Yohe and Woods 2000). Excavations at Owl Cave, however, produced fluted points from a buried context, providing one exception to the broader pattern (Butler 1978; Miller and Dort 1978). Viewing these materials within the context of the region’s terminal Pleistocene record makes understanding the Owl Cave record all the more important and motivates the analyses we present below.

Owl Cave is one of three collapsed lava tubes collectively called the Wasden site (Butler 1963, 1968, 1978). Wasden sits on the eastern SRP approximately 48 km west of Idaho Falls, Idaho (Figure 1). While Owl Cave may be one of the most important terminal Pleistocene archaeological sites in western North America, its research potential has yet to be realized. Despite the intensity of Idaho State University’s original excavations in the 1960s and 1970s, and evidence suggesting the use of late Pleistocene fauna (Miller 1982, 1989; Miller and Dort 1978), we lack a synthesis of the excavation data. Furthermore, the fluted points recovered from the same stratigraphic unit as a number of fragmentary pieces of mammoth bone (Mammuthus sp.) appear most consistent with those typed as Folsom elsewhere (Titmus and Woods 1991) and, consequently, their association with mammoth appears anomalous given the known Paleoindian culture history of North America. If these artifacts are both morphologically and chronologically Folsom and they associate contextually with the mammoth remains, then Owl Cave contains an unexpected record of mammoth hunting that postdates Clovis. However, lacking a comprehensive report or a complete account of the site’s fauna assemblage and its relationships with the fluted points, researchers have dropped Owl Cave from the list of sites with...
unambiguous associations between people and extinct megafauna (Grayson and Meltzer 2002).

In this study, we evaluate the associations between the fluted points and the mammoth remains. Our analysis is based, in part, on the provenience data contained in a copy of the Idaho Museum of Natural History (IMNH) artifact catalog, photographs of maps and profiles obtained during a visit to the museum, and copies of original 1960–1970s documents (Miller 1977). We augment this information with a series of new AMS dates, a recent bone-surface modification study of the mammoth remains, and a protein residue analysis of the projectile points in an effort to understand the formational history responsible for the artifacts and animals contained in the Owl Cave deposits.

Excavations at Owl Cave

Idaho State University and the Upper Snake River Prehistoric Society first tested Owl Cave in 1964, with the approval and support of private landowners Leonard Wasden and Ken Huskinson. Full-scale excavations continued with occasional interruptions between 1965 and 1977 (Butler 1968, 1978; Butler et al. 1971). During this time, excavators removed deposits in 2 × 2 m units and natural levels (Figure 2). The archaeological deposits in Owl Cave proved to have substantial depth and include a bison (Bison sp.) bone bed associated with lanceolate projectile points at roughly 3 m below main site datum (BMD). Excavators encountered the fluted points and extinct mammal remains about 5 m below surface (Butler 1968, 1978; Butler et al. 1971). The bison bone assemblage has seen little research and no accelerator dates. Instead, attention has focused on the terminal Pleistocene component that we explore here.

The discovery of a large roof/wall collapse at about 5 m BMD initially provided a difficult obstacle to further excavation. However, the promise of older deposits led to the removal
of this debris in 1971. Soon after, the discovery of fragmentary mammoth (Mammuthus sp.) remains prompted additional excavation. Miller and Dort (1978) returned to the site in 1975 and 1976. This subsequent investigation recovered the fragmentary remains of a range of extinct fauna (including cf. Camelops sp., Canis dirus, and Bison sp.), as well as more mammoth bone recovered within an east trending alcove at roughly 6 m BMD (recorded as “level 18” in the IMNH catalog; Miller 1977). In 1977, these excavations also recovered seven fluted point fragments, a biface fragment, and a small amount of debitage from the alcove, reportedly within the same stratum as the Pleistocene animal remains.

The Terminal Pleistocene Chronology of Owl Cave

We begin by discussing the chronological placement of the terminal Pleistocene faunal material from Owl Cave, with a focus on the ages of the mammoth and bison in the collection. To do so, we collate and calibrate the available radiocarbon data from the Wasden site (Table 1) and compare them with the currently accepted ages for Clovis and Folsom. We calibrated all dates using the OxCal 4.2 software and the IntCal13 calibration curve (Bronk Ramsey 2009). Figure 3 presents the summed probability distributions (SPD) for the conventional radiocarbon dates on mammoth and bison run by the original excavators, new AMS dates on the same taxa, in addition to SPDs for Clovis and Folsom (raw data from Collard et al. 2010 and Surovell et al. 2016, respectively).

Immediately following the original excavations, Butler (1978; see also Miller 1989; Plew and Pavesic 1982) ran several radiocarbon dates on mammoth and bison bone collagen. Butler’s original dates on mammoth consisted of three assays, two of which he discarded because they were not pretreated to remove humic acid. A third treated sample returned a date of 13,090–12,593 cal B.P. Three dates on bison remain from the terminal Pleistocene deposits span the period 12,654–10,701 cal B.P. (Plew and Pavesic 1982). Taken together, these results suggest a Clovis age for the mammoth remains and a Folsom age for the bison. However, these dates were run over 30 years ago using conventional radiocarbon methods, with uncertainty about the preparation protocols employed, as well as relatively large sigma values. Fortunately, more recent AMS results exist for the Owl Cave faunal material.

Between 2012 and 2016, one dentin and six collagen samples from Owl Cave were AMS dated by various investigators (Table 1). These derive from one bison (Bison sp.) tooth, one Dire wolf (Canis dirus) bone, one camel (Camelops sp.) bone, and four different pieces of fragmentary mammoth bone. The SPDs derived from the Owl Cave mammoth and bison peak at different times. The oldest derives from the four mammoth specimens and represents a calibrated time span of 14,013–13,146 cal B.P. In contrast, the younger distribution derives from the bison tooth, spans the calibrated 2σ range from 12,594–12,404 cal B.P, and does not overlap with the dates on mammoth, wolf, and camel in a statistically meaningful way. In fact, the 95 percent confidence interval around the bison tooth is separated from the next youngest specimen (wolf) by 464 years and from the youngest mammoth date by 552 years.

The AMS data also suggest that the collection contains the remains of least two mammoths. In this case, the oldest and youngest mammoth dates fail to overlap at 2-sigma and, instead, at least 220 years separates the 95 percent confidence intervals of each measure. If we accept Butler’s mammoth date, a suggestion made with all due caution, then three mammoths might be present at Owl Cave, since Butler’s date is both younger than and fails to overlap at 2-sigma with the youngest of the new AMS dates on mammoth presented here. Further dating of these remains might resolve the number of individuals present.

Finally, the SPDs derived from the AMS dates on the Owl Cave fauna becomes even more interesting when compared with the age ranges for Clovis and Folsom. As shown in Figure 3, the mammoth remains largely predate the Clovis period, while the bison tooth falls squarely within the SPD describing Folsom. This latter finding is consistent with Butler’s earlier dates suggesting the introduction of bison remains into the cave during the Folsom period.
Table 1. Owl Cave Radiocarbon Dates.

<table>
<thead>
<tr>
<th>Lab No.</th>
<th>IMNH No.</th>
<th>Taxa</th>
<th>BMD</th>
<th>δ^{13}C</th>
<th>δ^{15}N</th>
<th>C/N</th>
<th>14C Age B.P.</th>
<th>Cal. Age B.P.</th>
<th>2σ Cal. B.P.</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>AMS UGAMS-25078</td>
<td>1642-1033</td>
<td>Bison</td>
<td>E5D</td>
<td>-19.0</td>
<td>5.8</td>
<td>3.2</td>
<td>10,530 ± 35</td>
<td>12,496 ± 56</td>
<td>12,594–12,404</td>
<td>This article</td>
</tr>
<tr>
<td>Beta-322085</td>
<td>10BV0030-186</td>
<td>Dire Wolf</td>
<td>E4E</td>
<td>n/a</td>
<td>n/a</td>
<td>n/a</td>
<td>11,280 ± 50</td>
<td>13,142 ± 50</td>
<td>13,251–13,058</td>
<td>IMNH</td>
</tr>
<tr>
<td>Beta-322086</td>
<td>10BV0030-244</td>
<td>Camel</td>
<td>E4C</td>
<td>n/a</td>
<td>n/a</td>
<td>n/a</td>
<td>11,500 ± 50</td>
<td>13,348 ± 53</td>
<td>13,456–13,252</td>
<td>IMNH</td>
</tr>
<tr>
<td>Aeon-1182</td>
<td>1642-22</td>
<td>Mammoth</td>
<td>E3D/E</td>
<td>-19.7</td>
<td>n/a</td>
<td>3.3</td>
<td>11,550 ± 110</td>
<td>13,382 ± 109</td>
<td>13,581–13,146</td>
<td>Steve Holen†</td>
</tr>
<tr>
<td>ISGS-A2303</td>
<td>10BV30D12-1</td>
<td>Mammoth</td>
<td>E3F/G</td>
<td>-19.6</td>
<td>n/a</td>
<td>n/a</td>
<td>11,650 ± 70</td>
<td>13,480 ± 75</td>
<td>13,610–13,300</td>
<td>This article</td>
</tr>
<tr>
<td>Beta-322088</td>
<td>10BV0030-979</td>
<td>Mammoth</td>
<td>E4E/F</td>
<td>n/a</td>
<td>n/a</td>
<td>n/a</td>
<td>11,840 ± 50</td>
<td>13,657 ± 59</td>
<td>13,765–13,555</td>
<td>IMNH</td>
</tr>
<tr>
<td>Beta-322087</td>
<td>10BV0030-256</td>
<td>Mammoth</td>
<td>E3E</td>
<td>n/a</td>
<td>n/a</td>
<td>n/a</td>
<td>12,000 ± 50</td>
<td>13,863 ± 77</td>
<td>14,013–13,738</td>
<td>IMNH</td>
</tr>
<tr>
<td>Conventional</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>WSU-2483</td>
<td>n/a</td>
<td>Bison</td>
<td>n/a</td>
<td>n/a</td>
<td>n/a</td>
<td>n/a</td>
<td>9735 ± 115</td>
<td>11,082 ± 190</td>
<td>11,404–10,701</td>
<td>Plew and Pavesic 1982</td>
</tr>
<tr>
<td>WSU-2485</td>
<td>n/a</td>
<td>Bison</td>
<td>n/a</td>
<td>n/a</td>
<td>n/a</td>
<td>n/a</td>
<td>10,145 ± 170</td>
<td>11,798 ± 310</td>
<td>12,390–11,250</td>
<td>Plew and Pavesic 1982</td>
</tr>
<tr>
<td>WSU-2484</td>
<td>n/a</td>
<td>Bison</td>
<td>n/a</td>
<td>n/a</td>
<td>n/a</td>
<td>n/a</td>
<td>10,470 ± 150</td>
<td>12,355 ± 165</td>
<td>12,654–12,050</td>
<td>Plew and Pavesic 1982</td>
</tr>
<tr>
<td>WSU-1786</td>
<td>n/a</td>
<td>Mammoth</td>
<td>n/a</td>
<td>n/a</td>
<td>n/a</td>
<td>n/a</td>
<td>10,920 ± 150</td>
<td>12,845 ± 129</td>
<td>13,090–12,593</td>
<td>Butler 1978</td>
</tr>
<tr>
<td>WSU-1259†</td>
<td>n/a</td>
<td>Mammoth</td>
<td>n/a</td>
<td>n/a</td>
<td>n/a</td>
<td>n/a</td>
<td>12,250 ± 200</td>
<td>14,337 ± 362</td>
<td>15,075–13,740</td>
<td>Butler 1978</td>
</tr>
<tr>
<td>WSU-1281†</td>
<td>n/a</td>
<td>Mammoth</td>
<td>n/a</td>
<td>n/a</td>
<td>n/a</td>
<td>n/a</td>
<td>12,850 ± 150</td>
<td>15,348 ± 256</td>
<td>15,846–14,804</td>
<td>Butler 1978</td>
</tr>
</tbody>
</table>

†Idaho Museum of Natural History specimen number.
‡Below main datum.
§Personal communication 2016.
¶Butler (1978) rejected these dates because they were not treated for humates.
Fluted Points and Mammoth Bone: Spatial Associations

Illustrating the spatial relationships between the fluted points and mammoth remains presents another avenue for understanding Owl Cave’s formational history. Figure 4 documents the excavation block in plan view. We generated this map from provenience data provided in the original notes, sketches, and catalogues (Miller 1977), and it highlights those units containing mammoth bone and/or flaked stone tools. Using the unit profiles drafted during the excavations, Miller produced a generalized stratigraphic profile of Owl Cave showing the overall depth of the deposits, as well as significant features including abundant ice wedge casts, the bison bone bed, and the stratum containing the fluted points and extinct fauna (Figure 5). Excavators took several photographs to document the fluted points in situ, but these show no other items, including bone, in close association with the points (Figure 6).

While the photographic evidence does little to aid in sorting out the relationships of interest, horizontal and vertical provenience to the nearest centimeter is available for both the projectile points and many of the mammoth remains. These data, however, key horizontally only to the 2 × 2 m excavation unit. The seven Folsom point fragments found in the cave distribute across...
five different excavation units (Table 2), with six of the artifacts located in the rear of the alcove. Also of note is the spatial distribution of conjoining fragments IMNH-76394 and IMNH-76437, which were separated horizontally by over two meters. Why the Folsom points are concentrated in the back of the cave is unclear, as it is a relatively cold and wet portion of the site today, but their locations might represent a toss zone into which broken, low-utility, and expended tools were discarded.

Figure 7 plots the vertical distributions of both the mammoth bone and the fluted points. These data show that mammoth bone distributes vertically from about 5.8 m BMD to basalt bedrock at about 6.5 m BMD. Of interest is the distribution of mammoth bone in Unit E4F. In this instance, mammoth specimens concentrate into two vertically discrete and dense groups of bone, suggesting two different depositional events. Unfortunately, only one specimen was dated (Beta-322088, 11,840 ± 40 B.P.; Table 1)
from this unit and it was sampled from the lower concentration.

Arraying the distribution of the fluted points against that of the mammoth bone indicates that both sets occupy the same range of vertical space. For example, Unit E5F produced at least 19 pieces of mammoth bone and two projectile points. In this case, the fluted points rest about 10 cm above all but two of the mammoth specimens, and about 10 cm below the piece of mammoth bone highest in Unit E5F. A broader comparison between the site-wide data for both groups finds that all but one of the fluted points (IMNH-76403, Unit E3D) fall within the vertical range of mammoth bone and the single outlier lay above it.

**Mammoth-Bone Surface Modifications**

A study of bone surface modifications provides another avenue for evaluating behavioral connections between humans and mammoth at Owl Cave, and ours is not the first study to evaluate the Owl Cave mammoth remains for butchery traces. Since many of the shaft fragments exhibit breakage and flaking to some degree, Miller’s (1983, 1989) analysis concluded that people had used many of the heavier cortical pieces as cores for the manufacture of sharp bone flakes. Miller also attributed some modifications to butchery for two reasons. First, long bones dominate the assemblage, suggesting to her that Pleistocene hunters transported specific elements into the site for marrow processing and tool production. Second, she saw the patterning and sequence of breakage as the intentional manipulation of the mammoth bone rather than the outcome of noncultural taphonomic processes (Miller 1983, 1989). Others, however, have expressed skepticism, as do we, both because of the absence of hammerstones (cf. Grayson and Meltzer 2002; Haynes 1991) and the potential for roof fall in Owl Cave to modify mammoth long bone in ways mimicking human butchery (Karr 2012). Because Miller’s (1983, 1989) analyses of the mammoth bone from Owl Cave generated some controversy, we inspected these specimens for
bone modifications. In this instance, however, instead of focusing on fracture patterns, we targeted surface modifications as a way to identify human agency in the creation of the mammoth bone assemblage.

We conducted our analysis of the Owl Cave mammoth assemblage at the IMNH during April 2014 (see DeCarlo 2017 for an exhaustive description of all surface modifications to the Owl Cave mammoth bone). We inspected the collection first with a 10× hand lens and then selected 46 specimens displaying modifications of interest. We examined these more closely at the Idaho State University Center for Archaeology, Materials, and Applied Spectroscopy (CAMAS) using an Olympus SZ61 stereo...
microscope at varying magnifications (6.7×–45×), angles, and lighting, resulting in the documentation of 102 surface modifications (DeCarlo 2017). To differentiate butchering marks from noncultural modifications resulting from, for example, carnivore gnawing or trampling, we compared the characteristics of each mark to a series of modification-specific rubrics derived from a review of replicative studies (Behrensmeyer et al. 1985; Blumenschine et al. 1996; Blumenschine and Selvaggio 1988; Bunn 1981; Dominguez-Rodrigo et al. 2009; Fisher 1995; Greenfield 2006; Haynes 1980, 1983, 1991; Potts and Shipman 1981; Redmond et al. 2012; Shipman 1981; Shipman and Rose 1983; Shipman et al. 1984).

Ninety-five (93 percent) of the 102 Owl Cave mammoth bone modifications possess features readily attributable to a causal agent (DeCarlo 2017). Of these, one (1 percent) modification represents trowel or shovel damage, two (2 percent) represent rodent modification, 16 (16 percent) are consistent with carnivore ravaging, and 76 (75 percent) modifications likely result from either trampling or, more likely, roof fall, which is consistent with the excavation photographs. In this study, we use “trampling marks” as a generic term for non-culturally, non-carnivore generated striations. The remaining seven (7 percent) modifications were initially classified as “anomalous” marks (Figure 8). Although intriguing, none of these modifications held up under scrutiny as unambiguous evidence for butchery. It is important to note, however, that experimental studies demonstrate that butchers can fully process an elephant with stone tools without leaving any surface modifications on the skeleton (Frison, 1989:778; Frison and Todd, 1986:42, 130; Haynes 1991:185–186). We offer a brief discussion of these anomalous marks here.

Roof fall best explains several of the anomalously marked specimens. IMNH-018 is a long bone shaft fragment exhibiting numerous abrasions and incisions (Figure 8a). Close inspection showed that the specimen possessed both a flake scar and associated abrasion that might document a cultural modification. The flake scar shows the removal of a broad flake, including a feather
Figure 8. Selected bone modifications to Owl Cave mammoth bone: (a) surface abrasion to IMNH-018, (b) surface abrasion to IMNH-210, (c) surface incisions to IMNH-719, (d) surface striations displayed by IMNH-195, (e) surface striation on IMNH-216, (f) surface striation on IMNH-216, (g) surface striation on IMNH-216.
termination and a negative bulb of percussion scar. At the striking platform, there is a short abrasion leading to the flake scar’s point of impact. The flake scar appears to result from a percussion impact, but such impacts can result from non-cultural processes. This is important since this piece of mammoth bone was one of several described in the IMNH catalog (Miller 1977) as resting among rock fall. Consequently, it is possible that the basalt boulders that landed on IMNH-018 may well account for the flake scar and associated surface abrasions (see also Karr 2012).

Specimen IMNH-210 is a mammoth rib fragment displaying numerous curved abrasions whose trajectories intersect each other (Figure 8b). However, IMNH-210 displays one anomalous mark, which is reminiscent of cultural percussion battering. This modification consists of an oval-shaped depression located within 5 mm from a fractured edge, a location suggestive of a butchery-related impact (Blumenschine et al. 1996:496). This mark also displays striations emanating from the depression as if a hammerstone came into gradual contact with IMNH-210, incising the surface in the process. While this modification may document percussion battering, the depositional context of IMNH-210 offers a competing, noncultural agent. Considering that excavators recovered IMNH-210 from under roof fall, it is possible that this abrasion was the point of impact between the bone and a falling basalt boulder. In addition, percussion battery of bone often results in many depressions or a collapsed platform (Haynes 1991:240), making the single depression inadequate evidence of human percussion.

IMNH-719 is a weathered long bone fragment that was also located below rock fall according to Miller (1977; see Figure 8c). This specimen displays one anomalous mark, which includes three, closely spaced parallel incisions. Under magnification, only the middle incision displays a consistent depth and width at its center, but then tapers as it terminates; features that might suggest butchery (Greenfield 2006:161; Redmond et al. 2012:87; Shipman and Rose 1983:75). In contrast, the right incision flattens and widens into multiple, shallow incisions, while the left one widens and gouges the bone, all features indicative of an agent such as trampling or roof fall (Dominguez-Rodrigo et al. 2009:2650). Due to their spatial context, these three marks are most consistent with sediment or some object forced across the bone during a single event.

IMNH-195 also displayed an anomalous striation (Figure 8d). In this case, an incision displaying several features consistent with a cut mark. Though interrupted by some type of puncture, the incision was straight on either side of the puncture, suggesting that the puncture mark was unrelated to the initial incision. This modification also displays an adjacent incision that briefly parallels the main one and, consequently, is reminiscent of a shoulder effect (Dominguez-Rodrigo et al. 2009:2646; Shipman and Rose 1983). Under closer inspection, however, it also possesses many features of a noncultural abrasion. For example, the base of the incision appears fairly wide and flat and the supposed shoulder effect does not change in its width when the main incision narrows. If this second incision was the result of the tilting of the cutting blade, its shape would change as the main incision changes, again suggesting that this is the result of a separate projection or granule being dragged across the bone’s surface.

IMNH-216 derives from a partial scapula and displays three anomalous surface modifications. Unmagnified, these modifications appear as parallel incisions consistent with cut marks. Under magnification, however, these markings possess features consistent with noncultural modifications. The most prominent of the three (Figure 8e) displays a flat incision base, a sinuous trajectory, and a crushed, rounded terminus, all features indicative of trampling (Dominguez-Rodrigo et al. 2009:2646, 2650; Redmond et al. 2012:87). Similarly, the second incision (Figure 8f) also possesses a broad incision base, while magnification revealed the third (Figure 8g) as a series of shallow incisions indicative of some type of abrasion.

**Projectile Point Protein Residue Analysis**

Several researchers have described the Owl Cave projectile points (Figure 9), with the consensus that they are morphologically consistent with the Folsom cultural complex best known from
the Great Plains and Southwest (Henrikson and Yohe 2009; Miller 1982, 1983, 1989; Titmus and Woods 1991; Yohe and Woods 2000). Seven of the projectile point fragments come from the interior of Owl Cave, while two additional specimens (IMNH-75503 and IMNH-76425) come from the surface in the immediate area of the cave entrance. Of these, four fragments conjoin into two unique specimens. Table 2 lists these specimens, their proveniences, and the obsidian source from which they originated.

To test for the presence of blood protein, we used crossover immunoelectrophoresis (CIEP; Culliford 1964, 1971; Dorrill and Whitehead 1979; Kooyman et al. 1992; Newman 1990, 1996; Shanks et al. 2001; Tuller and Saunders 2013). This method is a biomolecular/immunological technique that archaeologists have successfully used to identify protein residues on flaked stone, ground stone, and ceramic artifacts (e.g., Barnard et al. 2007; Hogberg et al. 2009; Newman and Julig 1989; Newman et al. 1993; Yohe et al. 1991). Moreover, archaeologists have used this technique in the analysis of Paleoindian material, including fluted points, resulting in the identification of horse, camel, elephant, and reindeer protein (Duke 2015; Gilbert et al. 2008; Kooymen et al. 2001; Parr 2006; Seeman et al. 2008; Yohe and Bamforth 2012; Yost 2013).

CIEP provides results by testing protein residues for reactions to various taxa specific antisera. In this instance, we tested the Owl Cave projectile points against a broad range of antisera expected to represent the suite of prey animals potentially available to SRP foragers. In all but one instance, the antisera used in this study were acquired commercially (Table 3). Given that CIEP requires the use of antisera specific to taxonomic families, and no antisera for elephants was readily available, a custom antiserum for elephants was produced from fresh, whole blood extracted from a modern Asian elephant (Elephas maximus) housed at the Fresno Zoo in Fresno, California, by LAMPIRE Biological Laboratories.

The nine fluted point fragments from Owl Cave were submitted for immunological analysis to the Laboratory of Archaeological Sciences (LAS) at California State University, Bakersfield (Henrikson and Yohe 2009). Yohe removed potential residues from the artifacts as discussed above and recorded three positive reactions (Table 2). IMNH-75503 tested positive for sheep indicating the presence of proteins from any species of sheep or goat. However, because this
Table 3. Antisera Used in This Study.

<table>
<thead>
<tr>
<th>Antiserum</th>
<th>Source</th>
<th>Reacts With</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bear</td>
<td>Cappel Research</td>
<td>Black, grizzly, etc.</td>
</tr>
<tr>
<td>Bovine</td>
<td>Cappel Research</td>
<td>Bison, cow</td>
</tr>
<tr>
<td>Camel</td>
<td>Sigma-Aldrich</td>
<td>All camelids</td>
</tr>
<tr>
<td>Cat</td>
<td>Cappel Research</td>
<td>Bobcat, cougar, lynx, etc.</td>
</tr>
<tr>
<td>Chicken</td>
<td>Sigma-Aldrich</td>
<td>Quail, grouse, and other gallinaceous fowl</td>
</tr>
<tr>
<td>Deer</td>
<td>Cappel Research</td>
<td>Deer, elk, moose</td>
</tr>
<tr>
<td>Dog</td>
<td>Cappel Research</td>
<td>Coyote, dog, wolf</td>
</tr>
<tr>
<td>Elephant</td>
<td>Lampire</td>
<td>Elephant and extinct relatives</td>
</tr>
<tr>
<td>Guinea Pig</td>
<td>Sigma-Aldrich</td>
<td>Beaver, marmot, porcupine, squirrel</td>
</tr>
<tr>
<td>Horse</td>
<td>Cappel Research</td>
<td>Donkey, horse, kiang, etc.</td>
</tr>
<tr>
<td>Rabbit</td>
<td>Cappel Research</td>
<td>Rabbit, hare, pika</td>
</tr>
<tr>
<td>Rat</td>
<td>Cappel Research</td>
<td>All rat and mouse species</td>
</tr>
<tr>
<td>Sheep</td>
<td>Cappel Research</td>
<td>Bighorn and other sheep, goats</td>
</tr>
</tbody>
</table>

point fragment is an unprovenienced surface find and sheep have pastured there in recent history, this result may document modern contamination from urine or feces. IMNH-76436 tested positive for horse. Horse antiserum will react positively with species of horse or donkey, and we assume that extinct equids would also test positive. IMNH-76438 tested positive for elephant, indicating the presence of proteins from some related species, extinct or extant. The negative reactions for identifiable proteins on the remaining artifacts may result from poor preservation resulting in loss of antigenicity, insufficient trace protein quantities, or the possibility that the artifacts never contacted blood or any other proteins from any of the taxa included in this study. Finally, we would normally process associated soil samples as contamination controls. While the IMNH houses several soil samples, none came from the same units as projectile points, leaving their utility as controls dubious at best, and no such analysis is presented here.

Discussion

The analyses presented above fail to associate the fluted points with the mammoth remains in a way signaling mammoth hunting by Folsom foragers. While some of the data—AMS dates, for example—suggest one narrative, other information, such as the protein analysis, suggest another. Simply put, the dated mammoth remains predate Clovis, let alone Folsom, yet the Folsom points from Owl Cave were found mixed in with the mammoth remains and tested positive for elephant and horse protein residues. Faced with this quandary, we envision several competing formational histories that each might explain the composition of the Owl Cave terminal Pleistocene materials.

First, the fluted points and at least some of the mammoth remains associate directly through a hunting event. If true, and if the points represent a Folsom occupation, then, depending on whether or not one accepts Butler’s mammoth date, either (1) Folsom predates Clovis on the SRP, (2) Folsom was contemporary to Clovis on the SRP, or (3) the fluted points from Owl Cave represent a morphologically similar but earlier and otherwise unrelated technology to the Folsom phenomenon (see Beck and Jones 2010 for a related discussion of Western Fluted points). In this latter instance, “Folsom” becomes a definitional issue and one that no longer refers to a distinctive point style, but instead a chronological interval.

This scenario is problematic since widely accepted cases of mammoth hunting from the Great Plains and Southwest associate exclusively with Clovis points, which predate Folsom in every context in which they have either been dated or found in a buried, stratigraphic sequence. We are further troubled by the lack of documentation for direct associations between the Owl Cave projectile points and mammoth bone, the absence of horse remains, the presence of Folsom-age bison remains in the same deposit, the lack of cultural bone-surface modifications, and the absence of “Folsom” points from any other securely dated pre-Clovis or Clovis-period contexts. Such concerns lead us to explore alternative explanations for the Owl Cave materials.

A substitute, but related, explanation accommodating a direct contextual association between fluted points that are chronologically Folsom and mammoth remains would suggest that Owl Cave documents a rare example of Folsom predation on lingering populations of horse and...
mammoth. Consequently, Owl Cave may contain a Folsom-age mammoth that was not dated due to sampling error. Conversely, the extinct mammal remains from Owl Cave and the Folsom points may be unrelated, yet the protein residue accurately records Folsom-age hunting, scavenging, or other uses of mammoth and horse products, without any carcass parts from those events having been transported back to the cave. While these scenarios are not impossible, they are improbable, because such circumstances would suggest an undated and previously unrecognized remnant population of Pleistocene mammals in the northern Great Basin and Snake River Plain.

The Folsom points and mammoth remains may associate contextually in another way, one not requiring the contemporaneous presence of the two populations. In this scenario, much later Folsom foragers modified frozen or otherwise well-preserved mammoth bones into bone tools. Important to this scenario is recent research indicating that very old, but well-preserved, bone can break in ways fully consistent with the expectations for green-bone fractures. Haynes (2016), for example, documents recent green-bone fractures caused by earthmoving equipment to mammoth bone at the Ingelwood mammoth site.

Recognizing that old bone can break in such ways accommodates many of the conflicting lines of evidence discussed above. In this scenario, Miller (1989) could well be correct in suggesting that the fragmentary nature of the mammoth bone represents Pleistocene foragers using long bone diaphyses as cores for tool production, but in this case at the hands of Folsom, not Clovis or pre-Clovis, foragers. This scenario also suggests why the mammoth remains do not display any unambiguously cultural surface modifications. Simply put, there would have been little or no soft tissue to remove. Nor would this scenario require the improbable assertion that Folsom peoples were hunting mammoth at a time when the available data indicate that those animals were extinct. Instead, Folsom foragers may have sheltered in the cave and produced bone tools from well-preserved mammoth remains. Along the way, these visiting foragers would have deposited several broken projectile points as part of ordinary tool maintenance activities, contaminating these artifacts with persisting mammoth protein in the process. We note, however, that this scenario fails to explain the presence of horse protein on one of the Folsom points.

Finally, the fluted points and megafauna may represent completely unassociated events and the protein residue results reflect contamination from noncultural animal remains residing in the deposits. Cryoturbation represents one taphonomic process that might have contaminated the fluted points with otherwise unassociated protein. According to Dort (1968:33), the formation of ice wedges in Owl Cave signified extended periods of extreme cold (with ground temperatures consistently reaching −15°C). Based on the size of the largest Owl Cave ice wedge casts, Dort surmised that these intense cold periods may have lasted for several hundred years. The most significant ice wedging events are bracketed by the bison bone bed and the roof fall episode above the cave floor, with evidence for wedging in the lowest deposits as well.

Although not captured in the schematic profile, sediments near the ice wedge casts and below the roof fall in the alcove are characterized by “involutions,” which, according to Dort (1968:34), were likely the result of continuous seasonal freezing and thawing. Consequently, cryogenic distortion of the sediments could have led to the vertical movement of both animal protein and stone artifacts in the lower levels of the deposits, eventually leading to the contamination of the fluted points. Cryoturbated sediments have been noted in other Idaho lava tubes (Henrikson and Long 2007), and in at least one case, Scaredy Cat Cave, decomposed, frozen bison meat was recovered from deposits dating to roughly 4000 cal B.P. (Henrikson 2002), suggesting that this contamination is not impossible. Yet, this scenario is problematic because no horse remains are known from the cave, and experimental data suggesting that protein contamination can occur through cryoturbation are lacking.

Conclusions
In the end, our study of the Owl Cave materials fails to establish a case for mammoth hunting at the site. Instead, one can read the available data in several different ways, leaving us with little more
than a number of largely untestable suppositions about site formation processes at Owl Cave. Consequently, we present several competing scenarios illustrating how the terminal Pleistocene deposits could have formed.

To be clear, we take no position at this time on what Owl Cave truly documents, and instead have tried to give an even hand to discussing the various ways one might read the terminal Pleistocene materials from the site. It is apparent to us that Owl Cave contains a palimpsest of events. While our research efforts thus far have focused strictly on interpreting existing data from the lowest deposits, the bison bone bed has been, once again, neglected. However, a recent study (Guenther 2014) does suggest that the assemblage of bone and Agate Basin–style projectile points reflect an early Holocene bison drive. Based on Guenther’s (2014) analysis, additional bison remains are likely present in adjacent Coyote and Dry Cat Caves.

Although we are not compelled to accept any of the possible scenarios presented here, future research at the Wasden site has tremendous potential to resolve long-standing ambiguities in the existing documentation. The original excavators focused on deposits deep inside the alcove of Owl Cave; they did not excavate sediments in the cave’s sunny, western portion, possibly leaving half of the bison bone bed intact. Likewise, if the fragmentary points in the alcove represent a discard zone, open areas just inside the drip line may contain more substantial evidence of a terminal Pleistocene occupation associated with fluted points, the only one of its kind in the desert West. In the near future, we hope to excavate the remaining deposits with modern methods designed to capture the fine-grained spatial data needed to properly evaluate associations between extinct animals and the material residues of the early peoples inhabiting the SRP.

Acknowledgments. The authors thank the numerous individuals who assisted in preparation of this manuscript. Susanne J. Miller and Wakefield Dort provided essential information and invaluable institutional knowledge on the original excavations. Lynn Murdock, Amy Commendador-Dudgeon, and Amber Tews of the Idaho Museum of Natural History allowed access to the collections and original field notes, and provided the loan of Owl Cave materials for these analyses. Ruthann Knudson graciously provided her illustrations of the fluted points. Wakefield Dort, the Idaho Heritage Trust and IMNH funded the new accelerator dates and Craig Skinner completed the XRF analysis in a professional and timely manner. Norman Henrikson of the Laboratory of Archaeological Science at CSUB conducted the immunological analysis of the Owl Cave points. Marissa Guenther, graduate student at CSUB, assisted with the illustrations. Shaunzi the elephant, assisted by the friendly staff from the Fresno Zoo, provided a blood sample in the generation of antisera for this study. Arantza Zubala translated the English abstract into Spanish. Lastly, we would like to thank Steve Croft (former Owl Cave landowner) for allowing unrestricted access to the site and the Archaeological Conservancy for recognizing the significance of the property and pursuing its acquisition.

Data Availability Statement. The original field notes and associated documentation used to support the findings of this study are available at the Idaho Museum Natural History, Pocatello. Recently generated AMS assays, as well as the reports associated with the CIEP and XRF analyses, are available at the Cultural Resource Management Office, Idaho National Laboratory, Idaho Falls.

References Cited

Armstrong, L., Trent, L., Suzann Henrikson, Hollie K. Gilbert, and Brenda R. Pace


Barnard, Hans, Lori Shoemaker, Micala Rider, Oliver E. Craig, Robert E. Farr, Mark Q. Sutton, and Robert M. Yohe II


Beck, Charlotte, and George T. Jones


Behrensmeyer, Anna K., and Susan M. Kidwell


Blumenschine, Robert J., Curtis W. Marean, and Salvatore D. Capaldo


Blumenschine, Robert J., and Marie M. Selvaggio


Bronk Ramsey, Christopher


Bunn, Henry T.

Butler, B. Robert

Butler, B. Robert, Helen Gildersleeve, and John Sommers

Cannon, Michael D., and David J. Meltzer

Collard, Mark, Briggs Buchanan, Marcus J. Hamilton, and Michael J. O’Brien

Culliford, Bryan J.

DeCarlo, Matthew M.

Dominguez-Rodrigo, Manuel, S. de Juana, Ana B. Galán, and M. Rodríguez

Dorrill, Marion, and P. H. Whitehead

Dort, Wakefield, Jr.

Duke, Daron

Fisher, John W., Jr.

Frisson, George C.

Frisson, George C., and Lawrence C. Todd
1986 *The Colby Site*, University of New Mexico Press, Albuquerque.


Grayson, Donald K., and David J. Meltzer

Greenfield, Haskel J.

Guenther, Marissa Anne

Haynes, Gary


Henrikson, Norman D., and Robert M. Yohe II
2009 *Protein Residue Analysis on Nine Folsom Points from Archaeological Site 10BV30*, California State University, Bakersfield, Laboratory of Archaeological Science Report No. 219. Submitted to the Center for Archaeological Research, California State University, Bakersfield, California. Copies available from the Laboratory of Archaeological Science, California State University, Bakersfield.

Henrikson, L. Suzann

Henrikson, L. Suzann, and Montana M. Long

Hogberg, Anders, Kathryn Puseman, and Chad Yost

Karr, Landon P.
Kooyman, Brian, Margaret E. Newman, and Howard Ceri

Kooyman, Brian, Margaret E. Newman, Christine Cluney, Murray Lobb, Shayne Tolman, Paul McNeil, and L. V. Hills

Long, Montana M.

Miller, Susanne J.


Miller, Susanne J., and Wakefield Dort Jr.

Newman, Margaret E.


Newman, Margaret E., and Patrick J. Julig

Newman, Margaret E., Robert M. Yohe II, Howard Ceri, and Mark Q. Sutton

Parr, Robert

Plew, Mark G., and Max G. Pavesic

Potts, Richard, and Pat Shipman

Redmond, Brian G., H. Gregory McDonald, Haskel J. Greenfield, and Matthew L. Burr


Shanks, Orin C., Robsin Bonnichsen, Anthony T. Vella, and Walt Ream

Shipman, Pat

Shipman, Pat, Daniel C. Fisher, and Jennie J. Rose

Shipman, Pat, and Jennie Rose

Surovell, Todd A., Joshua R. Boyd, C. Vance Haynes Jr., and Gregory W. L. Hodgins

Titmus, Gene L., and James C. Woods

Tuller, Hugh, and Rebecca Saunders

Yohe, Robert M., II, and Douglas Bamforth

Yohe, Robert M., Margaret E. Newman, and Joan S. Schneider
Yohe, Robert M., II, and James Woods

Yost, Chad

Submitted January 23, 2017; Revised March 8, 2017; Accepted March 11, 2017