

Shallow Subsurface Evidence for Postglacial Holocene Lakes at Ivanpah Dry Lake: An Alternative Energy Development Site in the Central Mojave Desert, California, USA

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Abstract

Dry lakes, or playas, in the American Southwest have become important locations for alternative energy solar electric generation systems. Many of these playas have not been investigated for past environmental conditions prior to their development, causing alterations to the shallow sediment profile and altering subsurface chemical and physical data important to recent geologic and hydrologic histories of these areas. Recent studies of these features commonly have focused on surficial data with little or no subsurface geologic data to delineate wet and dry cycles of playa lakes. Many of the playa margins are covered with a surficial layer of alluvial and/or aeolian sands varying in thickness, which could interfere with correct interpretations of data gathered only on the surface material on these lands. Over the past 13,000 years Ivanpah Dry Lake has had at least three episodes of lake highstands up to ~10m in response to climate changes, based on data in this study with the oldest episode dates to ~13,000 cal yrs BP. This study has a link to other sites as many other playas are slated for energy development and could endure subsurface data alteration during site reworking and occupation, resulting in a loss in their collective historical geologic and hydrologic records if more complete studies of important sites are not carried out before developments begin.

Keywords: Ivanpah, pluvial lakes, playa, Lake Mojave, Holocene, Pleistocene, lacustrine, California

1. Introduction

Desert dry lakes and their margins in the American Southwest and elsewhere are considered potential sites for the development of alternative energy projects to supplant current energy sources that adversely affect climate, and climate change awareness in action policies by potentially affected institutions. Many of these areas are located within regions where there has been little focus into historical environmental conditions. In response to increased alternative energy needs, there are strategies in place to create solar energy zones that encroach onto and could disturb buried deposits across the deserts of the American Southwest, including Ivanpah Dry Lake (Fig. 1) (“Lake Ivanpah”). These developments are impacting the very sedimentological records that could provide information on the historical environmental conditions, and potential future environmental impacts. In other words, it seems that the vast magnitude of paleohydrologic changes that can occur in these basins is under-appreciated when facilities performance over decades are considered.

Understanding long-term environmental change in the arid Southwest rests in part on paleohydrologic studies, mostly of the Quaternary Period, pluvial lakes fed by rivers originating from beyond the desert (e.g. Blunt and Negrini, 2015; Enzel et al., 2003; Litwin et al., 1999; Nishizawa, 2010; Negrini et al., 2006; Wells et al., 2003), and paleospring deposits that reflect more local histories of recharge within the Great Basin and Mojave deserts (Quade et al., 1995; 2003). Such records of hydrologic responses to precipitation allow for greater delineation of the effects of changing climate. The effects of these changes have been profound on both surface processes and biotic communities in the region (e.g., Bull 1991; Spaulding 1990).

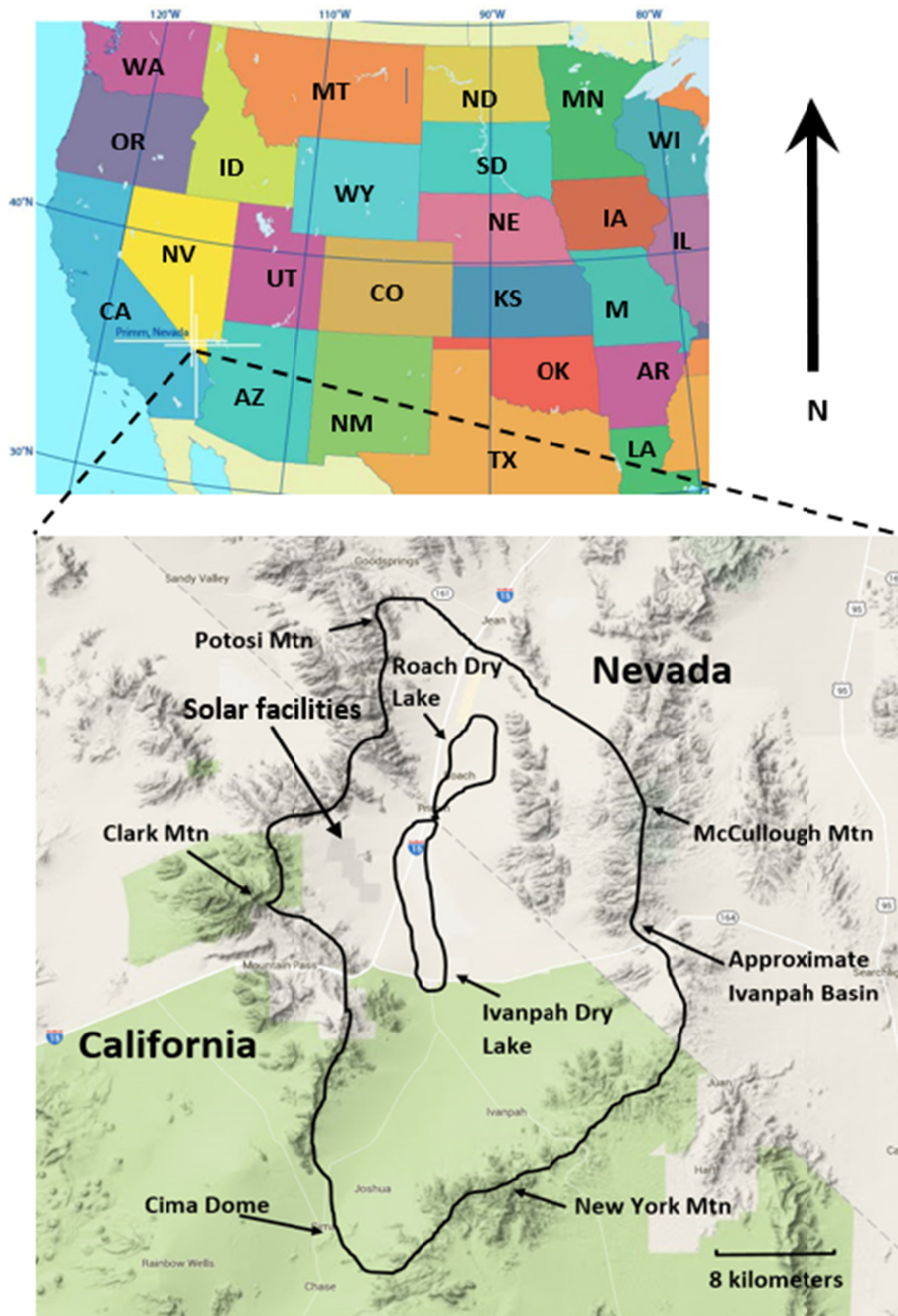


Figure 1. Location of Lake Ivanpah, California and approximate Ivanpah hydrologic basin

Desertification of the American Southwest followed the retreat of the continental ice sheets in the Northern Hemisphere, as an increase in thermal regimes accompanied a northward shift in circulation patterns to those characteristic of the present arid Southwest (Thompson et al., 1993). There have been studies in this region of late Quaternary Period precipitation and its effects on groundwater, climate change, and pluvial lake systems

(Quade et al., 1995; Pigati et al., 2011; Waring, 1920; Wells et al., 2003). Other research in this area evaluated Pleistocene Camelid tracks and aeolian distributions as they relate to climate change at Mesquite Lake, California, a dry lake system located 16.5 km northwest of Primm, Nevada (Shapiro, 2016; Whitney et al., 2015). Finally, environmental variability within this region, and more notably, Ivanpah Valley, is complex.

To better appreciate the environmental changes inherent to Lake Ivanpah, it is important to recognize that episodes of higher lake stands and increased paleospring discharge, occurred not only during the latest Pleistocene, but also during the succeeding Holocene (e.g. Wells et al., 2003; Noble et al., 2016; Quade et al., 1995; Pigati et al., 2011). The Pleistocene to Holocene transitional period experienced reversals in the desiccation of ancient lakes and extinct spring systems (“paleolakes” and “paleosprings”). So, while it has been reported that the “final” desiccation of paleolakes and ancient springs occurred between ca. 9,300 and 7,800 calibrated years BP (cal yrs BP), essentially concordant with the extirpation of mesic vegetation in current deserts (Wells et al., 2003; Pigati et al., 2011; Spaulding and Graumlich, 1986), there were important reversals during the succeeding postglacial. According to Noble et al. (2016), pluvial events of sufficient magnitude to fill some basins for decades or centuries occurred during the middle and late Holocene. Refilling of basins along the Mojave River reportedly occurred at least twice: ca. 3,600 yrs BP and again during the Little Ice Age, ca. 600 to 200 yrs BP (Enzel et al., 1992; Tchakerian and Lancaster, 2002). Other studies documented that the late Holocene Period witnessed a rebirth of paleospring systems throughout this region (e.g. Mehringer and Warren, 1976; Jones et al., 1999).

There has been comparatively little research on dry lakes with drainage basins lying entirely within the Mojave Desert. Unlike lakes Owens and Searles, on the Owens River that heads in the eastern Sierra Nevada, or Lakes Manix and Mojave on the Mojave River that heads in the Transverse Ranges, isolated dry lake basins show few geomorphological signs of highstands (i.e. wave-cut benches), fluvial deposits and/or terraces (Mifflin and Wheat, 1979; Reheis and Redwine, 2008). Comparing the apparent absence of pluvial lakes in the southern Great Basin to their relative abundance and great size in the north, Mifflin and Wheat (1979) developed a model relating the combined effects of an increased thermal regime in the south, lower elevations, and smaller mountains to insufficient runoff to support persistent lakes, even during glacial conditions. As a result, little research has focused on the paleohydrology of internal dry-lakes of the southern Great Basin. Studies have shown that the mere absence of surface evidence does not preclude the possibility of ancient lakes. Rather, studies require interpretation from the examination of aeolian dunes, infrared photographs, subsurface conditions, grain size, and sediment chemistry-proxy data to identify buried remnants of ancient lake beds (Sack, 1995; 2001).

2. Environmental Setting

Three solar electric energy generation projects are currently being built, or operational, in the Ivanpah Valley in the last five years. In order of their proximity (farthest from) to the current playa margin are: Stateline Solar Farm Development on the northeastern margin of Lake Ivanpah, on the toe of the Clark Mountain alluvial fan in California; Silver State and Silver State South Solar Energy Centers, east of Roach Dry Lake, northern portion of the Lucy Gray alluvial fan toe in Nevada; and Ivanpah Solar Electric Generating System, located further up the Clark Mountain alluvial fan. The Ivanpah facility is currently the world's largest concentrated solar thermal power station in operation while the other two facilities are large solar photovoltaic systems (CH2MHILL, 2009).

2.1 Hydrologic Setting

Although the Ivanpah basin is not part of the Mojave River drainage (see Enzel et al., 2003), tributaries to the lower Mojave River lie only ~25 km to the west in Shadow Valley. Approximately 20 km to the east on the other side of the McCullough Mountains, the Paiute Valley drains into the Colorado River. The Ivanpah basin incorporates a drainage network extending from the Teutonia Peaks area on Cima Dome, 25 km to the southwest in California, to Potosi Mountain, about 40 km to the east-northeast in the Spring Mountains of Nevada (Fig. 1). The watershed also includes the north flank of the New York Mountains and Mid Hills to the south and southwest. The Ivanpah Range, Mescal Range, and Clark Mountain provide runoff from the west, and, the Clark Mountain Range and southern-most Spring Mountains provide runoff from the north.

The Ivanpah basin also includes Roach Dry Lake, immediately to the northeast in Nevada, separated from Ivanpah playa only by the toe of the alluvial fan extending west from the Lucy Gray Mountains. Because this permeable sill is the only separation between the two playas, Roach Dry Lake and its drainage are part of the Ivanpah hydrologic system. Other studies of the hydrological processes of Lake Ivanpah have determined that Ivanpah Valley is entirely internally drained (House et al., 2010). Lake Ivanpah's surface drainage in California

is approximately 1,128 km²; its catchment in Nevada is approximately 831 km², for a total drainage basin area of 1,959 km² (756 mi²). The surface area of Ivanpah playa is about 53 km² while Roach playa is approximately 15 km².

2.2 Topographic Setting

Ivanpah playa has a sinusoidal shape that is the result of two large alluvial fans intruding on its margins from the northeast and from the west; i.e. Lucy Gray and Wheaton Wash alluvial fans (Figs. 1-3). Both alluvial fans create a convex-shaped shoreline at their terminal edges. The arcuate margin of Lucy Gray Fan terminal edge projects west and extends for ~5 km south of the sill it creates, terminating at the playa in the north. Approximately 10 km south, on the opposed side of the playa, the toe of the Wheaton Wash Fan forms an eastward projecting convex shoreline. Fans are composed of moderately sorted, well stratified sands and gravels of pebble and boulder sizes with minor amounts of aeolian silts and sands at depth, with an estimated age of deposits of ~2.6 million years according to House et al. (2010). Both the Lucy Gray and Wheaton Wash Fan are lacking active fluvial processes, with recent flows confined to ephemeral systems nearer the source of materials (House et al., 2006; 2010).

The relief of the topography encompassing a basin's drainage is a critical factor in determining runoff; i.e. the higher mountains cause greater amounts of orographically induced precipitation (Mifflin and Wheat, 1979). Studies of Mojave River runoff for instance, show that water contained in Lake Mojave originated chiefly from surrounding mountains outside the Mojave Desert (Enzel et al., 2003). With the Mojave River in mind, the Ivanpah drainage basin includes some of tallest mountain ranges in the central Mojave Desert (i.e. New York Mountains [~ 2,296 m], Clark Mountain [~2,417 m], Potosi Mountain in the southern Spring Mountains [~2,595m], and ~2,142 m for McCullough Mountain) and therefore, runoff that entered the Ivanpah basin originates from the surrounding mountains.

The barren part of Ivanpah Basin's elongate and sinusoidal playa occupies only the northern 55% of the 19.3 km-long basin floor. Generally not more than 3 km wide, the playa extends south about 11 km from the vicinity of Primm, Nevada, to an ill-defined southern limit at about the 795 m contour. The barren northern and central portions of the playa are mapped as essentially level at ~794 m amsl (USGS, 1985). The playa surface continues farther south, but is mantled by an increasingly dense cover of saltscrub hummocks to the south. These hummocks, usually not exceeding a meter in height, consist of grey-brown to pale brown (10YR7/4), silty fine sand anchored by saltscrub vegetation, and typified by Allscale (*Atriplex polycarpa*), a species of saltbush. South of the Nipton Road, blow-outs that have stripped back the mantling saltscrub hummocks reveal a mosaic of playa surface and carbonate-rich paleospring outcrops armored by nodular tufa concretions (Robinson et al., 1999). Saltscrub rims the entire playa except for its north end, and it gives way upslope to a creosote bush (*Larrea tridentata*) - Burrobush (*Ambrosia dumosa*) scrub transition zone.

Moving south, the floor of Ivanpah playa rises in elevation well before its southern end is reached. Because the surface of Ivanpah playa is not level, evidence of shallow water in the south indicates deeper water to the north. Previous research near Nipton Road identified oxidized lake muds in yardangs at 800.1 m amsl, only ~0.9 m above the adjacent playa but >6 m above the elevation of the pan in the north (Robinson et al., 1999). Their investigation found bedded, sandy alluvium yielding abruptly to a charcoal-rich, dark-grey lake mud at ~1 m depth. The top of this lake mud was surveyed at 799.9 m amsl, again ~6 m above the playa floor in the northern Ivanpah basin, yielding an age of 9,560 ± 70 ¹⁴C yrs BP (Robinson et al., 1999). Assuming inundation is needed for playa-mud deposition, and that no substantial tectonic warping of the basin floor has occurred, their study indicated that a lake of at least moderate depth occupied the northern Ivanpah basin.

There is no abrupt southern boundary to Ivanpah playa. Rather, it is characterized by an increasingly dense cover of saltscrub hummocks and tongues of gravelly alluvium, the latter from the broad delta-fan expanding onto the basin floor from the south at the mouth of Cima Wash. The name Cima Wash was applied to the system of anastomosing sandy washes that extends ~25 km south and west from the southern playa margin to the southeast flank of Cima Dome. As this broad drainage runs down the southwestern Ivanpah Valley, it is joined by washes draining the north flank of the New York Mountains and Mid-Hills. Remote imagery and field assessment places the distal toe of the Cima Wash delta-fan, and the southern limit of Ivanpah playa, ~1 km south of Nipton Road.

2.3 Ivanpah Playa

The location of Lake Ivanpah is a topographic depression at the base of the Ivanpah, Mescal, and Clark Mountain ranges, just west of the Nevada state line. The study area is hot and dry with temperatures fluctuating between ~1°C in winter (January/December) to >41°C in summer (July/August). Precipitation averages <11.5 cm annually near the basin bottom and higher in the surrounding mountain ranges (Robinson et al., 1999).

Geographically, this relatively high basin (playa surface is ~ 794 m above mean sea level [amsl]) is located within the Mojave Desert. Roach Dry Lake is connected to the Ivanpah playa at the border of Nevada and California (Primm, NV) and is part of the Ivanpah Catchment Basin. It lies at ~794 m amsl and is separated from Ivanpah by a narrow sill produced by the merger of the Lucy Gray alluvial fan and a limestone ridge extending south from the Spring Mountains Range (Carr and Pinkston, 1987; Garside et al., 2009). The Ivanpah catchment in California is approximately 1,128 km² with the Nevada portion being ~831 km² (totaling ~1,959 km²) with Ivanpah playa covering ~53 km² and Roach playa ~15 km².

Lacustrine sediments have been described as having distinct lithological units with defined beds (laminated) of reworked material containing ubiquitous amounts of carbonates, much greater than sediment organic carbon, with large amounts of soluble salts (Eugster and Kelts, 1983). Other authors have defined lacustrine sediments as originating from incoming sediments due to wetting events, depositing sediments in an aqueous environment which in turn becomes a playa surface with aridity (Kliem et al., 2012; May et al., 2015). Thus, the superficial portion of a lacustrine unit will appear as a classic playa surface, the upper most portion of a lacustrine unit according to Kliem et al., (2012).

2.4 Ancient Shorelines and Lacustrine Features

Identifying, correlating and mapping shoreline features in an area where the climate has been relatively stable since the end of the Pleistocene can be challenging (Sack, 1995). Shorelines usually leave demarcations (“highstands”) of their farthest advance and later, may be buried or obliterated by successional advances but leaving visible geomorphic evidence (Adrian et al., 1999). Ivanpah has no such obvious preserved visible lake highstands as describe by other authors (i.e. Negrini et al., 2006; Enzel et al., 1992; Wells et al., 2003). Studies have however, documented the reworking and/or burial of shorelines and their associated lacustrine sediments by active alluvial fans and evidence of fluvial and aeolian processes that obliterate and bury geomorphic evidence making them difficult to locate surficially with aerial photographs, topographic maps and ground reconnaissance activities (Sack, 1995).

There was surficial geological mapping of portions of this basin that identified no obvious ancient shorelines (House et al., 2010; 2006; Ramelli et al., 2006; Schmidt et al., 2006). These studies however did map “playa fringe deposits” to elevations >10m above the current playa margin. Visual evidence in the form of shorelines is only one factor in pluvial lake demarcation: soils, sedimentologic, sediment chemistry, subsurface stratigraphy, and geomorphic evidence also can be used to identify pluvial lakes lacking obvious shorelines (Sack, 1995; 2001; Schnurrenberger et al., 2003). Current efforts do not contradict prior mapping efforts (House et al., 2010, 2006; Ramelli et al., 2006; Schmidt et al., 2006), but are complementary to them. However, because the possibility of a pluvial lake having occupied the basin is absent from their interpretive geological framework (House et al., 2010, 2006; Ramelli et al., 2006; Schmidt et al., 2006.), and because their work was restricted to the Nevada portion of the basin, those data sets have limited transfer value to this effort.

Within the Ivanpah basin nonetheless, there are yardangs, loose and un-weathered surface sand, strath surfaces (Fig. 2A), hummocks (Fig. 2B), both modern (Fig. 2C) and ancient (Fig. 2D) beach surfaces, north-south trending low lying aeolian dunes (Fig. 2E) with discontinuous sand sheets, and exposed lacustrine sediments just off the perimeter of the playa margin. These features are subtle evidence of preserved landforms that could easily be overlooked by analysis of aerial photography and topographic map review, even supported with some ground reconnaissance. This research does not map geomorphic evidence on the surface, but rather, evaluates subsurface conditions at one location on the eastern side of Lake Ivanpah.

3. Objectives

The purpose of this research is to demonstrate that a large (depth >5 m) body of water episodically occupied the basin in lieu of a pluvial lake, and expanded well beyond the edge of the current playa as supported predominantly by data gathered from subsurface sediments. To do this, subsurface conditions on the playa margins were investigated using sedimentary stratigraphy and chemical proxies to identify subsurface anomalies potentially related to lake transgressions over alluvial fan toes. These chemical proxies (metals, SO₄²⁻, Cl⁻, CaCO₃, pH, Mg/Ca ratios, radiocarbon dating (¹⁴C), δ¹³C, and δ¹⁸O) were used to locate a suitable location for further excavation so that subsurface sedimentological, geomorphic, and lacustrine event evidence could establish chronology, and be used to describe the magnitude of environmental change within the Ivanpah basin. Data was comparable with records of lake-level changes from other paleolakes, particularly those that received most of their runoff from mountains both inside and outside the Mojave Desert.

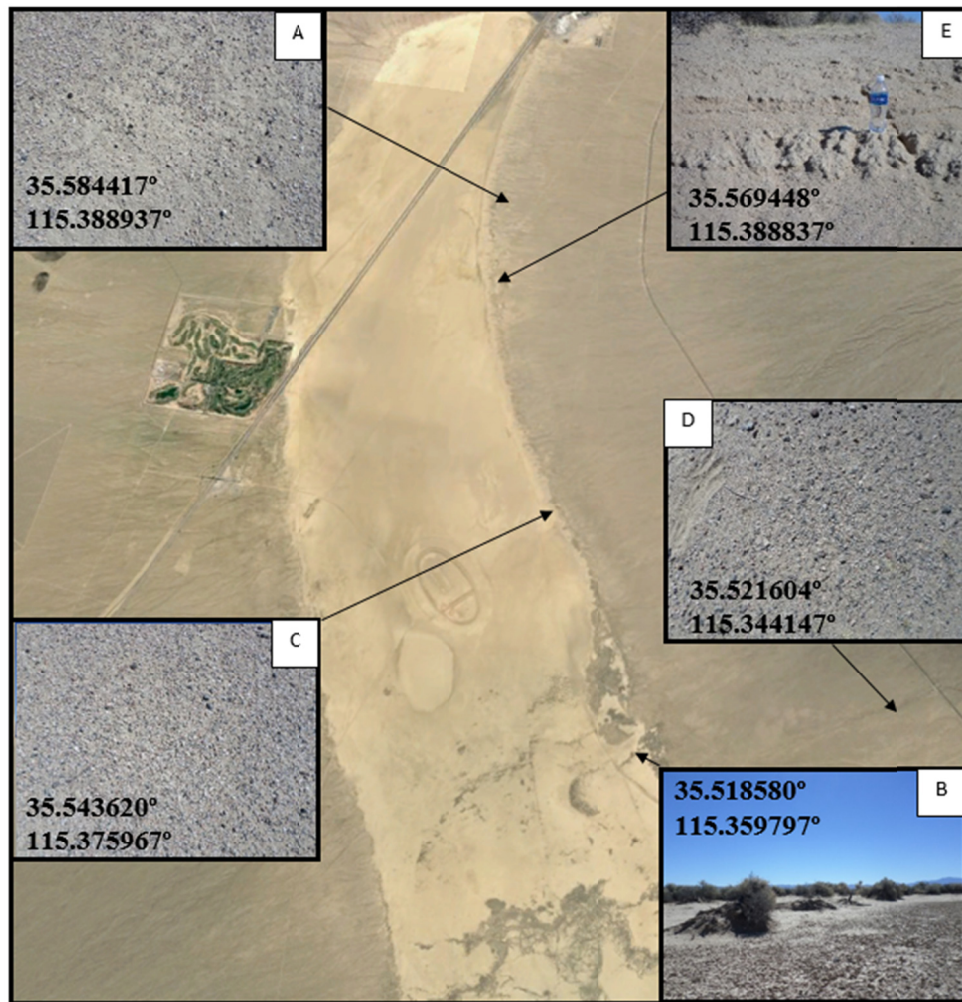


Figure 2. Subtle geomorphic evidence across Lake Ivanpah

4. Methods and Materials

4.1 Field Sampling Methods

Methods employed are similar to Rodrigues-Filho and Muller (1999) and adapted for this region and environment. Their study involved coring to a depth of 12 m with mechanical assistance for sediment characterization. This study was however, limited to hand-driven cores by the BLM due to the culturally sensitive environment of the eastern side of Ivanpah. With this limitation, coring was used only for the identification of subsurface anomalies and to determine the optimal location for further excavation for sediment characterization.

The eastern side of Ivanpah was chosen for this study because the western side of the lake has been heavily impacted by large solar energy projects (Fig. 3A). The first phase consisted of the 12 shallow cores to ~50 cm depth (see supplementary data for geolocations) starting near the center of the dry lake and moving eastward onto the Lucy Gray Mountains bajada (Fig. 3B). Core samples were collected with a hand driven portable AMS hollowstem auger (2.54 cm x 60 cm) kit with a split spoon insert and a high-density polyethylene (HDPE) liner to collect a continuous 50 cm profile. Liners were removed, capped and placed in a cooler for transport to the laboratory. After cores were analyzed and an anomaly was identified (chemical proxy), one location (LGL 5) was excavated by hand to 1.96 m bgs to identify and document substrata features where chemical proxies (the anomaly) were identified. LGL 5 substrata were described and catalogued so that the processes that emplaced the fluvial and aeolian sediments could be linked by their depositional development and chemical proxies detected.

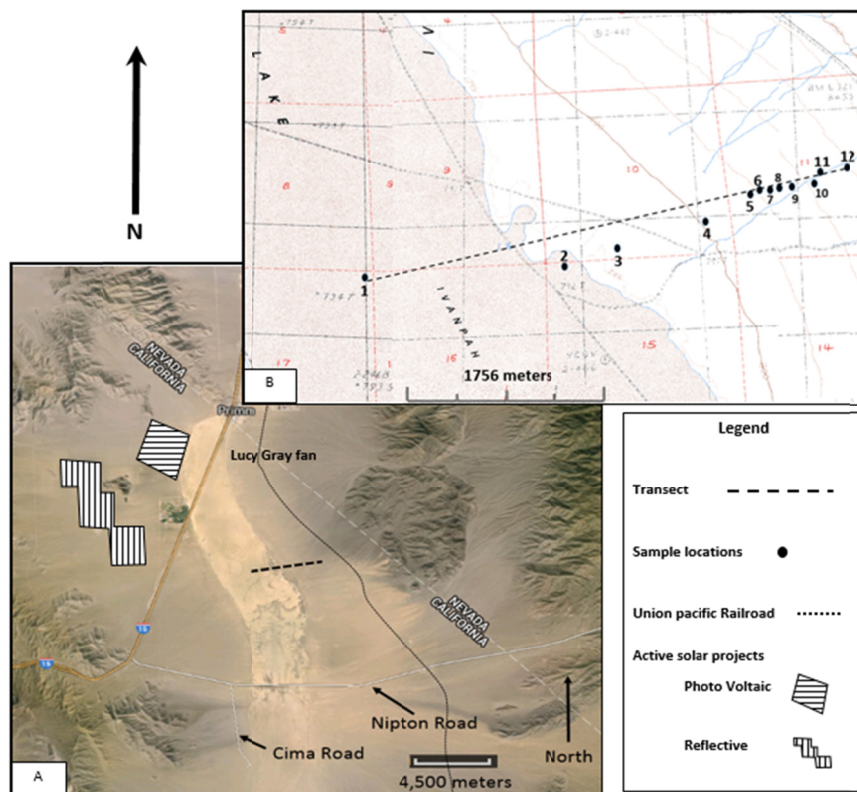


Figure 3. Aerial overview of Lake Ivanpah and features

Note: (A); Lucy Gray Transect (LGL) and sample locations (B). Lat./Long. for locations (1-12) are available in supplementary data.

Samples were collected between May 2014 and December 2015 from lands administered by the United State Department of Interior, Bureau of Land Management (BLM), Needles, California, Field Office. The approach was designed to assess chemical proxies such as trace metals (As, Ba, Cr, Cu, Li, Mn, Ni, Pb, Ti, V) and crustal metals (Al, Ca, Fe, K, Mg, Zn) and wet chemistry (Cl^- , SO_4^{2-} , pH, CaCO_3), Mg/Ca ratios, C^{14} , $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ in sediments across a ~4.5 km transect.

Google Earth was not utilized for surface elevations within the Ivanpah basin because of an error observed. To plot sampling locations and the single excavation site accurately, each sampling site was recorded with a Trimble Survey R10 GNSS rover and base station receiver, GLONASS L1/L2 signal, Trimble HD-GNSS™ processing engine, Trimble SurePoint™ electronic bubble and traceable tilt values with a Trimble 360 satellite tracking. The Trimble xFill™ is designed to work in radio and cellular black spots with an accuracy of ± 0.1 m with all locations tied to United States Geological Survey benchmarks.

4.2 Laboratory Procedure

Each core sample was dissected into 10 cm sections to provide separate data locations at depth. Core dissection was performed by scoring around the HDPE insert, then breaking at the scoring location so that each 50 cm core could be processed as 10 cm sections (0-10, 10-20, 20-30, 30-40, 40-50), resulting in 60 separate samples along the LGL transect. Sediment samples were removed from each section, placed in clean HDPE sample containers, marked with location and depth, and transported to the laboratory.

4.3 Metals Analysis

Researchers Rodrigues-Filho and Muller (2001) have demonstrated that chemical weathering in saturated sediments causes the accumulation of less mobile metals and soluble constituents in a horizontally bedded fashion. Horizontally distributed metals in lake bed sediments are described by Rodrigues-Filho and Muller (1999) as a reliable indicator of previous lake bed features. Therefore, trace elements (As, Ba, Cr, Cu, Li, Mn, Ni, Pb, Ti, V) and crustal metals (Al, Ca, Fe, K, Mg, Zn) in sediments were analyzed using United States Environmental Protection Agency (USEPA) Solid Waste 846 (SW-846) protocols (USEPA, 1997) were

performed at the College of Southern Nevada. These protocols included quality control measures to validate data integrity. Sediment samples (~1 gram) were digested by USEPA method 3050B (hot aqua-regia digestion), diluted to 50 ml after digestion with deionized water, allowed to cool, settle, then filtered to remove particulate matter. Analysis was performed according to USEPA method 6010B; inductively coupled plasma-optical emission spectrometer.

Instrument calibration consisted of six points across a concentration range, including a blank, with a linear regression (R^2) of >0.995 (USEPA, 1997). Analytical integrity was further verified by using a USEPA certified reference sample (RTC Corporation: CRM022-020, Sample 5, lot D522) purchased from a USEPA certified reference laboratory. The certified reference sample measures within specified windows ($\pm 15\%$) and cannot exceed specified requirements. Additionally, samples were analyzed in triplicate with a required USEPA relative percent difference (RPD) of <20% (USEPA, 1997).

4.4 Wet Chemistry

Analysis for wet chemistry were performed at the College of Southern Nevada. Sediment pH is an important measurement for assessing potential availability of beneficial nutrients in sediment. USEPA method 9045 was applied using a H^+ ion-selective glass electrode in a saturated paste to measure sediment pH (USEPA, 1997). The pH meter was calibrated by selection of pH standards that bracketed the sediment pH and measured to within 0.1 standard units.

Sulfate (SO_4^{2-}) was determined by USEPA method 9038 (USEPA, 1997). Ten (10) g of a sample was placed into a 100 mL HDPE bottle, 50 mL of 24°C distilled water was added and agitated for 60 minutes to dissolve SO_4^{2-} ions, then allowed to settle, followed by filtration through WhatmanTM 42 filter paper. The SO_4^{2-} ion was precipitated (with barium-chloride [$BaCl_2$] to barium-sulfate ($BaSO_4$) forming a suspension under controlled conditions to determine the concentration using a nephelometer and a calibration curve for sulfate (USEPA, 1997). Sample weight and solution volume were accounted in the data.

Sediment chloride was determined potentiometrically using a chloride ion (Cl^-) selective electrode (ISE) in conjunction with a chloride combination ISE on a pH meter with an expanded millivolt scale. Each sample was determined by measuring ~10 g into a 100 ml HDPE bottle, adding 50 ml of 24°C distilled water and agitated for 60 minutes to dissolve Cl^- ions. Once the sediment was agitated, the solution was filtered through a WhatmanTM 42 filter paper. Each sample was compared to a calibration curve with final concentration determined by accounting for sample weight and solution volume.

Calcium carbonate ($CaCO_3$) was measured by gravimetric techniques outlined by Bauer et al. (1972). One (1) g of sediment was measured into a tared 25 mL beaker to the nearest 0.001 g. Ten (10) mL of 5N hydrochloric acid (HCl) were added to the beaker covered with a watch glass to prevent splashing. The sample was swirled twice at 10 minute intervals for 30 minutes to allow the HCl and sediment reaction to dissolve all available carbonates. Samples were placed into an oven for 12 hours at 105 °C, removed and allowed to cool before the weight was recorded on both the beaker and the sediment. The difference between the original beaker and sediment weight and the final weight, after the HCl addition, was recorded as $mg\ kg^{-1}\ CaCO_3$.

4.5 Mg/Ca ratios, ^{14}C , $\delta^{13}C$ and $\delta^{18}O$

Mg/Ca ratios were used to examine possible links between past environments and associated temperatures. Magnesium is incorporated into calcite deposited in sediments upon evaporation of water. With the incorporation of Mg as an impurity in calcite (i.e. endothermic), more is incorporated into growing crystals at higher temperatures (Morse et al., 1997). Mg is a divalent cation that substitutes for Ca during the development of biogenic calcium-carbonate (Barker et al., 2005; Barker et al., 2003). When Mg is incorporated into calcite, its incorporation is influenced by the temperature of the surrounding water; increasing with rising temperatures (Ries, 2004; Barker et al., 2005).

Korber et al. (2007), Morse et al. (1997), and Sun et al. (2015) suggest Mg/Ca ratios >3.5 represent modern freshwater. Higher atmospheric temperatures will produce higher Mg/Ca ratios from the aquatic chemical system, whereas lower atmospheric temperatures will produce lower Mg/Ca ratios from the same aquatic chemical system (Barker et al., 2005; Cleroux et al., 2008; Regenberg et al., 2009). For this study, ratios < 2.0 represent a cooler period with decreased evaporation; ratios > 3.5 represent higher temperatures with increased evaporation.

All ^{14}C samples were performed by Beta Analytic Inc. in Miami, Florida and the University of Arizona Accelerator Mass Spectrometry Laboratory in Tucson, Arizona. Beta Analytic Inc. and the University of Arizona Accelerator Mass Spectrometry Laboratory data are conveyed as Conventional Radiocarbon Age (CRA) reported in ^{14}C and cal yrs BP (2 σ age range). Samples for ^{14}C dating were composed of one bulk sample (charcoal

present) and four grab samples (detrital carbon). The bulk sample was ~10 kg collected from within a subsurface beach zone (LU3) layer that was floated for visible charcoal and sent to the laboratory for analysis. The four grab samples came from single locations of ~300 g of sediment collected at each location from within the excavation (LGL5), and sent for analysis.

Laboratory procedures for ^{14}C analysis first visually inspected samples for size, homogeneity, debris, inclusions, clasts, grain size, organic constituents and potential contaminants. Their first step was to disperse sediment samples in de-ionized water, homogenized through stirring and sonication, then sieved through a 180 μm sieve with materials passing through the sieve used for the analysis. The next step in preparation was soaking samples in 2.5N HCl at 90°C for a minimum of 1.5 hours to ensure removal of carbonates (inorganics) so that a measure of organic carbon could be achieved. Following the HCl at 90°C, serial rinses with 70°C de-ionized water were performed until neutrality was reached. Any debris (i.e. micro-rootlets) was discarded during rinsing. Samples were then dried in an oven at 100°C for 12-24 hours and then concentrated HCl was applied to a representative sub-sample under magnification to confirm the absence of carbonates. If carbonate was still present, the laboratory would repeat the HCL processes to remove any inorganic carbonates from the sample. As a final point, although the use of ^{14}C dating does present possible erroneous results, though slight, it is the standard used by researchers for bulk dating of detrital carbon in dry lake sediments, even if no visual carbon is present (i.e. Enzel et al., 2003; Negrini et al., 2006; Quade et al., 2003; Wells et al., 2003).

The University of Arizona Environmental Isotope Laboratory, Department of Geosciences analyzed sediment samples for $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ in mineral carbonates. Values for $\delta^{13}\text{C}$ are important for their understanding of past environments; an increase in prime productivity causes a corresponding rise in $\delta^{13}\text{C}$ as supplementary ^{12}C is locked up in plants. When organic carbon is buried, more ^{12}C is locked out of the sediment system than is found in background systems. Samples analyzed for $\delta^{18}\text{O}$ ratios were used to indirectly determine the temperature of the surrounding water at the time the sediment was deposited. When $\delta^{18}\text{O}$ ratios vary slightly, it is a response to temperature of the surrounding water. Other factors related to $\delta^{18}\text{O}$ are salinity and volume of water in a given area. Studies have used this technique on gastropods, endocarps, and other carbonate matrices; this study used $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ to detect chemical anomalies.

4.6 Quality Control/Quality Assurance

Quality assurance/quality control protocols followed USEPA methods (USEPA, 1997). All analyses required replicated certified reference material purchased from a USEPA certified supplier. Instrument detection limits (IDL) were calculated by multiplying the standard deviation (SD) of a fortified reagent blank by three times. Statistical data were calculated excluding values that were below IDL values. Sample quality control was processed per protocols as outlined in USEPA SW-846 procedures (USEPA, 1997). Finally, all samples were analyzed in triplicate following USEPA requirements for relative percent deviation (RPD) of <20% to be acceptable (USEPA, 1997).

4.7 Stratigraphy and Chronologic Context

Chemical proxy data (“the anomaly”) were compared to this local stratigraphy as well as broader geomorphic features identified along the LGL transect for selection of a point for further excavation. To document the stratigraphy contributing to the anomalies in initial chemical proxy data, a larger (1.5 m x 1.5 m wide) test pit was excavated at LGL 5 (Fig. 3B). Strata were exposed in this excavation on two vertical walls with north-south and east-west axes to a depth of 1.96 m, sketched and photographed. Samples for isotopic analyses were collected from different strata, taking care to avoid sampling across stratigraphic discontinuities. Sediment from identified layers were sieved for gradation and recorded. Water depth is expressed in m, and changes with difference in depth (Δ_w) from the current Ivanpah playa floor of ~794 m amsl. For this study, Rehies et al. (2012) definition was applied for “shallow” lake depths of less than 5 m ($\Delta_w \leq 5$ m), “moderate” depth to Δ_w of 5-8 m, and “deep” lake conditions of $\Delta_w > 8$ m. For the purpose of this paper, the end of the Pleistocene is placed at 12,000 yrs BP, the early to middle Holocene boundary at 7,800 yrs BP, and the beginning of the late Holocene at 4,000 yrs BP.

5. Results

5.1 Sediment Chemical Proxy Anomaly

Coring was used only for the identification of subsurface anomalies so that a location could be identified for further excavation. Elevation change at Ivanpah is ~12.7 m from LGL 1 (794.4 m), at the playa base surface, to 807.1 m amsl at LGL 12: the Lucy Grey bajada creosote bush scrub transition zone (Fig 2b). Table 1 shows a distinct anomaly in data with Al, Fe, K, and V following a uniform distribution below the surface across LGL 1

through 12, as did other constituents such as metals, SO_4^{2-} , Cl^- , pH, CaCO_3 (see [supplementary tables](#)). Aluminum, Fe, K, and V are higher within the horizontally positioned (“bedded”) anomaly and bracketed by lower concentrations above and below the max concentrations (Table 1; highlighted in grey and lighter grey) based on $\pm 1\sigma$. Data shows Al, Fe, K and V have similar distributions at approximately -20 to -40 cm bgs at LGL 1 through LGL 6, surfacing at LGL 7 (799.7 m), descending to 20-40 cm bgs by LGL 8, surfacing again at LGL 11 (804.2 m) and descending to 20-40 cm bgs at LGL 12. Largely, this anomaly is positioned bgs with a uniform depth except at LGL 7 and LGL 11, where it surfaces.

Crustal (e.g. Al, Fe, Ca) and trace metals (e.g. Cr, Cu, Pb, V) ranged between 3,500 and 46,010 mg kg^{-1} Al; 5,150 and 29,780 mg kg^{-1} Fe; and 8.7 and 83 mg kg^{-1} V across the LGL transect (see [supplementary tables](#)). These findings are not unusual as oxidizing environments increase the presence of FeOOH in horizontally bedded sediments (Stumm and Morgan, 1996; Sanchi et al., 2015), providing adsorptive capacity for charged species dissolved in surface and percolating waters. Data presented in Table 1, including supplementary tables, shows Al, Fe, Zn and other metals (e.g. As, Ba, Ca, Cr, Cu, Li, K, Mg, Mn, Ni, Pb, Ti, V) tend to follow this horizontal bedding (“an anomaly”) at depth (~20-40 cm) from LGL 1 out ~4.5 km east to LGL 12.

Sediment pH also measured along the LGL transect ranged between 6.81 and 8.50 standard units (SU) with a distribution comparable to metals and other chemical proxies (Table 2). Core sample pH, although not significantly different across the LGL transect, shows the anomaly at depth and followed metal accumulations presented in Table 1, a common association know between pH and metals (Tardy and Roquin, 1992; Watanabe et al., 2001). These findings are not unusual as sediment pH will support similar chemistries in horizontal beds where there are analogous redox conditions (Tardy and Roquin, 1992).

Data for $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ were also used to identify patterns below the surface with two locations (LGL 3 and 5; Fig. 3B) along the LGL transect selected for measurements. Table 3 displays $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ with comparable alignments as presented in Tables 1 and 2. LGL 3 is located within the playa (794.5 m amsl) base surface, whereas LGL 5 is located on the Lucy Grey bajada (796.4 m amsl) ~1.9 m above LGL 3. Data for $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ at LGL 3 and 5 have a similar orientation, although with minor differences; -2.20, -2.56 $\delta^{13}\text{C}$ and -4.71, -5.10 ‰ $\delta^{18}\text{O}$, respectively.

Table 1. Selected chemical proxies for locations 1 - 12 of the LGL transect with Max $\pm 1\sigma$ ranges

Location	1	2	3	4	5	6	7	8	9	10	11	12	
Elevation	794.4	794.7	794.5	796.3	796.4	797.4	799.7	800.6	801.9	803.2	804.2	807.1	
Depth													
Al	10	11,600	12,500	16,500	18,750	28,860	27,850	33,160	4,015	36,920	6,700	44,580	31,200
	20	20,550	21,000	17,000	16,600	36,290	41,920	44,760	7,200	38,670	3,500	43,010	45,920
	30	20,600	20,300	24,800	19,500	49,670	40,010	44,750	9,350	41070	7,650	41,920	46,010
	40	20,900	23,500	24,100	17,100	39,370	32,550	34,760	8,200	40,960	7,360	36,410	33,010
	50	20,400	21,050	20,800	16,200	32,220	35,600	26,130	5,800	36,000	6,080	31,440	32,000
	STDV	4035	4188	3863	1426	7990	5674	8028	2081	2301	1657	5443	7638
Max +1 σ	24935	27688	28663	20926	57660	47594	52788	11431	43371	9307	50023	53648	
Max -1 σ	16865	19312	20937	18074	31380	36246	36732	7269	38769	5993	39137	38372	
Fe	10	17200	15560	15700	9050	23340	21810	25920	5150	24800	8100	29000	23400
	20	17350	16265	16805	13350	25020	27250	27190	8305	25800	8950	29100	28630
	30	16550	24550	26400	24800	29780	26600	27450	18250	26210	9250	28700	29110
	40	27650	27800	26900	25900	26690	21880	24080	29750	25830	7800	25600	23000
	50	27500	26350	26500	17750	25050	24680	21760	16800	24270	7900	23300	22800
	STDV	5782	5774	5684	7251	2434	2554	2377	9632	812	657	2591	3190
Max +1 σ	33432	33574	32584	33151	32214	29804	29827	39382	27022	9907	31691	32300	
Max -1 σ	21868	22026	21216	18649	24256	24696	25073	20118	25398	8593	26509	25920	
K	10	22270	27820	26790	10150	8815	8280	9581	1920	9144	9250	9513	8065
	20	22405	30330	29300	17705	9734	9576	9611	2614	9753	9575	9404	9439
	30	29480	25040	28000	21395	9620	9600	9784	2831	9461	16200	9564	9217
	40	32890	34050	28650	24980	9532	8635	9341	2214	9312	10800	9093	8900
	50	30380	31670	27390	20150	9070	8556	7913	2402	9448	8800	1280	8850
	STDV	4863	3479	993	5539	393	616	762	352	224	3041	3633	522
Max +1 σ	37753	37529	30293	30519	10127	10216	10546	3183	9977	19241	13197	9961	

Max -1 σ	28027	30571	28307	19441	9341	8984	9022	2479	9529	13159	5931	8917	
V	10	46	39	39	16	55	46	65	8.7	51	12	79	57
	20	44	37	37	28	57	65	71	14	69	15	83	78
	30	48	42	41	34	77	61	71	19	67	16	82	78
	40	48	44	43	41	67	49	62	18	61	11	68	69
	50	44	45	40	31	61	43	58	13	63	14	57	66
STDV	2.0	3.4	2.2	9.2	8.9	9.7	5.7	4.1	7.0	2.1	11.1	8.8	
Max +1 σ	50	48	45	50	86	75	77	23	76	18	94	87	
Max -1 σ	46	42	41	32	68	55	65	15	62	14	72	69	

Note: Dark shading is maximum value in vertical profile; lighter shading is $\pm 1\sigma$ of maximum value detected in vertical profile; elevation in meter; depth in cm below surface; concentrations in mg kg⁻¹; σ is standard deviation. Additional data available in supplementary tables.

Table 2. Selected pH values found in the Lucy Grey Line

Depth	LGL Locations			
	3	5	7	11
10	7.21	7.39	7.62	7.31
20	7.41	7.30	7.63	7.27
30	7.54	7.62	7.27	7.19
40	8.05	7.15	7.16	7.22
50	7.96	7.81	7.34	7.54

Note: pH values are in S.U.; depth in cm below surface; shaded values match Table 1

The lowest $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ values detected at LGL 3 (-2.56 ‰, -5.42 ‰, respectively) were at 0-10 cm bgs. LGL 5 $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ revealed a comparable pattern to LGL 3; $\delta^{13}\text{C}$ ranged between 1.39 ‰ and -1.33 ‰ with 1.39 ‰ at 50 cm bgs and -1.33 ‰ at 0-10 cm bgs. This alignment shows the highest $\delta^{13}\text{C}$ at 40-50 cm bgs with the lowest values at 0-10 cm bgs. Distribution of $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ align with other chemical proxies with a distinct change (Δ) in values based on depth. The $\Delta\delta^{18}\text{O}$ is similar to $\Delta\delta^{13}\text{C}$; highest at 40-50 cm bgs with the lowest values detected at 0-10 cm bgs. Data for both $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ illustrate a similar anomaly below the surface as seen with other constituents. The observed pattern is consistent when the local system is under similar hydrologic balances caused by environmental changes (e.g. Watanabe et al., 2001, Regenberget al., 2009). Data also show the highest $\delta^{13}\text{C}$ value (1.39 ‰) at LGL 5 at 40-50 cm bgs with the lowest value (-1.33 ‰) at 0-10 cm bgs. At shallower depths (0-10 cm), $\Delta\delta^{13}\text{C}$ decreases, likely the result of increased temperatures over the past 1.0 ka yrs BP. Moreover, $\delta^{13}\text{C}$ data indicate that the deeper levels (40-50 cm bgs) were likely long-standing, water-rich environments. LGL 5 $\delta^{18}\text{O}$ values were highest (-2.85 and -5.30 ‰, respectively) at 40-50 cm and 20-30 cm bgs with the lowest (-5.66 and -5.42 ‰, respectively) at 0-10 cm bgs. Data show $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ within the LGL5 anomaly are consistent with mid-Holocene. The $\delta^{13}\text{C}$ data ranged from -0.11 to 0.03‰ and $\delta^{18}\text{O}$ data ranged from -3.56 to -2.85‰.

Table 3. $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ ratios for LGL 3 and 5

Location	depth	$\delta^{13}\text{C}$	STDV	$\delta^{18}\text{O}$	STDV
3	10	-2.56	0.036	-5.42	0.022
	20	-2.21	0.012	-5.10	0.037
	30	-2.48	0.020	-4.71	0.062
	40	-2.20	0.017	-5.34	0.021
	50	-2.30	0.005	-5.30	0.034
	0	-1.33	0.024	-5.66	0.029
5	20	-1.33	0.025	-4.92	0.035
	30	-0.11	0.031	-3.56	0.045
	40	0.03	0.011	-4.23	0.018
	50	1.39	0.017	-2.85	0.072

Note: Depth: cm below surface, values are in ‰; Laboratory ID: AA K883

Metal ratios (Mg/Ca) shown in Table 4 have comparable alignments to other sediment chemistry presented in Tables 1-3; following the detected anomaly. Data show 81% of Mg/Ca ratios are <3.5 and 21% are >3.5; values set by other authors to indicate higher temperatures and increased evaporation (Barker et al., 2005; Cleroux et al., 2008; Regenberg et al., 2009). Only 5% of ratios are <2.0; a value symptomatic of cooler temperatures with decreased evaporation (Barker et al., 2005; Regenberg et al., 2009). Data for LGL 3: 30-50 cm; LGL 5: 20-40 cm; and LGL 7 and 11: 0-20 cm, show elevated Mg/Ca ratios within the anomaly, indicative of long-standing water. Table 4 displays a comparable pattern as shown in other data tables; lowest Mg/Ca ratios are situated within the identified chemical-proxies. These findings show that Ivanpah had periods of moist-humid conditions followed by aridity caused by environmental fluctuations.

Table 4. Mg/Ca ratios of the LGL transect

	Locations											
Depth*	1	2	3	4	5	6	7	8	9	10	11	12
10	3.83	3.64	4.03	2.07	2.89	2.45	2.44	2.54	2.32	2.08	2.85	1.97
20	3.89	3.94	3.86	2.83	2.48	2.99	2.36	2.30	2.35	2.03	2.95	2.29
30	3.94	4.32	3.19	2.73	3.38	3.12	2.31	2.08	2.82	2.24	2.62	2.31
40	3.16	4.21	3.60	1.86	3.29	2.61	2.29	2.83	2.80	2.09	2.61	2.41
50	3.72	4.38	3.60	1.82	2.99	2.79	2.31	2.72	2.99	2.13	2.49	2.47

Note: * cm below surface

5.2 Geomorphic and Stratigraphic Features at LGL-Test Pit 5 Discussion

Chemical proxies (shallow coring) led to a 1.5 x 1.5 m wide by ~2 m depth excavation at LGL 5 (796.4 m amsl; 35.523617°, -115.341933°) to explore the anomaly identified at 20-40 cm bgs (Table 1, Fig. 4A). Research shows the presence of possible lacustrine units below the surface; these units are designated as Lake Unit (LU) 0, 1, 2 and 3 with LU3 subdivided into LU3a, b, c, and d. Five samples were collected from LGL 5 and sent for ¹⁴C analysis and are presented within their respective sections below. The excavation sidewalls displayed a mixture of compositions dominated by sand, silt, and clay, with some clastic materials at shallow depths, unconformities indicative of three possible lake beds, and two apparent playa surfaces exhibiting mud cracks.

The surface (unit ALØ) of LGL5 is composed of a relatively thin (~10 cm bgs) layer of light brown (Color hue 7.5YR 6/4), very fine to fine sandy loam with granules, pebbles and small gravels at the base (Fig. 4F). ALØ is dominated by an upward sequence of fine-earth textures and a non-laminated matrix, aeolian in origin, becoming coarser at depth, grading to pea-sized pebbles, clasts, and small gravels situated on a boundary surface (LU3d) at depth (-10cm). ALØ represents a late Holocene-to-modern deposit situated on an older unit (LU3).

LU3 is a relatively thick unit (LU3a, b, c, d) ranging from ~10 to 60 cm bgs with a base elevation of 795.8 m amsl when adjusted for ground elevation. Materials within LU3d (~10 to 20cm bgs) is an aridisol composed of light brown to reddish brown, sandy loam (color hue 7.5YR 6/4) with subrounded to subangular texture, showing classic oxidation (Fig. 4B; Retallack, 1988). This unit contains 19.4% pea-sized gravels (0.5-1.0 cm), 72.9% sands, and 8.0 % silts (Table 5), consolidated by clays with carbonate strings forming a weak vesicular surface that appears erosionally truncated. The surface of LU3d displays a disconformity (truncated) representing a rapid regressive phase overlain by ALØ and leaving no subsequent visual evidence. The lack of well-developed soils in LU3d is consistent with a younger age for the most recent transgressional phase situated on a base of gravels and sands of LU3a, b, and c. Lastly, LU3d deposition is difficult to determine, the structure reduces vertical infiltration, making it problematic to date owing to very high inorganic carbon, though it reflects the cumulative effects of pedogenic processes with time.

LU3a, b, and c consist of 27.5% gravels, 71.4% sands and 1.1% silts, sedimentologically characteristic of a fluvial deposit beach facies containing a poorly sorted matrix (Table 5; Figs. 4). The unit grades upward from sub-rounded basal gravels (~2-5 cm) and clasts (~2-5 cm) with minor sand in LU3a (Fig. 4C); pea-sized beach gravels (~0.7-2 cm) in LU3b (Fig. 4F); and sand/loamy sand in LU3c (Fig. F) yielding a composite (LU3a, b, c) age of 1,050-1,025 cal yrs BP (Table 6). All subunits within LU3 (a-c) are light brown with a color hue 7.5YR 6/3. The uppermost portion of LU3c is positioned at 796.2 m amsl when adjusted for current surface elevation. Evidence shows that a shoreline advanced eastward from changing lake volumes, water deepened at LGL 5, depositing sandy loam with pea-sized beach gravels within unit LU3a-c forming a classic transgressional sequence of poorly sorted, lacustrine gravel beach facies (Bacon et al., 2006). Structurally, LU3 (a-c) is ~30 cm

thick fluvial deposit of sands, pea-sized gravels and clasts, all positioned on an older lacustrine unit boundary surface (LU2).

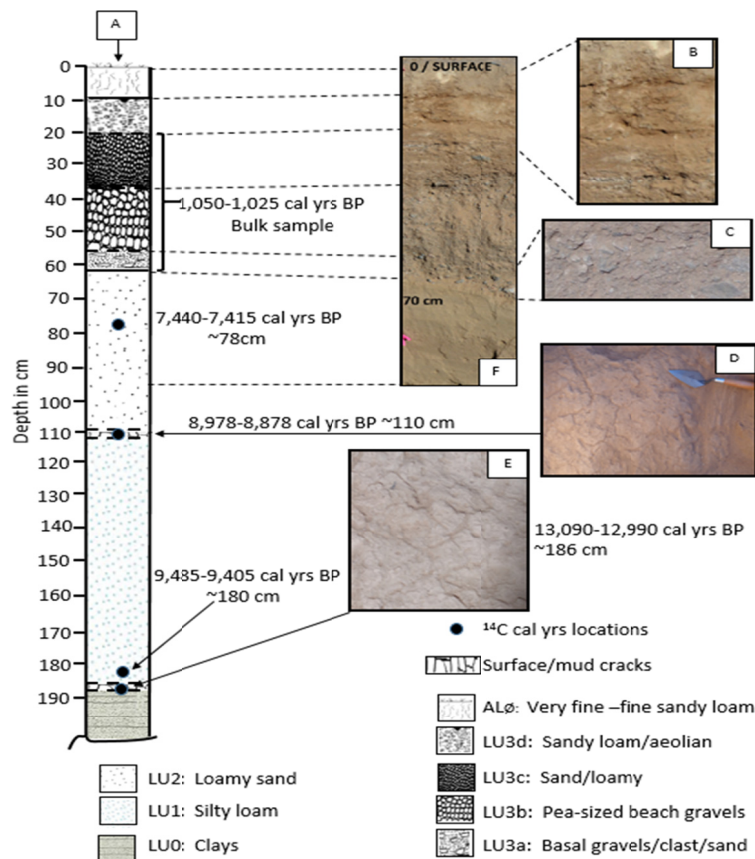


Figure 4. LGL 5 stratigraphic cross section from transect

Unit LU2 is positioned at 795.8 m amsl when adjusted for elevation and consists of <0.1 % gravels, 89.1% sands and 10.4% silts and <0.5% clays (Table 5). This unit consist of a light brown loamy sand (color hue 7.5YR 6/4) containing carbonate strings within the upper few (~3) cm of a thicker component ranging from ~63 cm to ~109 cm bgs yielding a date of 7,440-7,415 cal yrs BP at ~78 cm bgs (Table 6). Under magnification (4x), bedding in LU2 consists of loamy sand-sized grains arranged in parallel lines (partings and intervening) with fine to very fine sands erosionally truncated at ~63 cm bgs (Fig. 4F) exhibiting minor surface cracking. LU2 sediments are very well sorted and interbedded, with very thin (i.e. 1-3 cm thick) lamination throughout, interpreted as beach sand overlain by a coarser fluvial unit LU3 as described by McKee and Weir (1953). LU2 is likely the result of a low energy suspension event overlain LU1.

Table 5. Grain size, color hue of LGL5 sediments

Unit ID	~Depth (cm)	Color hue	Gravel %	Sand %	Silt %	Clay%
AL0	0-10	7.5YR 6/4	1.0	83.4	15.5	<0.1
LU3d	10-21	7.5YR 6/4	19.4	72.9	8.0	<0.1
LU3c	21-36	7.5YR 6/3	20.8	74.1	5.1	<0.1
LU3b	36-55	7.5YR 6/3	23.7	74.8	1.8	<0.1
LU3a	55-63	7.5YR 6/3	37.6	62.3	0.1	<0.1
LU2	63-109	7.5YR 6/4	<0.1	89.3	10.5	<0.1
LU1	109-186	7.5YR 7/3	<0.1	25.7	74.2	<0.1
LU0	186-unknown	7.5YR 6/4	<0.1	44.2	53.1	2.6

Note: LU0 was explored only to 196 cm below surface

Table 6. Radiocarbon dates from LGL5 sediments

Lab ID	Unit ID	Depth ¹ (cm)	Materials dated	$\delta^{13}\text{C}$ (‰)	¹⁴ C age ² (yrs BP)	cal yrs BP ³ (2 σ age range)
Beta432398	LU3(a-c)	-30 to -61	Bulk C _{org}	-11.5	1,060 ± 30	1,050-1,025
Beta434682	LU2	- 78	Grab MC _{org}	-17.5	6,500 ± 31	7,440-7,415
AA107397	LU1	-110	Grab MC _{org}	-18.1	7,940 ± 31	8,978-8,878
Beta434906	LU1	- 180	Grab MC _{org}	-21.6	8,400 ± 30	9,485-9,405
Beta436547	LU0	- 186	Grab MC _{org}	-17.0	11,160 ± 40	13,090-12,990

Note: C: carbon, M: microscopic, org: organic, ¹ cm below surface; ² reported in conventional radiocarbon years; ³ reported in calibrated years BP with 95% probability

LU1 is a well preserved lacustrine unit commencement at ~110 cm bgs, extending to ~186 cm bgs, and appearing to have been deposited by low energy suspension. The superficial portion of this unit dated to 8,978-8,878 cal yrs BP at ~110 cm bgs and at depth (~180 cm bgs) dated 9,485-9,405 cal yrs BP. The surface of LU1 (~110 cm) is desiccated, showing signs of cracking (“mud-cracks”) with a crust rich in swelling clays exhibiting evaporite-staining of the surface (~2 cm), indicative of an established desert playa (Fig. 4D). The sediment matrix consists of very thin laminated pink silt loam (color hue 7.5YR 7/3) containing <0.4 % gravels, 24.2% sands, 74.6% silts, and 0.8% clays (Table 5). In contrast to other units, LU1 appears more sorted with overlapping of edges (“imbricated”), and sand-size grains that are marginally rounded. The appearance of this unit is likely the effects of a repeated influx of water followed by episodic evaporation resulting in rough-crusts surfaces similar to those described by Adams and Sada (2014) and Tollerud and Fantle (2014). LU1 is dissimilar to units LU2, LU3 and ALØ; sediment matrix is very thin laminated silt loam, indicative of a lacustrine water facies overlain a deeper unit (LU0).

The lowest unit (LU0) encountered was a preserved lacustrine feature of light brown clay (color hue 7.5YR 6/4) beginning at ~186 cm bgs, dating to 13,090-12,990 cal yrs BP at 186 cm bgs. It is important to note that this unit was not explored beyond ~196 cm bgs owing to limited excavation techniques and safety. Sediments consisted of very thin laminated, fine-grained, imbricated clay-rich matrix composed of <0.1 % gravels, 44.2% sands, 53.1% silts and 2.6% clays (Table 5). The surface of LU0 contained surface-cracking (“mud-cracks”) with evaporite-staining (~2.5 cm), composed of well-sorted, firm, illuviation of argillic clays that were bedded, very plastic, and with particle sizes ranging between 1 mm and 2 μm (Fig. 4E). This unit is progressively thicker with well-developed grains, ped-cutans, and blocky to prismatic structure. LU0 is clearly the result of chemical weathering of local silicate-bearing materials that produced primary clay deposits. LU0 is in stark contrast to other units; the matrix is composed of firm, very thin, laminated clays indicative of a long-standing, lacustrine, deep-water facies. Finally, the age of LU0 (13,090-12,990 cal yrs BP at ~186 cm bgs) places the surface firmly within the Pleistocene - Holocene transitional period.

6. Discussion

Major environmental events inferred from the sedimentological evidence at Ivanpah were compared to those of other studies from localities within or bordering the Mojave Desert. Documented events are likely associated with other recognized climatic shifts in the region, for instance, Wells et al. (2003) and others describe climatic shifts (drying events) within the region at ca. 9.2, 7.6, 8.2, 3.2 ka yrs BP with several minor shifts (wetting events) occurring ca. 2.5, 2.0, 1.5, 1.0 and 0.8 ka yrs BP, respectively (Cook et al., 2007; Harvey et al., 1999; Holmquist et al., 2016). Sediments from the LGL 5 excavation yielded data documenting multiple sequences with LU3d characterized as an aridisol sandy loam showing minor pedogenic processes (Figs. 4F/B). This unit’s (LU3d) deposition and age is problematic though, evidence suggests it is younger than LU3a-c (1,050-1,025 cal yrs BP). Within this time frame, LU3d is probably linked to the wetting period (~800 to 600 cal yrs BP) described by Mensing et al. (2008) (Fig. 4B). They documented that after the wetting period a drier, arid climate settled into this region. This drier period is likely responsible for ALØ; an aeolian sandy-silt bed deposited on the north-northeastern lateral of the Ivanpah valley.

Unit LU3a-c is a classic, near-shore littoral deposit of interbedded gravel and sand beach facies grading upward from basal to pea-sized gravels containing sands and minor silts as described by Bacon et al. (2006). LU3 (a-c) age of 1,050-1,025 cal yrs BP (Figs. 4F/B/C) places this unit within Holmquist et al. (2016) and Cook et al. (2007) period of warming over the Western United States, promoting increased monsoonal activities ~1,000 yrs BP. Other authors also describe sporadic wetting periods around the same time that caused monsoonal events (spring to winter) beginning ~1,900 and ending ~1,100 cal yrs BP (Negrini et al., 2006). Holmquist et al. (2016)

further describes a period ending ~1,280 cal yrs BP when this region became significantly wetter; perhaps providing ideal stormy lake conditions to establish the beach zone of LU3_(a-c) that overlays the unit of LU2.

The surface unit LU2 is erosionally truncated with very minor desiccation cracking dating to 7,440-7,415 cal yrs BP at ~78 cm bgs (Fig. 4F). Researchers Tollerud and Fantle (2014) studied similar environments documenting minor desiccation of playa surface and the stages at which mud-cracks form (Tollerud and Fantle, 2014). They also describe the subsurface of similar mud-cracked units as very thin laminated sediments resulting from a low energy suspension environment. Their efforts described surfaces similar to the minor desiccation of LU2 where the evolution of the mud-cracks promotes a smooth, evaporite-stained, strong surface with minor cracking of ~3-5 cm long 0.5-1.5 cm thick, as seen with the surface of LU2.

Seismological evidence for LU2, a unit ~46 cm thick, is indicative of a period of drying that occurred after a wetting phase (Fig. 4F). This unit is composed of classic fine, very thin laminated sediments commencing at ~3-4 cm below the desiccated surface (~63 cm bgs) to a depth of ~109 cm bgs. It is dissimilar to LU3_(a-d) as it contains substantially more sands and silts, with no gravels. Evidence indicates that it is the result of a classic low energy depositional lacustrine environment similar to that described by Tollerud and Fantle (2014). Murchison and Mulvey (2000) and Patrickson et al. (2010) documented a drying events between 8,200 and 7,650 yrs BP causing recessional phases of the Great Salt Lake located northeast of Ivanpah. Furthermore, Blunt and Negrini et al. (2015) recognized events west-northwest of Ivanpah where a Tulare Lake shoreline (ca. 8,166-7,984 and 8,857-9,008 cal yrs BP) began a recessional phase due to aridity as well. It is probable that LU2 is the result of those documented drying events and subsequent desiccation of LU2.

Below LU2 is LU1, a unit composed of very fine, very thin laminated sediments commencing at ~109 cm to a depth of ~185 cm bgs. Of importance at LU1 is the surface mud-cracks, which are ~3-10+ cm long by a ~2-5 cm thick; they are smooth, contain minor evaporite-staining, and have a strong surface (Fig. 4D). Below the surface cracking are very thin laminated sediments suggestive of a classic low energy depositional lacustrine environment as described by others (Negrini et al., 2006; Tollerud and Fantle, 2014). The surface (~109 cm bgs) of LU1 yielded an age of 8,978-8,878 cal yrs BP with a lower depth (~180 cm) dating to 9,485-9,405 cal yrs BP (Fig. 4A).

LU1 links with documented regional shifts that affected the southwestern United States (e.g. Great Salt and Tulare lakes). Harvey et al. (1999) documented that during the early Holocene (ca. 9,500 -8,600 cal yrs BP), the jet stream moved north, and allowing monsoonal activity to enter the American Southwest inundating geographic basins. Such a shift in environmental conditions would have been the reason for the existence of LU1: a thick lacustrine facies. Following this event (Harvey et al. 1999), a drying episode described by Murchison and Mulvey (2000) likely led to the desiccation of the surface of LU1. Data suggests that the surface of LU1 is the product of the early Holocene lake termination event described by Murchison and Mulvey (2000).

The deepest unit, LU0 (13,090–12,990 cal yrs BP), represents a period of a long-standing water environment during the Pleistocene – Holocene transition (ca. 14,600–8,000 yrs BP) as described by Antinao and McDonald (2013). Moreover, at Tulare Lake, a period of increased wetting was documented (ca. 12,605 to 12,971 cal yrs BP), promoting perennial lakes within the region (Bacon et al., 2006, Kirby et al., 2015), possibly driving the wetting events at Lake Ivanpah. Furthermore, authors Harvey et al. (1999) describe an arid phase (ca. 13,000-10,500 yrs BP) following the perennial lake period, likely related to Bacon et al. (2006) and Kirby et al. (2015) work, resulting in the termination of pluvial lakes in this region. These drying events are probably the cause of the recessional phases identified at LU0 dating to 13,090 – 12,990 cal yrs BP (Fig. 4E). Evidence clearly puts unit LU0 within the early Pleistocene – Holocene transition; however, it is also feasible that LU0 is the result of the end of a late pluvial lake phase. Because of the limitation on the excavation imposed by the BLM, it is difficult to extrapolate much more from LU0, as additional work including deeper excavations would be required to fully describe this unit.

7. Summary and Conclusions

Dry lake systems located within the Mojave Desert are potentially good sites for alternative energy such as solar energy collection however, they have not been sufficiently studied for late Pleistocene - early Holocene lacustrine environments. Development could adversely impact valuable sedimentological resources contained within the shallow subsurface. This study used shallow sediment chemical proxies and characterization to assess subsurface conditions for buried lakebed features at Lake Ivanpah. The number and depth of shallow cores, and uses of trenching, to describe shallow sedimentary facies for this paper were adequate for limited delineation of wet and dry periods associated with lacustrine lake occupations, but does need more comprehensive data collection (especially subsurface data) and interpretation of this important period of desertification in recent

geologic history.

Results at Lake Ivanpah illustrated that transitional lakes existed in the basin for two Holocene and one Pleistocene-Holocene transitional periods. Although there are no signs of visual wave-cutting at Ivanpah, chemical proxies and sediment characterization of site LGL5 showed a buried shoreline (“beach zone”) and two lacustrine features. Furthermore, chemical proxies also identified two possible truncated shorelines at sites LGL7 and 11. With this in mind, evidence shows lake high-stands likely occurred at water depths of ~1.8, ~5.3 and ~9.8 m, respectively, in the Ivanpah valley. Lacustrine units identified beneath the surface at LGL5 were dated to ca. 1,050-1,025 (LU3a-c), 7,440-7,415 (LU2), and 9,485-9,405 (LU1) cal yrs BP with the deepest lake unit (LU0) dating to ca. 13,090-12,990 cal yrs BP. This evidence clearly shows that Lake Ivanpah has a significantly more complex history than just a single continuous lake episode of mid- to late-Holocene in age but rather, multiple episodes of lake stands, the oldest being a Pleistocene-Holocene (or older) transitional lake.

It is likely that other lacustrine units exist beneath the Ivanpah playa. The lack of visual evidence is most likely the result of an abrupt regional climatic shift <1,000 yrs BP, resulting in shoreline truncation and obliteration. Understanding environmental change in these basins is needed to better recognize climate changes during the late Pleistocene-early Holocene and regional desertification. Further studies of the surface of the Ivanpah and Roach dry lakes and surrounding landscapes is required to better characterize the overall basin. Finally, a more comprehensive surface and subsurface study is required to describe past conditions and depth of lacustrine environments at Ivanpah before extensive alternative energy developments disrupt shallow sedimentary evidence necessary for future research.

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Supplementary Data

		Locations												
		Depth	1	2	3	4	5	6	7	8	9	10	11	12
Al Units mg kg-1	10	11,600	12,500	16,500	18,750	28,860	27,850	33,160	4,015	36,920	6,700	44,580	31,200	
	20	20,550	21,000	17,000	16,600	36,290	41,920	44,760	7,200	38,670	3,500	43,010	45,920	
	30	20,600	20,300	24,800	19,500	49,670	40,010	44,750	9,350	41070	7,650	41,920	46,010	
	40	20,900	23,500	24,100	17,100	39,370	32,550	34,760	8,200	40,960	7,360	36,410	33,010	
	50	20,400	21,050	20,800	16,200	32,220	35,600	26,130	5,800	36,000	6,080	31,440	32,000	
	Mean	18810	19670	20640	17630	37282	35586	36712	6913	38724	6258	39472	37628	
	STDEV	4035	4188	3863	1426	7990	5674	8028	2081	2301	1657	5443	7638	
	MAX	20900	23500	24800	19500	49670	41920	44760	9350	41070	7650	44580	46010	
	MIN	11600	12500	16500	16200	28860	27850	26130	4015	36000	3500	31440	31200	
	<hr/>													
		Depth	1	2	3	4	5	6	7	8	9	10	11	12
Fe Units mg kg-1	10	17200	15560	15700	9050	23340	21810	25920	5150	24800	8100	29000	23400	
	20	17350	16265	16805	13350	25020	27250	27190	8305	25800	8950	29100	28630	
	30	16550	24550	26400	24800	29780	26600	27450	18250	26210	9250	28700	29110	
	40	27650	27800	26900	25900	26690	21880	24080	29750	25830	7800	25600	23000	
	50	27500	26350	26500	17750	25050	24680	21760	16800	24270	7900	23300	22800	
	Mean	21250	22105	22461	18170	25976	24444	25280	15651	25382	8400	27140	25388	
	STDEV	5782	5774	5684	7251	2434	2554	2377	9632	812	657	2591	3190	
	MAX	27650	27800	26900	25900	29780	27250	27450	29750	26210	9250	29100	29110	
	MIN	16550	15560	15700	9050	23340	21810	21760	5150	24270	7800	23300	22800	
	<hr/>													
		Depth	1	2	3	4	5	6	7	8	9	10	11	12
K Units mg kg-1	10	22270	27820	26790	10150	8815	8280	9581	1920	9144	9250	9513	8065	
	20	22405	30330	29300	17705	9734	9576	9611	2614	9753	9575	9404	9439	
	30	29480	25040	28000	21395	9620	9600	9784	2831	9461	16200	9564	9217	
	40	32890	34050	28650	24980	9532	8635	9341	2214	9312	10800	9093	8900	
	50	30380	31670	27390	20150	9070	8556	7913	2402	9448	8800	1280	8850	
	Mean	27485	29782	28026	18876	9354	8929	9246	2396	9424	10925	7771	8894	
	STDEV	4863	3479	993	5539	393	616	762	352	224	3041	3633	522	
	MAX	32890	34050	29300	24980	9734	9600	9784	2831	9753	16200	9564	9439	
	MIN	22270	25040	26790	10150	8815	8280	7913	1920	9144	8800	1280	8065	

V Units mg kg-1	Depth	1	2	3	4	5	6	7	8	9	10	11	12
	10	46	39	39	16	55	46	65	8.7	51	12	79	57
20	44	37	37	28	57	65	71	14	69	15	83	78	
30	48	42	41	34	77	61	71	19	67	16	82	78	
40	48	44	43	41	67	49	62	18	61	11	68	69	
50	44	45	40	31	61	43	58	13	63	14	57	66	
Mean	46	41	40	30	63	53	65	15	62	14	74	70	
STDEV	2	3.4	2.2	9.2	8.9	9.7	5.7	4.1	7	2.1	11.1	8.8	
MAX	48	45	43	41	77	65	71	19	69	16	83	78	
MIN	44	37	37	16	55	43	58	9	51	11	57	57	

Mg Units mg kg-1	Depth	1	2	3	4	5	6	7	8	9	10	11	12
	10	6700	6700	6650	4480	8640	8299	9180	3519	9698	3952	10680	8814
20	6710	6750	6600	6100	9790	11170	10510	4370	9535	4025	10480	10520	
30	6600	6675	6800	6600	12200	11050	10710	4380	10430	4143	9030	10630	
40	6850	8100	6900	6650	10790	9114	9178	4092	10370	3870	9100	9100	
50	7400	7700	6700	6100	9170	9490	7849	3700	9553	3658	8574	8900	
Mean	6852	7185	6730	5986	10118	9825	9485	4012	9917	3930	9573	9593	
STDEV	319	668	120	882	1412	1251	1164	391	446	182	944	903	
MAX	7400	8100	6900	6650	12200	11170	10710	4380	10430	4143	10680	10630	
MIN	6600	6675	6600	4480	8640	8299	7849	3519	9535	3658	8574	8814	

Ca Units mg kg-1	Depth	1	2	3	4	5	6	7	8	9	10	11	12
	10	25700	24400	22800	4300	25050	20300	21300	8950	22940	7420	30430	17360
20	26100	26650	25450	17550	24310	33480	24840	9000	22900	7950	30910	24070	
30	26050	28490	24450	16950	41210	34500	27850	9100	29500	9270	27320	24590	
40	21700	34100	22050	12370	35600	23750	23810	10330	29050	8100	25340	22100	
50	27600	33750	22150	11080	31390	26510	18120	9070	21120	7800	21360	22010	
Mean	25430	29478	23380	12450	31512	27708	23184	9290	25102	8108	27072	22026	
STDEV	2210	4312	1504	5352	7148	6153	3678	584	3883	697	3927	2852	
MAX	27600	34100	25450	17550	41210	34500	27850	10330	29500	9270	30910	24590	
MIN	21700	24400	22050	4300	24310	20300	18120	8950	21120	7420	21360	17360	

Zn Units mg kg-1	Depth	1	2	3	4	5	6	7	8	9	10	11	12
	10	85	77	75	32	59	59	75	20	69	21	82	66
20	85	80	81	61	73	82	83	24	70	23	82	84	
30	80	69	82	70	98	80	82	34	77	31	81	87	
40	92	89	83	76	83	62	67	32	76	28	72	80	
50	88	84	74	62	80	65	56	23	68	24	71	77	
Mean	86	80	79	60	79	70	73	27	72	25	78	79	
STDEV	4.4	7.5	4.2	16.9	14.3	10.6	11.3	6.1	4.2	4	5.6	8.1	
MAX	92	89	83	76	98	82	83	34	77	31	82	87	
MIN	80	69	74	32	59	59	56	20	68	21	71	66	

As Units mg kg-1	Depth	1	2	3	4	5	6	7	8	9	10	11	12
	10	2.5	0.9	2	1.1	0.03	0.01	2.9	0.02	3.60E-05	0.9	0.06	0.06
20	5.5	1.4	1.3	1.1	0.09	0.03	0.4	0.01	1.60E-05	2.8	0.01	0.01	
30	0.8	5.3	5.4	1.8	0.05	0.03	1.9	0.02	3.60E-05	0.9	0.03	0.04	
40	1.8	6.1	4.3	2.3	0.01	0.01	1.8	0.01	1.60E-05	1	0.01	0.01	
50	4.3	1.1	3.2	1.9	0.04	0.04	1.4	0.01	1.60E-05	0.9	0.01	0.01	
Mean	3	3	3	2	0	0	2	2	0	0	1	0	
STDEV	1.9	2.5	1.7	0.5	0	0	0.9	0.9	0	0	0.8	0	
MAX	6	6	5	2	0	0	3	3	0	0	3	0	
MIN	1	1	1	1	0	0	0	0	0	0	1	0	

	Depth	1	2	3	4	5	6	7	8	9	10	11	12
Ba Units mg kg-1	10	255	352	246	217	310	353	275	133	210	102	315	273
	20	215	198	313	266	364	424	308	148	270	114	283	287
	30	208	241	312	269	432	418	306	160	275	123	270	291
	40	360	238	320	234	362	407	239	97	269	121	243	260
	50	350	205	235	204	357	331	260	81	234	117	187	250
	Mean	278	247	285	238	365	387	278	124	252	115	260	272
	STDEV	73	61.9	41.1	29	43.6	41.9	29.7	33.7	28.4	8.3	48.1	17.4
	MAX	360	352	320	269	432	424	308	160	275	123	315	291
	MIN	208	198	235	204	310	331	239	81	210	102	187	250

	Depth	1	2	3	4	5	6	7	8	9	10	11	12
Cr Units mg kg-1	10	29	25	22	12	36	31	40	6.2	30	9.9	46	33
	20	29	24	26	19	39	41	42	9.7	43	11	50	46
	30	31	21	25	22	47	38	42	14	32	12	41	42
	40	32	29	27	25	41	30	37	14	32	10	42	38
	50	27	27	23	22	40	37	29	8.6	31	9.8	32	33
	Mean	30	25	25	20	41	35	38	11	34	11	42	38
	STDEV	1.9	3	2.1	4.9	4	4.7	5.4	3.4	5.3	0.9	6.7	5.7
	MAX	32	29	27	25	47	41	42	14	43	12	50	46
	MIN	27	21	22	12	36	30	29	6	30	10	32	33

	Depth	1	2	3	4	5	6	7	8	9	10	11	12
Cu Units mg kg-1	10	37	33	35	14	28	25	31	7.3	29	10	37	26
	20	38	35	42	27	43	34	35	10	33	12	37	36
	30	39	36	40	30	39	30	35	13	32	13	33	37
	40	45	42	41	34	31	27	28	12	32	10	30	28
	50	42	24	39	30	32	30	29	18	29	10	24	25
	Mean	40	34	39	27	35	29	32	12	31	11	32	30
	STDEV	3.3	6.5	2.7	7.7	6.2	3.4	3.3	4	1.9	1.4	5.4	5.7
	MAX	45	42	42	34	43	34	35	18	33	13	37	37
	MIN	37	24	35	14	28	25	28	7	29	10	24	25

	Depth	1	2	3	4	5	6	7	8	9	10	11	12
Li Units mg kg-1	10	38	36	29	11	10	10	9.3	7.4	10	9.8	13	8.9
	20	41	40	32	23	21	17	11	9.5	12	10.5	12	12
	30	39	43	30	29	26	18	12	11	11	11	9.6	11
	40	45	43	33	30	26	12	10	10	9.7	9.9	10	9.5
	50	65	27	34	24	24	10	9.6	9.7	9.5	9.5	9.5	9.6
	Mean	46	38	32	23	21	13	10	10	10	10	11	10
	STDEV	11.2	6.7	2.1	7.6	6.7	3.8	1.1	1.3	1	0.6	1.6	1.3
	MAX	65	43	34	30	26	18	12	11	12	11	13	12
	MIN	38	27	29	11	10	10	9	7	10	10	10	9

	Depth	1	2	3	4	5	6	7	8	9	10	11	12
Mn Units mg kg-1	10	800	720	709	322	535	527	741	432	589	474	745	536
	20	810	718	703	610	585	650	723	502	613	468	748	740
	30	800	703	710	579	700	610	691	550	633	476	671	756
	40	883	820	779	706	588	571	609	474	603	402	576	700
	50	860	767	780	685	650	582	472	400	599	396	493	695
	Mean	831	746	736	580	612	588	647	472	607	443	647	685
	STDEV	38.4	48	39.6	153.6	64.1	45.8	110.2	58.7	16.7	40.5	110.8	87.5
	MAX	883	820	780	706	700	650	741	550	633	476	748	756
	MIN	800	703	703	322	535	527	472	400	589	396	493	536

	Depth	1	2	3	4	5	6	7	8	9	10	11	12
Ni	10	26	20	23	12	21	20	21	7.4	23	10	29	22
Units	20	24	20	23	20	24	27	27	10	26	11	29	30
mg kg-1	30	25	23	26	22	33	26	27	13	27	12	30	30
	40	28	25	25	25	27	21	22	12	27	11	25	27
	50	25	20	24	20	24	24	18	9.1	22	10	21	23
	Mean	26	22	24	20	26	24	23	10	25	11	27	26
	STDEV	1.5	2.3	1.3	4.8	4.5	3	3.9	2.2	2.3	0.8	3.8	3.8
	MAX	28	25	26	25	33	27	27	13	27	12	30	30
	MIN	24	20	23	12	21	20	18	7	22	10	21	22

	Depth	1	2	3	4	5	6	7	8	9	10	11	12
Pb	10	33	14	21	9.5	11	13	12	6.2	14	5.2	19	17
Units	20	24	13	20	10	14	16	17	7.9	18	6.1	17	18
mg kg-1	30	23	17	24	13	15	16	19	8.6	16	6.4	15	18
	40	32	17	25	15	13	12	10	5.5	14	5.1	13	16
	50	30	16	20	15	12	14	9.8	4.6	12	5.3	12	15
	Mean	28	15	22	13	13	14	14	7	15	6	15	17
	STDEV	4.6	1.8	2.3	2.6	1.6	1.8	4.2	1.7	2.3	0.6	2.9	1.3
	MAX	33	17	25	15	15	16	19	9	18	6	19	18
	MIN	23	13	20	10	11	12	10	5	12	5	12	15

	Depth	1	2	3	4	5	6	7	8	9	10	11	12
Ti	10	480	292	323	253	741	609	860	145	810	270	1100	780
Units	20	439	384	387	206	680	941	871	237	860	311	1130	1058
mg kg-1	30	455	636	470	386	1250	841	990	409	961	371	1008	1008
	40	480	591	416	331	1100	625	810	391	851	331	940	950
	50	473	526	389	276	921	610	648	267	800	307	740	860
	Mean	465	486	397	290	938	725	836	290	856	318	984	931
	STDEV	18	144.2	53.2	69.9	239.4	155.6	124	110.4	63.9	36.9	155.6	112.1
	MAX	480	636	470	386	1250	941	990	409	961	371	1130	1058
	MIN	439	292	323	206	680	609	648	145	800	270	740	780

	Depth	1	2	3	4	5	6	7	8	9	10	11	12
SO₄²⁻	10	6878	1523	1520	687	3531	1189	1081	854	690	1854	1520	690
Units	20	9556	7046	9891	680	687	1844	1523	1687	1356	1520	1525	1390
mg kg-1	30	12066	29303	8531	12903	695	6870	1021	1189	352	520	680	1300
	40	30290	29500	12066	20250	519	1022	884	855	190	499	180	520
	50	26950	9054	9054	2010	2528	1691	851	519	180	488	181	510
	Mean	17148	15285	8212	7306	1592	2523	1072	1021	554	976	817	882
	STDEV	10697	13178	3977	8869	1361	2454	269	441	494	660	675	430
	MAX	30290	29500	12066	20250	3531	6870	1523	1687	1356	1854	1525	1390
	MIN	6878	1523	1520	680	519	1022	851	519	180	488	180	510

	Depth	1	2	3	4	5	6	7	8	9	10	11	12
Cl	10	88553	162170	301068	1280	530	870	1060	226	282	207	2081	2295
Units	20	41920	87950	172407	3018	1238	4925	730	2230	664	197	630	4858
mg kg-1	30	38374	158900	170078	2700	220	392	910	730	668	227	186	123
	40	46097	140400	53182	1980	520	1031	377	150	179	214	146	197
	50	50360	58900	117818	1690	280	640	550	110	160	211	143	210
	Mean	53061	121664	162911	2134	558	1572	725	689	391	211	637	1537
	STDEV	20343	45973	91245	716	405	1890	273	897	256	11	833	2071
	MAX	88553	162170	301068	3018	1238	4925	1060	2230	668	227	2081	4858
	MIN	38374	58900	53182	1280	220	392	377	110	160	197	143	123

	Depth	1	2	3	4	5	6	7	8	9	10	11	12
CaCO₃	10	172	153	148	132	177	482	490	141	474	120	549	501
Units	20	169	156	186	165	470	465	483	127	468	143	470	477
mg kg⁻¹	30	124	105	149	151	410	475	420	153	470	120	453	470
	40	163	225	158	169	422	480	485	155	460	122	480	420
	50	223	289	141	155	478	469	474	133	470	123	463	360
	Mean	170	186	156	154	391	474	470	142	468	126	483	446
	STDEV	35.3	71.9	17.6	14.5	123.4	7.2	28.8	12.2	5.2	9.8	38.2	56.2
	MAX	223	289	186	169	478	482	490	155	474	143	549	501
	MIN	124	105	141	132	177	465	420	127	460	120	453	360

	Depth	1	2	3	4	5	6	7	8	9	10	11	12
pH	10	7.21	7.3	7.21	7.46	7.39	6.94	7.62	8.05	6.81	8.15	7.31	6.84
Units	20	7.41	7.32	7.41	7.66	7.3	6.84	7.63	8.02	7.35	8.29	7.37	6.89
S.U	30	7.54	7.23	7.54	7.86	7.62	7.33	7.27	8.25	6.97	8.5	7.47	7.23
	40	8.05	7.43	8.05	7.67	7.15	7.42	7.16	8.34	7.02	8.31	7.22	7.01
	50	7.96	7.47	7.96	7.68	7.81	7.62	7.34	8.25	7.64	8.39	7.54	6.97
	Mean	7.6	7.4	7.6	7.7	7.5	7.2	7.4	8.2	7.2	8.3	7.4	7
	STDEV	0.4	0.1	0.4	0.1	0.3	0.3	0.2	0.1	0.3	0.1	0.1	0.2

Supplementary Tables

Core Locations

Sample ID	Latitude	Longitude
1	35.518350°	-115.379900°
2	35.518400°	-115.359616°
3	35.519900°	-115.354383°
4	35.521400°	-115.345967°
5	35.523617°	-115.341933°
6	35.523900°	-115.340566°
7	35.524267°	-115.338217°
8	35.524367°	-115.337933°
9	35.524600°	-115.336166°
10	35.524483°	-115.333917°
11	35.525400°	-115.333417°
12	35.525933°	-115.330558°

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